

Drug Recognition Expert 7-Day School Student Manual HS 172 R9/02 (September 2002)

Document Description	Page(s) Withheld	Exemption	Comments
Drug Recognition Expert 7-Day School Instructor Manual HS 172 R09/02 (September Revision) The DRE School Pre-Test	53-58	Exam information Test questions - Employment and Licensing - RCW 42.56.250(1)	Test questions, scoring keys, and other examination data used to administer a license, employment, or academic examination are exempt from production.
Drug Recognition Expert 7-Day School Instructor Manual HS 172 R09/02 (September Revision) Pre-Test Answer Key	59	Exam information scoring keys - Employment and Licensing - RCW 42.56.250(1)	Test questions, scoring keys, and other examination data used to administer a license, employment, or academic examination are exempt from production.
Drug Recognition Expert 7-Day School Instructor Manual HS 172 R09/02 (September Revision) Quiz Number One	60-65	Exam information Test questions - Employment and Licensing - RCW 42.56.250(1)	Test questions, scoring keys, and other examination data used to administer a license, employment, or academic examination are exempt from production.
Drug Recognition Expert 7-Day School Instructor Manual HS 172 R09/02 (September Revision) Answer Key for Quiz Number One	66	Exam information scoring keys - Employment and Licensing - RCW 42.56.250(1)	Test questions, scoring keys, and other examination data used to administer a license, employment, or academic examination are exempt from production.
Drug Recognition Expert 7-Day School Instructor Manual HS 172 R09/02 (September Revision) Quiz Number Two	67-72	Exam information Test questions - Employment and Licensing - RCW 42.56.250(1)	Test questions, scoring keys, and other examination data used to administer a license, employment, or academic examination are exempt from production.
Drug Recognition Expert 7-Day School Instructor Manual HS 172 R09/02 (September Revision) Answer Key for Quiz Number Two	73	Exam information scoring keys - Employment and Licensing - RCW 42.56.250(1)	Test questions, scoring keys, and other examination data used to administer a license, employment, or academic examination are exempt from production.
Drug Recognition Expert 7-Day School Instructor Manual HS 172 R09/02 (September Revision) Quiz Number Three	74-79	Exam information Test questions - Employment and Licensing - RCW 42.56.250(1)	Test questions, scoring keys, and other examination data used to administer a license, employment, or academic examination are exempt from production.
Drug Recognition Expert 7-Day School Instructor Manual HS 172 R09/02 (September Revision) Answer Key for Quiz Number Three	80	Exam information scoring keys - Employment and Licensing - RCW 42.56.250(1)	Test questions, scoring keys, and other examination data used to administer a license, employment, or academic examination are exempt from production.
Drug Recognition Expert 7-Day School Instructor Manual HS 172 R09/02 (September Revision) Quiz Number Four	81	Exam information Test questions - Employment and Licensing - RCW 42.56.250(1)	Test questions, scoring keys, and other examination data used to administer a license, employment, or academic examination are exempt from production.

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Drug Recognition Expert 7-Day School Instructor	87	Exam information scoring	Test questions, scoring keys, and other
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Answer Key for Quiz Number Four	88-97	Exam information Test	Test questions, scoring keys, and other
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Answer Key for Quiz Number Five			examination are exempt from production.
Drug Recognition Expert 7-Day School Instructor	99-100	Exam information Other	Test questions, scoring keys, and other
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Drug Recognition Expert 7-Day School Instructor	124-128	Exam information scoring	Test questions, scoring keys, and other
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Drug Recognition Expert 7-Day School Instructor Manual	129-150	Exam information Test questions - Employment and	Test questions, scoring keys, and other examination data used to administer a
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DRE3A Feedback Report (Final Exam B Answer Sheet)			examination are exempt from production.
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Manual		Examination Data used to	examination data used to administer a
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HS 172 R09/02 (September Revision) Final Knowledge Exam		Licensing - RCW 42.56.250(1)	license, employment, or academic examination are exempt from production.
Drug Recognition Expert 7-Day School Instructor Manual HS 172 R09/02 (September Revision) Knowledge Examination Answer Key	169-181	Exam information scoring keys - Employment and Licensing - RCW 42.56.250(1)	Test questions, scoring keys, and other examination data used to administer a license, employment, or academic examination are exempt from production.
Drug Recognition Expert 7-Day School Instructor Manual HS 172 R09/02 (September Revision) A Self-Test for Review and Study	821-827	Exam information Test questions - Employment and Licensing - RCW 42.56.250(1)	Test questions, scoring keys, and other examination data used to administer a license, employment, or academic examination are exempt from production.
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Acknowledgments

The NHTSA and IACP would like to thank the following individuals for their contributions in updating and revising the DRE curricula.

Sgt. Murray Conrad

Albuquerque Police Department

Deputy Mark George Boulder County Sheriff's Office

Sgt. Art Haversat, Retired Los Angeles Police Department

Sgt. Tom Page, Retired Los Angeles Police Department

Sgt. Doug Paquette New York State Police Lt. Mike Pryor

Tucson Police Department

Sgt. Brian Schafer

Minneapolis Police Department

Sgt. Deb Schroder

California Highway Patrol

Sgt. Dan Webb

Texas Department of Public Safety

Sgt. Jose Zarraga

Miami Dade Police Department

Thanks to the states of Arizona, Colorado, Maryland, New York, Oregon and Texas for pilot testing the training curricula.

A special thanks to Pam McCaskill, TSI in Oklahoma City for her patience and tireless efforts in typing and correlation of the DRE manuals.

DRUG EVALUATION AND CLASSIFICATION TRAINING "THE DRUG RECOGNITION EXPERT SCHOOL"

ADMINISTRATOR'S GUIDE

SEPTEMBER, 2002 EDITION

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A. Purpose of this Document

This Administrator's Guide provides an introduction to and an overview of the seven-day classroom training course on drug evaluation and classification. This course is perhaps better known as **The DRE School**. It is the second in a series of three stages of training that, collectively, prepare persons to serve as Drug Recognition Experts (DREs).

Throughout this manual, the term "DRE" is used to designate an individual who is specially-trained to conduct examinations of drug-impaired drivers. In some participating agencies, the term stands for "Drug Recognition Expert"; in others, it means "drug recognition evaluator", and in others, "drug recognition examiner". In addition, some agencies use the term "DRT" -- Drug Recognition Technician -- and others prefer "DRS" -- Drug Recognition Specialist. All of these and similar terms are acceptable and considered synonymous. But for this training program, the standard term is DRE.

It is worth repeating that this seven-day DRE School is neither the beginning nor the end of an officer's preparation to serve as a DRE. No one can be admitted to this course unless he or she has successfully completed the two-day program titled "Preliminary Training for Drug Evaluation and Classification" (the "PRE-School"), or demonstrates that he or she has mastered the subject-matter of that PRE-School via previous training and experience. And, the fact that an officer successfully completes this seven-day program does <u>not</u> qualify him or her to serve as a DRE. He or she still must complete the Certification Phase of training, a supervised on-the-job phase in which the trainee conducts examinations of persons actually under arrest on suspicion of drug impairment.

This seven-day course, then, is only the middle phase of DRE training. But it is a very important phase. It is during this phase that the student will learn to conduct systematic and standardized examinations of persons suspected of drug impairment to determine:

- (1) Whether the suspect actually is impaired; and if so,
- (2) Whether the impairment is drug- or medically-related; and if drugs,
- (3) The broad category or combination of categories of drugs that is the likely cause of the observed impairment.

This Administrator's Guide is concerned only with the second phase of training. During this phase, the student becomes familiar with the various types of drugs that people use and -- too often -- abuse. The student learns how the different drugs affect people, and especially how they affect a person's ability to operate a vehicle. The student learns how the different drugs manifest their presence in an individual. In particular, the student learns how to examine a suspect's eyes and vital signs to detect evidence of various kinds of drugs. By the time the student successfully completes the training, he or she is able to conduct a complete drug evaluation and classification examination, and is able to describe the evidence that the examination will disclose to help determine if the suspect suffers a medical condition or if a suspect is under the influence of a particular category or combination of categories of drugs.

This Administrator's Guide is intended to facilitate planning and implementation of the Drug Evaluation and Classification Classroom Training Program. The Guide overviews the 7-day course of instruction, and the documents and other materials that make up the curriculum package for the course. It describes course administrative requirements and offers guidelines for discharging those requirements satisfactorily. It outlines the preparatory work that must be accomplished by a law enforcement agency before the course can be offered to that agency's personnel. And, it outlines the follow-up work that should be undertaken to ensure that the highest possible quality of instruction continues to be delivered, during all phases of a DRE's training.

Before addressing the details of this classroom training in Drug Evaluation and Classification Procedures, a few words are appropriate concerning the procedures themselves. In particular, it is important to make clear what the Drug Evaluation and Classification Procedures are <u>not</u>:

These procedures are <u>not</u> a field test, or a pre-arrest investigative tool. It is highly unlikely that they could be conducted with adequate care in an outdoors, scene-of-investigation setting. In any event, they are not designed to provide probable cause for a suspect's arrest. Rather, they are a post-arrest investigative tool, intended for application to arrestees for whom there is at least some articulable suspicion of drug use or drug impairment.

- These procedures do <u>not</u>, generally speaking, disclose what <u>specific</u> drug or drugs the suspect has used. That may seem to be a startling, and upsetting statement. Nevertheless, it is true. What the procedures <u>will</u> do, however, is to disclose (with reasonable accuracy) the broad category or combination of categories that produce distinguishable "signatures" visible to a qualified DRE. Some of the categories include relatively few individual drugs. Others include many drugs. The DRE can tell, usually, if a particular category is present. But except in special circumstances, he or she cannot tell which individual member of that category is the drug in question. Thus for example, a DRE usually will not be able to distinguish a person impaired by diazepam from a person impaired by secobarbital. Will not be able to tell the difference between a codeine-impaired subject and someone under the influence of Demerol. Won't see a difference between someone under the influence of peyote and someone under the
- The procedures are <u>not</u> a substitute for chemical testing. Laboratory analysis of blood samples by qualified personnel remains an important step in the acquisition of evidence in drug-related cases. The drug evaluation and classification procedures provide articulable bases for requesting a suspect to supply the urine or blood sample; guide the laboratory technicians toward the general categories of drugs they can expect to find in the sample; and, disclose important evidence to supplement the laboratory analysis. But the drug recognition expert does <u>not</u> eliminate the need for the laboratory technician.

None of the foregoing remarks is intended to lessen the importance of the drug evaluation and classification procedures. A cadre of skilled DREs definitely will enhance a department's ability to recognize and convict persons under the influence of drugs. The DRE is a very important "weapon" in law enforcement's anti-drug arsenal. But the DRE is not the entire arsenal.

One final word of introduction: the primary orientation of this course is toward traffic law enforcement. Without doubt, persons under the influence of drugs endanger society in many ways. But it is the danger they cause as drivers of motor vehicles that is of principal interest here. This course assumes that the DRE will devote his or her skills in large part to conducting examinations of suspected impaired drivers. This is not to say that the skills that this training seeks to develop do not have many non-traffic applications. Nevertheless, it is the traffic applications that will receive most of the student's attention.

B. Overview of the Course

1. For whom is the training intended?

This training definitely is not intended for just anyone. The candidate DRE isn't just any police officer, but an officer who already has some very special knowledge and skills, and a very definite commitment to DWI and drug enforcement. And, that officer isn't employed by just any department. Instead, he or she works for a department that has taken pains to provide the command and logistics support needed to allow the DRE to function at maximum effectiveness. And the department has concrete proof of its commitment to deterring impaired driving. Finally, that department doesn't serve just any community or state. Instead, it operates in a jurisdiction that has a legal and political framework that is consistent with effective enforcement of drug-impaired driving violations.

The following lists the prerequisites and desirable characteristics of the students for whom this training is intended; of the departments that employ those students; and, of the communities served by those departments.

a. Student Prerequisites

To be considered a qualified candidate for this training, the proposed student must be a law enforcement officer or an employee of a public criminal justice agency or an institution providing law enforcement training, and must:

- o have achieved the learning objectives of the two-day PRE-School;
- o have demonstrated proficiency in the use of the Standardized Field Sobriety Tests (i.e., Horizontal Gaze Nystagmus, walk and turn and one leg stand);
- o have good communications skills, and a demonstrated ability to testify in court;
- o be willing to continue to serve as a DRE for at least two years following completion of the training.

Of course, it is highly <u>desirable</u>, although not essential, that the proposed student have prior knowledge of drug symptomatology and experience in drug enforcement.

b. <u>Departmental Prerequisites</u>

To be considered qualified to submit students for this training, the interested law enforcement agency <u>must</u>:

- o have active drug enforcement and DWI enforcement programs;
- be pro-active in training officers in Standardized Field Sobriety Testing; also, the training must be consistent with NHTSA guidelines, and the agency must maintain records of officers' Standardized Field Sobriety Testing enforcement activities;
- o have access to adequate chemical testing resources to support the drug evaluation and classification program, and ensure effective prosecution of drug-impaired subjects;
- o have adequate facilities and equipment to support the drug evaluation and classification examinations;
- o have an management information system (MIS) capable of accurately tracking alcohol and drug enforcement activities:
- o demonstrate the firm support and commitment of the chief law enforcement officer and other appropriate officials for the drug evaluation and classification program. Evidence of this support includes but is not limited to:
 - Willingness to assign at least one person of supervisory rank to become a certified DRE and to manage and coordinate the agency's Drug Evaluation and Classification Program.
 - Willingness to upgrade the agency's MIS, as necessary, to track progress of DRE training; drug and DWI arrests; DRE evaluations; results of toxicological examinations; and, case filings and dispositions.
 - Willingness to conduct DRE training in a manner that complies fully with NHTSA curricula and guidelines.
 - Willingness to adopt NHTSA-approved DRE evaluation forms.
 - Willingness to authorize DREs and DRE candidates to devote sufficient time to the DRE function to develop and maintain proficiency.

- Willingness to provide the services of qualified DRE instructors to assist NHTSA in training candidate DREs from other agencies.

c. <u>Legal and Political Prerequisites</u>

To be considered qualified to recommend a law enforcement agency for this training, a state or community <u>must</u> have laws or courtestablished precedents that:

- o specifically allow for the analysis of chemical samples obtained from persons suspected of impaired driving, to determine the presence and/or concentration of drugs other than alcohol;
- o allow the arresting officer or law enforcement agency to specify the chemical test or tests (e.g., blood, breath or urine) to be given to suspected impaired drivers;
- o specifically facilitate testing for drugs other than alcohol.

In addition, it is <u>desirable</u> that the state or community have laws that:

- o make the fact of the driver's refusal to submit to the test or tests admissible in court;
- o make it an offense to be under the influence of alcohol and/or illicit drugs, whether or not the person is operating a vehicle.

Furthermore, the state's or community's prosecutors must:

- o demonstrate a willingness to introduce Standardized Field Sobriety Test evidence in alcohol/drug cases;
- o express a willingness to participate in this training to become familiar with drug evaluation and classification procedures and related information.

The state's or community's judges must:

- o demonstrate a willingness to accept and consider Standardized Field Sobriety Test evidence in alcohol/drug cases;
- o express a willingness to consider drug evaluation and classification evidence in alcohol/drug cases.

Finally, it is desirable that the jurisdiction's political and community leaders express support for the drug evaluation and classification program.

2. What are the purposes of the course?

The ultimate goal of this course is to help prevent crashes, deaths and injuries by improving enforcement of drug-impaired driving violations. It is not exactly clear how many drug-impaired drivers are on our nation's roads, or how many crashes they cause. But even the most conservative estimates indicate that these drivers kill thousands of Americans, and injure at least tens of thousands of others each year.

3. What will the students get out of this course?

The classroom training course is designed to help the students achieve three broad goals, and eight specific learning objectives.

<u>Goals</u>: The student who successfully completes this phase of DRE training will be able to...

- ... distinguish if an individual is under the influence of a drug or drugs other than alcohol, or under the combined influence of alcohol and other drugs, or suffering from some injury or illness that produces signs similar to alcohol/drug impairment;
- ... identify the broad category or categories of drugs inducing the observable signs of impairment; and,
- ... progress to the Certification Phase of the training.

Objectives: In order to pass this course, the student must be able to...

- ... describe the involvement of drugs in impaired driving incidents;
- ... name the seven categories of drugs and recognize their effects;
- ... describe and properly administer the psychophysical and physiologic evaluations used in the drug evaluation and classification procedures;
- ... document the results of the drug evaluation and classification examination;
- ... properly interpret the results of the examination;

- ... prepare a narrative drug influence report;
- ... discuss appropriate procedures for testifying in typical drug evaluation and classification cases; and,
- ... maintain an up-to-date relevant resume.
- 4. What subject matter does the course cover?

The course focuses primarily on two broad topics:

- (1) The examinations, observations, measurements, etc. that constitute the drug evaluation and classification procedures.
- (2) The nature, effects, signs and symptoms of each of the seven categories of drugs, and of the combination of categories.

More specifically, the course provides formal presentations on:

- o Drugs in Society and in Motor Vehicle Operation.
- o Development and Effectiveness of the Drug Evaluation and Classification Procedures.
- o An Overview of Physiology and Drugs.
- o An Overview of the Drug Evaluation and Classification Procedures.
- o Eye Examinations (Horizontal Gaze Nystagmus; Vertical Gaze Nystagmus; Lack of Convergence; Estimation of Pupil Size; Pupil Reaction to Light).
- o Vital Signs Examinations (Pulse Rate; Blood Pressure; Temperature)
- o The Physician's Desk Reference, and other reference materials.
- o The Seven Categories of Drugs (Central Nervous System Depressants; Central Nervous System Stimulants; Hallucinogens; Phencyclidine; Narcotic Analgesics; Inhalants; Cannabis).
- o Drug Combinations.
- o Narrative Arrest Report in Drug Evaluation Cases.

- o Case Preparation and Testimony.
- o Resume Preparation and Maintenance.

5. What activities take place during the training?

Formal presentations, or lectures, occupy approximately one-half of the course. These presentations cover the content topics outlined earlier. The presentations are supplemented by video tape segments, and by reading material contained in the Student's Manual.

Most of the remainder of the course is devoted to demonstrations and hands-on practice of the drug evaluation and classification procedures. Students repeatedly practice in teams, developing and sharpening their skills in administering eye examinations, vital signs examinations, and other components of the drug recognition expert's job. Students also participate in several test interpretation practice sessions, in which they review sample drug evaluation and classification reports and identify the category or categories of drugs responsible for the "evidence" described in the reports.

The remaining major activity is testing of the students' knowledge and proficiency. A written knowledge examination is administered, at the end of the course. A formal assessment of each student's skill in administering the drug evaluation and classification procedures is conducted during the next-to-last session.

6. How long does the training take?

This classroom training course occupies 7 training days. A typical schedule calls for each class day to begin at 8:00 am and conclude at 5:00 pm. A one-hour lunch period and hourly breaks of 10 minutes are accommodated in that schedule.

The course is divided into thirty-two (32) sessions. Of those, two are review sessions, conducted after normal class hours on the fourth and sixth days of the School. No student can progress to the Certification Phase of training until he or she has attended all mandatory sessions. In the event that some emergency causes a student to miss all or a portion of a session, after-hours tutoring must be conducted for that student prior to his or her enrollment in Certification training.

The titles, durations and sequence of the sessions are given below.

Session I

Introduction and Overview

(1 hour, 50 minutes)

Session II

Drugs in Society and in Motor Vehicle Operation (50 minutes)

Session III

Development and Effectiveness of the

DRE Program

(50 minutes)

Session IV

Overview of Drug Recognition Expert Procedures (2 hours, 30 minutes)

Session V

Eye Examinations

(1 hour, 45 minutes)

Session VI

Physiology & Drugs: An Overview

(2 hours)

Session VII

Examination of Vital Signs

(2 hours)

Session VIII

Demonstration of the Evaluation Sequence

(1 hour, 20 minutes)

Session IX

Central Nervous System Depressants

(1 hour, 45 minutes)

Session X

Central Nervous System Stimulants

(1 hour, 45 minutes)

Session XI

Practice: Eye Examinations

(1 hour)

Session XII

Alcohol Workshop

(1 hour, 45 minutes)

Session XIII

Physician's Desk Reference and Other

(30 minutes)

Reference Sources

Session XIV

Hallucinogens

(1 hour, 45 minutes)

Session XV

Practice: Test Interpretation

(45 minutes)

Session XVI

Phencyclidine (PCP)

(1 hour, 40 minutes)

Session XVII

Narcotic Analgesics

(3 hours)

REVIEW SESSION

(Mid-Course Review)

(2 hours, 30 minutes)

Session XVIII

Practice: Test Interpretation

(45 minutes)

Session XIX

Inhalants

(1 hour, 35 minutes)

Session XX

Practice: Vital Signs Examinations

(50 minutes)

Session XXI

Cannabis

(1 hour, 35 minutes)

Session XXII

Overview of Signs and Symptoms

(1 hour)

Session XXIII

Resume Preparation and Maintenance

(50 minutes)

Session XXIV

Drug Combinations

(1 hour, 50 minutes)

Session XXV

Practice: Test Interpretation

(45 minutes)

Session XXVI

Preparing the Narrative Report

(50 minutes)

Session XXVII

Practice: Test Administration

(1 hour, 45 minutes)

Session XXVIII

Case Preparation and Testimony

(1 hour 30 minutes)

REVIEW SESSION

Review of the DRE School

(2 hours, 30 minutes)

Session XXIX

Classifying a Suspect (Role Play)

(4 hours)

Session XXX

Transition to the Certification

(2 hours, 30 minutes)

Phase of Training

NOTE: All sessions of this course are absolutely essential. No short-cuts are permissible.

A model schedule for the seven-day course is given on the next page.

Alternate Schedule #1 combines the Pre-School and Seven-Day School.

Alternate Schedule #2 combines the DWI Detection and Standardized Field Sobriety Testing, Pre-School, and Seven-Day School.

If you use Alternate Schedule #1 or #2, you will need to make copies of those schedules for the students.

THE DRE SCHOOL - SCHEDULE (page 1)

WEDNEST	THURSDAY	FRIDAY
0800-0850 SESSION I: Introduction & Overview	0800-0850 SESSION V: (cont)	0800-0850 SESSION IX: Central Nervous System Depressants
0850-0900 BREAK	0850-0900 BREAK	0850-0900 BREAK
0900-1000 SESSION I. (cont)	0900-1005 SESSION VI: Physiology & Drugs (Overview)	0900-1000 SESSION IX: (cont)
1000-1010 BREAK	1005-1015 BREAK	1000-1010 BREAK
1010-1030 Pre-Test	1015-1110 SESSION VI: (cont)	1010-1100 SESSION X: Central Nervous System CNS Stimulants
1030-1120 SESSION II: Drugs In Society & In Motor Vehicle Operation	1110-1120 BREAK	1100-1110 BREAK
1120-1130 BREAK	1120-1200 SESSION VII: Examination of Vital Signs	1110-1200 SESSION X: (cont)
1130-1230 SESSION III: Development & Effectiveness of the Program	1200-1300 LUNCH	1200-1300 LUNCH
1230-1330 LUNCH	1300-1400 SESSION VII: (cont)	1300-1400 SESSION XI: Eye Examinations
1330-1440 SESSION IV: Overview of Drug Recognition Expert Procedures	1400-1410 BREAK	1400-1415 BREAK
1440-1450 BREAK	1410-1430 SESSION VII: (cont)	1415-1700 SESSION XII: Alcohol Workshop
1450-1550 SESSION IV: (cont)	1430-1515 SESSION VIII: Demonstrations of the Evaluation Sequence	
1550-1600 BREAK	1515-1530 BREAK	
1600-1630 SESSION IV: (cont)	1530-1605 SESSION VIII: (cont)	
1630-1730 SESSION V: Eye Examinations	1605-1635 QUIZ NUMBER ONE	

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THE DRE SCHOOL - SCHEDULE (page 2)

MONDAY	TUESDAY	WEDNESDAY	THURSDAY
0800-0830 SESSION XIII: Physician's Desk Reference & Other Reference Sources	0800-0820 QUIZ NUMBER TWO	0800-0915 SESSION XXIV: Drug Combinations	0800-1000 FINAL EXAM
0830-0915 SESSION XIV: Hallucinogens	0820-0850 SESSION XVII: (cont)	0915-0930 SESSION XXIV: (cont)	1000-1015 BREAK
0915-0930 BREAK	0850-0900 BREAK	1005-1050 SESSION XXV: Practice Test Interpretation	1015-1200 SESSION XXIX: Classifying a Suspect-Role Play
0930-1030 SESSION XIV: (cont)	0900-0945 SESSION XVIII: Practice Test Interpretation	1050-1100 BREAK	1200-1300 LUNCH
1030-1045 BREAK	0945-1020 SESSION XIX: Inhalants	1100-1200 SESSION XXVI: Preparing the Narrative Report	1300-1600 ADMINISTRATION OF THE TEST VALIDATION
1045-1130 SESSION XV: Practice Test Interpretation	1020-1030 BREAK	1200-1300 LUNCH	1600-1630 SESSION XXX: Transition to Certification Training
1130-1200 SESSION XVI: Phencyclidine (PCP)	1030-1130 SESSION XIX: (cont)	1300-1430 SESSION XXVII: Practice Test Interpretation	1630-1700 Course Critique; Closing Remarks; Presentation of Certificates
1200-1300 LUNCH	1130-1145 BREAK	1430-1445 BREAK	
1300-1410 SESSION XVI: (cont)	1145-1300 SESSION XX: Practice Vital Signs Examinations	1445-1530 SESSION XXVIII: Case Preparation and Testimony	
1410-1420 BREAK	1300-1400 LUNCH	1530-1545 BREAK	
1420-1515 SESSION XVII: Narcotic Analgesics	1400-1530 SESSION XXI: Cannabis	1545-1630 SESSION XXVIII: (cont)	
1515-1530 BREAK	1530-1540 BREAK	1630-1700 QUIZ NUMBER FOUR	
1530-1630 SESSION XVII: (cont)	1540-1640 SESSION XXII: Overview of Signs and Symptoms	1700-1800 BREAK	
1630-1730 SESSION XVII: (cont)	1640-1650 BREAK	1800-2000 OPTIONAL REVIEW SESSION#2	
1730-1800 BREAK	1650-1730 SESSION XXIII: Resume Preparation & Maintenance		
1800-2030 OPTIONAL REVIEW SESSION#1	1730-1800 QUIZ NUMBER THREE		
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ALTERNATE SCHEDULE #1 COMBINED PRE-SCHOOL AND 7-DAY SCHOOL

Time	Session Title	D - 7-day DRE School P - Pre-School	Duration
8:00A - 10:00A	Introduction and Overview	D	2hrs
10:00A - 11:00A	Drugs and Society	D	1hr
11:00A - 12:00P	Development and Effectiveness	D	1hr
12:00P - 1:00P	Lunch		1hr
1:00P - 3:30P	Overview of DRE Classification Procedures	D	2.5hrs
3:30P - 5:00P	Psychophysical Tests	P ·	1.5hrs ·
	END OF DAY		
·			
8:00A - 11:00A	Eye Examinations	D	3hrs
11:00A - 12:00P	Vital Signs	D	1hr
12:00P - 1:00P	Lunch		1hr
1:00P - 2:30P	Vital Signs (cont.)	D	1.5hrs
2:30P - 4:00P	Overview of Signs and Symptoms	Р	1.5hrs
4:00P - 5:00P	Alcohol as a Drug	P	1hr
	END OF DAY		
8:00A - 9:30A	Demonstration of the Evaluation Sequence	D	1.5hrs
9:30A - 12:00P	Physiology of Drugs	D	2.5hrs
12:00P - 1:00P	Lunch		1hr
1:00P - 2:30P	Central Nervous System Depressants	D	1.5hrs
2:30P - 5:00P	Alcohol Workshop All Instructors	P	2.5hrs
	END OF DAY		

Time	Session Title	D - 7-day DRE School P - Pre-School	Duration
8:00A - 9:00A	Central Nervous System Depressants (cont.)	D	1hr
9:00A - 11:30A	Central Nervous System Stimulants	D	2.5hrs
11:30A - 12:00P	Quiz Number One	D	.5hr
12:00P - 1:00P	Lunch		1hr
1:00P - 2:00P	Eye Examinations	D	1hr
2:00P - 2:30P	PDR and Other Drug References	D	.5hr
2:30P - 5:00P	Review and Pre-School Final Examination	P	2.5hrs
	END OF DAY		
8:00A - 10:00A	Hallucinogens	D	2hrs
10:00A - 11:00A	Practice Test Interpretation	D	1hr
11:00A - 12:00P	Phencyclidine	D	1hr
12:00P - 1:00P	Lunch		1hr
1:00P - 2:00P	Phencyclidine (cont.)	D	1hr
2:00P - 4:00P	Mid-Course Review All Instructors	D	2hrs
	END OF DAY		
8:00A - 11:00A	Narcotic Analgesics	D	3hrs
11:00A - 12:00P	Practice Test Interpretation	D	1hr
12:00P - 1:00P	Lunch		1hr
1:00P - 2:00P	Inhalants	D	1hr
2:00P - 3:00P	Practice Vital Signs All Instructors	· D	1hr
3:00P - 4:00P	Quiz Number Two	D	.5hr
	END OF DAY		

Time	Session Title	D - 7-day DRE School P - Pre-School	Duration
8:00A - 11:00A	Cannabis	D	3hrs
11:00A - 12:00P	Overview of Signs and Symptoms	D	1hr
12:00P - 1:00P	Lunch		1hr
1:00P - 2:00P	Drug Combinations	D	1hr
2:00P - 2:30P	Quiz Number Three	D	.5hr
2:30P - 5:00P	Alcohol Workshop All Instructors	D	2.5hrs
	END OF DAY		,
8:00A - 9:00A	Drug Combinations	D	1hr
9:00A - 10:00A	Practice Test Interpretation	D	1hr
10:00A - 11:00A	Preparing the Narrative Report	D	1hr
11:00A - 12:00P	Practice Test Administration All Instructors	D	1hr
12:00P - 1:00P	Lunch		1hr
1:00P - 2:30P	Case Preparation and Testimony	D	1.5hrs
2:30P - 3:00P	Quiz Number Four	D	.5hr
3:00P - 5:00P	Final Course Review All Instructors	· D	2hrs
	END OF DAY		
· ·			
8:00A - 11:00A	Final Examination All Instructors	D	3hrs
11:00A - 12:00P	Transition to Certification Training	D	1hr
12:00P - 1:00P	Lunch		1hr
1:00P - 3:00P	Classifying a Suspect (Role Play) All Instructors	D	2hrs
3:00P - 4:00P	Graduation		2hrs

ALTERNATE SCHEDULE #2 COMBINED DWI DETECTION AND STANDARDIZED FIELD SOBRIETY, PRE-SCHOOL AND 7-DAY SCHOOL

WEEK ONE Day One	DURATION
Block I - Introduction and Overview (merger of DWI Detection and SFST manual session I and the DRE manual session I)	2hrs
SFST and DRE School Pre-tests	, .
Block 2 - Definition of drug and overview of the drug categories (modified Pre-School session I, Introduction and Overview)	1hr
Block 3 - Detection and Deterrence (SFST manual session II)	1hr
Block 4 - The Legal Environment (SFST manual session III)	45min
Block 5 - Overview of Detection, Notetaking and Testimony (SFST manual session IV)	45min
Block 6 - Phase One: Vehicle in Motion (SFST manual session V)	1hr
Block 7 - Phase Two: Personal Contact (SFST manual session VI)	1hr
Block 8 - Phase Three: Pre-Arrest Screening (SFST manual session VII)	30min
DAY TWO	
Block 9 - Concepts and Principles of the SFST (SFST manual session VIII, segments A (development and validity) and B (types of nystagmus)	1hr
Block 10 - Eye examinations (Pre-School manual session IV, segments A (purposes of the eye examinations) and B 1, 2 and 3 (procedures and clues for HGN, VGN, and Lack of Convergence)	1hr
Block 11 - Psychophysical Tests (Pre-School manual session III, segments A and B, Romberg and Walk and Turn)	1hr
Block 12 - Psychophysical Tests (Pre-School manual session III, segments C and D, One Leg Stand and Finger to Nose)	1hr
Block 13 - SFST Battery Demonstrations (SFST manual session IX, plus Romberg and Finger to Nose, utilizing the DRE order)	1hr
Block 14 - SFST Dry Run Practice (SFST manual session X, plus Romberg and Finger to Nose, in the DRE order)	1hr
Block 15 - Alcohol Correlation Study #1 (merger of SFST manual session XI and Pre-School manual session V)	2hrs

DAY THREE	DURATION
Block 16 - Alcohol as a Drug (Pre-School manual session VIII)	2hrs
Block 17 - Overview of Signs and Symptoms (Pre-School manual session VII)	1hr
Block 18 - Eye Examinations (Pre-School manual session IV, beginning with B4 (estimation of pupil size) through 5 (reaction to light)).	1hr
Block 19 - Drugs in Society and in Motor Vehicle Operation (DRE manual session II)	1hr
Block 20 - Development and Effectiveness (DRE manual session III)	2hrs
Block 21 - Review Session - SFST curriculum	1hr
DAY FOUR	
Block 22 - SFST Course Final Examination (SFST manual session X)	30min
Block 23 - Eye Examinations - Practice Session (merger of the practice sessions in DRE manual session XI and Pre-School manual session IV)	30min
Block 24 - Examination of Vital Signs (merger of Pre-School manual session VI and DRE manual session VII)	3hrs
Block 25 - Overview of Drug Evaluation and Classification Procedures (merger of Pre-School manual session II and DRE manual session IV)	1hr
Block 26 - Demonstrations of the Evaluation Sequence (DRE manual session VIII)	2hrs
Block 27 - Review Session - Pre-School Curriculum	1hr
DAY FIVE	
Block 28 - Pre-School Final Examination (Pre-School manual session X)	30min
Block 29 - Physiology and Drugs: An Overview	4hrs
Block 30 - SFST Report Writing (SFST manual session XIII and SFST practice session)	1hr, 30min
Block 31 - Alcohol Correlation Study #2 (merger of Pre-School manual session V and SFST manual session XIV; includes SFST Proficiency Test)	2hrs

WEEK TWO DAY SIX	DURATION
Quiz #1	30min
Block 32 - Physician's Desk Reference, CPS and Additional Resources (DRE manual session XIII)	2hrs
Block 33 - Methods of Administration and Elimination (Note: This is not a current standard manual session, but is an LAPD curriculum addition)	30min
Block 34 - Central Nervous System Depressants (DRE manual session IX)	2hrs
Block 35 - Central Nervous System Stimulants (DRE manual session X)	3hrs
DAY SEVEN	
Quiz #2	30min
Block 36 - Hallucinogens (DRE manual session XIV)	2hrs
Block 37 - Practice: Test Interpretation (DRE manual session XV)	1hr
Block 38 - Phencyclidine - (DRE manual session XVI)	2hrs
Block 39 - Narcotic Analgesics (DRE manual session XVII, including examination of injection marks)	2hrs, 30min
DAY EIGHT	
Quiz #3	30min
Block 40 - Inhalants (DRE manual session XIX)	1hr, 30min
Block 41 - Practice: Test Interpretation (DRE manual session XVIII)	1hr
Block 42 - Cannabis (DRE manual session XXI)	2hrs
Block 43 - Resume Preparation and Maintenance (DRE manual session XXIII)	1hr
Block 44 - Practice: Vital Signs (DRE session XX)	30min
Block 45 - Alcohol Correlation Study #3 (DRE manual session XII)	1hr, 30min
DAY NINE	
Quiz #4	30min
Block 46 - Overview of Signs and Symptoms (DRE manual session XXII)	1hr
Block 47 - Drug Combinations (DRE manual session XXIV)	2hrs
Block 48 - Practice Session: Eye Examinations (Note: Students practice the pupil size examinations in this segment. There is no standard lesson plan for this segment.)	1hr

DAY NINE (cont)	
Block 49 - Practice: Test Interpretation (DRE manual session XXV)	1hr
Block 50 - Practice: Test Administration (DRE manual session XXVII)	30min
Block 51 - Review of the DRE School	2hrs
Quiz #5 is also incorporated into this session.	
DAY TEN	·
Block 52 - DRE School Final Examination (DRE manual session XXX)	1hr
Block 53 - Preparing the Narrative Report (DRE manual session XXVI)	1hr
Block 54 - Case Preparation and Testimony (DRE manual session XXVIII)	1hr
Block 55 - Classifying a Suspect (Role Plays) (DRE manual session XXIX)	3hrs
Block 56 - Transition to Certification Phase of Training (DRE manual session XXX)	1hr
Block 57 - Graduation - Presentation of Certificates and Achievement Awards (Note: Course critiques are finished during this segment.)	1hr

ALTERNATE SCHEDULE #3 ACCELERATED DRE SCHOOL

		W	eek One	
Day	Time	Manual	Session/Segment	Title
Monday	(1) 1000 to 1200	SFST DRE	Session I Session I	Introduction & Overview (SFST Script and Matrix Handouts); student/instructor introductions
	1200 to 1300			SFST & DRE Pre-tests
	(2) 1300 to 1400	Pre-School	Session I	Introduction
	1400 to 1500			Lunch Break
	(3) 1500 to 1545	SFST	Session II	Detection and Deterrence
	(4) 1545 to 1630	SFST	Session III	The Legal Environment
	(5) 1630 to 1730	SFST	Session IV	Overview of Detection, Notetaking & Testimony
	(6) 1730 to 1815	SFST	Session V	Phase One: Vehicle in Motion & Explanation of Divided Attention Impairment
	(7) 1815 to 1900	SFST	Session VI	Phase Two: Personal Contact
Tuesday	(8) 1200 to 1230	SFST	Session VII	Phase Three: Pre-Arrest Screening (modified PBT Session)
• .	(9) 1230 to 1330	SFST	Session VIII/A, B	Concepts and Principles of the SFST (development and types of nystagmus)
	(10) 1330 to 1400	Pre-School	Session IV/A & B, 1, 2, & 3	Eye Exams (Purpose of Eye examinations, procedures and clues for HGN, VGN and LOC)
	(11) 1400 to 1500	Pre-School	Session III/A & B	Romberg & Walk and Turn
	(12) 1500 to 1600	Pre-School	Session III/C&D	One Leg Stand & Finger to Nose
	1600 to 1700			Lunch Break
	(13) 1700 to 1800	SFST	Session IX	SFST Test Battery Demonstra- tions (includes Romberg, Finger to Nose in DRE order)
	(14) 1800 to 1900	SFST	Session X	SFST "Dry Run" Practice (includes Romberg, Finger to Nose, in DRE order)
	(15) 1900 to 2100	SFST Pre-School	Session IX Session V	Alcohol Correlation Study #1 - coordinator; wrap-up; bartender; log; vitals

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Wednesday	(16) 1000 to 1200	Pre-School	Session VIII	Alcohol as a Drug (Magic Mountain Video alcohol driving study)
	(17) 1200 to 1300	Pre-School	Session VII	Overview of Signs and Symptoms (distribution of blank drug matrix)
	(18) 1300 to 1400	Pre-School	Session IV/B4, 5	Eye Exams (pupil size & reaction to light)
	1400 to 1500			Lunch Break
	(19) 1500 to 1600	DRE	Session II	Drugs in Society and Motor Vehicle Operation
	(20) 1600 to 1800	DRE	Session III	Development and Effectiveness
	(21) 1800 to 1900			SFST Review Session
Thursday	(22) 1000 to 1030	SFST	Session X	Final Examination
	(23) 1030 to 1100	DRE Pre-School	Session XI Session IV	Eye Exams: Practice Session
	(24) 1100 to 1300	Pre-School DRE	Session VI Session VII	Examination of Vital Signs
	1300 to 1400			Vital Signs: Practice
	1400 to 1500			Lunch Break
	(25) 1500 to 1600	Pre-School DRE	Session II Session IV	Overview: Drug Evaluation and Classification Process (LETN & Chevron tapes)
	(26) 1600 to 1800	DRE	Session VIII	Demonstrations of the Evaluation Sequence
	(27) 1800 to 1900			Pre-School Review Session
Friday	(28) 1200 to 1230	Pre-School	Session X	Final Examination
	(29) 1230 to 1530	DRE	Session VI	Physiology and Drugs: An Overview
	1530 to 1630			Lunch Break
	1630 to 1730			Physiology and Drugs: Physiological Pursuit
	(30) 1730 to 1800	SFST	Session XIII	Report Writing
	1800 to 1900			SFST Practice
	(31) 1900 to 2100	Pre-School SFST	Session V Session XIV	Alcohol Correlation Study #2 & SFST Proficiency Test - coordinator; wrap-up; log; vitals; bartender

Week Two				
Day	Time	Manual	Session/Segment	Title
Monday	1000 to 1030			DRE Quiz #1
	(32) 1030 to 1230	DRE	Session XIII	Physician's Desk Reference & Additional Resources
	(33) 1230 to 1330	non- manual session		Methods of Administration & Elimination
	(34) 1330 to 1400	DRE	Session IX	CNS Depressants
	1400 to 1500			Lunch Break
	1500 to 1630	DRE	Session IX	continued
	(35) 1630 to 1900	DRE	Session X	CNS Stimulants
Tuesday	1000 to 1030			DRE Quiz #2
	1030 to 1130	DRE	Session X/E	continued
	(36) 1130 to 1230	DRE	Session XIV	Hallucinogens
	1230 to 1300	DRE	Session XIV	continued
	(37) 1300 to 1400	DRE	Session XV	Practice: Test Interpretation (includes Clinton Williams evaluation)
	1400 to 1500			Lunch Break
• .	(38) 1500 to 1600	DRE	Session XVI	Phencyclidine
	1600 to 1700	DRE	Session XVI/E	continued
	(39) 1700 to 1900	DRE	Session XVII/ includes E	Narcotic Analgesics
Wednesday	1200 to 1230			DRE Quiz #3
	1230 to 1330	DRE	Session XVII	Injection Marks Examination
	(40) 1330 to 1430	DRE	Session XIX	Inhalants
	(41) 1430 to 1530	DRE	Session XVIII	Practice: Test Interpretation
	(42) 1530 to 1700	DRE	Session XXII	Cannabis
	1700 to 1800			Lunch Break
	(43) 1800 to 1900	DRE	Session XXIII	Resume Preparation & Maintenance
	(44) 1900 to 1930	DRE	Session XX	Practice: Vital Signs
	(45) 1930 to 2100	DRE	Session XII	Alcohol Correlation Study #3 - coordinator; wrap-up; vitals; bartender; log

Thursday	1000 to 1030			DRE Quiz #4
	(46) 1030 to 1130	DRE	Session XXII	Overview of Signs & Symptoms
	(47) 1130 to 1330	DRE	Session XXIV	Drug Combinations
	(48) 1330 to 1430	non- manual session		Practice: Eye Exams
	1430 to 1530			Lunch Break
	(49) 1530 to 1630	DRE	Session XXV	Practice: Test Interpretation
	(50) 1630 to 1700	DRE	Session XXVII	Practice: Test Administration
	(51) 1700 to 1900			DRE Full Course Review "Your Brain on DRE"
				DRE Quiz #5
Friday	(52) 1000 to 1100			Final Examination: DRE School
	(53) 1100 to 1200	DRE	Session XXVI	Preparing the Narrative Report
	(54) 1200 to 1300	DRE	Session XXVIII	Case Preparation & Testimony
	1300 to 1400			Lunch Break
	(55) 1400 to 1700	DRE	Session XXIX	Classifying a Suspect: Role Plays - coordinator
	(56) 1700 to 1800	DRE	Session XXX	Transition to the Certification Phase of Training
	(57) 1800 to 1900			Graduation: Presentation of Certificates and Achievement Awards

C. Overview of the Curriculum Package.

In addition to this Administrator's Guide, the curriculum package for the classroom training program in drug evaluation and classification consists of the following documents and materials:

- o Instructor's Lesson Plans Manual
- o Audio-Visual Aids
- o Student's Manual
- o Set of Drug Evaluation Exemplars

1. Instructor's Lesson Plans Manual

The Instructor's Lesson Plans Manual is a complete and detailed blueprint of what the course covers and of how it is to be taught. It is organized into thirty-two modules, with each module corresponding to one of the training sessions.

Each module consists of a cover page, an outline page, the lesson plans themselves, and master (paper) copies of visual aids referenced in the lesson plans.

The cover page presents the module's (or session's) title and the estimated instructional time required to complete the module.

The outline page lists the specific performance objectives of the module, i.e., the capabilities that the participants will achieve once they have successfully completed the module. The outline page also lists the module's major content segments and the major types of learning activities that are employed during the module.

The lesson plans themselves are arranged in a standard, side-by-side content/instructional notes format. The "content" (left-side) of each page outlines <u>what</u> is to be taught. This content includes:

- o facts
- o concepts
- o procedural steps
- o rules and regulations
- o etc.

The "Instructional Notes" (right-side) portion of each page specifies how the content is to be taught. That is, it defines how the instructor is to present the material and involve the students in the presentation and ensure that they understand and assimilate the material. Typical entries under the "Instructional Notes" column include:

- o the approximate amount of time to be devoted to each major content segment
- o indications of what visual aids are to be used and when they are to be used
- o questions to be posed to students to involve them actively in the presentation
- o indications of points requiring special emphasis
- o guidelines for conducting particular demonstrations to clarify how drug examinations are to be performed
- o specifications of group exercises and other methods of involving students more actively in the lesson

The Instructor's Lesson Plans Manual serves, first, as a means of preparing the instructor to teach the course. He or she should review the entire set of lesson plans and become familiar with the content and develop a clear understanding of how the course "fits together". He or she is also expected to become thoroughly familiar with each module that he or she is assigned to teach, to prepare acetate copies of the visual aids, to assemble all "props" and other instructional equipment referenced in the lesson plans, and to augment the "instructional notes" as necessary to ensure that his or her own teaching style is applied to the content.

<u>Subsequently</u>, the Instructor's Lesson Plans Manual serves as an in-class reference document for the instructor, to help him or her maintain the sequence and pace of presentations and other learning activities.

It is worth emphasizing that the Instructor's Lesson Plans Manual does <u>not</u> contain the text of a speech. Although its outlines of content information are fairly well detailed and comprehensive, those outlines are <u>not</u> to be read verbatim to the participants. This training program is intended to be a dynamic, highly interactive learning experience in which the students are active participants. It should not be permitted to degenerate into a series of mere lectures.

2. Audio-Visual Aids

Five types of audio-visuals are used in this course:

- o wall charts
- o chalkboard/flip-chart presentations
- o "visuals" (overhead transparencies)
- o 35mm photographic slides
- o video tapes

The wall charts are permanently-displayed items. They consist of sketches with brief captions, intended to depict major themes and segments of the training. The wall charts should be handmade, using colored marker pens, on flip chart sheets. The sketches and text must be large enough so that they may be viewed from any seat in the classroom.

Standard-sized paper copies of the suggested wall charts are included in the Instructor's Lesson Plans Manual. The copies may be photocopied onto acetate, to produce overhead transparencies. The transparencies, in turn, can be projected onto flip chart sheets and traced with colored markers, to produce the wall charts themselves.

Wall charts should be placed high on the far left and right sides of the classroom's front wall, or on the side walls, where they will be visible without distracting from the screen or chalkboard.

The chalkboard/flip chart presentations, as recommended in the lesson plans, are self-explanatory.

The "visuals" or overhead transparencies are simple displays of graphic and/or narrative material that emphasize key points and support the instructor's presentation. Paper copies of those "visuals" are found in various modules of the Instructor's Lesson Plans Manual. Those paper copies must be photocopied onto acetate to produce the overhead transparencies. Each "visual" is numbered to indicate the session to which it belongs and its sequence within that session. For example, Visual VII-3 would be the third overhead transparency used in Session VII.

35mm photographic slides are available of all the overhead transparencies.

The video tapes consist of a number of segments that demonstrate the drug evaluation and classification procedures, and that exhibit the kinds of evidence associated with various categories of drugs. Some of these segments feature persons who are actually under the influence of various drugs and who have been arrested for offenses relating to their drug impairment.

3. Student's Manual

The Student's Manual is the basic textbook and study source for the course. It provides a session-by-session summary of the subject matter, and a list of study topics to help the students assimilate the material.

<u>During</u> the course, the Student's Manual will be primarily useful for <u>previewing</u> the sessions, and for studying the subject matter in preparation for the final knowledge and proficiency examinations. <u>After</u> the classroom training is completed, the student will find that the manual is a useful reference document, especially during the Certification Phase of training.

Students are expected to be familiar with all of the contents of their Student Manual. Instructors must encourage the students to study the manual carefully as they progress through the school. Note: Students are expected to be able to answer the "topics for study" review questions that appear at the end of various sections of their Student Manual.

4. Set of Drug Evaluation Exemplars

The exemplars are the documented results of simulated drug evaluation and classification examinations. A standardized reporting form is used for the exemplars. This is the same form that the students use as a test recording instrument when they practice administering and documenting the drug evaluation and classification examination.

The exemplars support learning activities that take place during eleven sessions:

o Sessions IX, X, XIV, XVI, XVII, XIX, and XXI cover the seven individual drug categories. Several exemplars have been prepared for each session, to illustrate the kinds of clues that can be expected when the examination is conducted for a person under the influence of that category. For example, the exemplars designed for Session IX illustrate the results of typical examinations of suspects under the influence of CNS depressants.

These exemplars will be found in the Instructor's and Student's Manual.

- o Session XV, XVIII and XXV are "Test Interpretation Practice" sessions. Students work in small groups, reviewing exemplars and determining, from the documented "evidence" they contain, what category or categories of drugs are present in each case. These exemplars also will be found in the Student's Manual.
- Session XXIX is the "role play" practice session. Instructors serve as O "test subjects". Students work in small groups, administering the entire drug evaluation and classification examination to each instructor. Each instructor uses an exemplar to inform the students as to what data they should record at each stage of the examination. For example, as part of the examination, the students will actually measure an instructor's blood pressure. The instructor will observe the students' technique and offer constructive criticism. The instructor will inquire as to the pressure readings that the students obtain. But, the instructor will tell the students to record the blood pressure readings documented on his or her assigned exemplar. Subsequently, the students must review their completed exemplars and determine what category or categories of drugs the instructor was "simulating". These exemplars are found at the end of the lesson plans for Session XXIX.

D. General Administrative Requirements

1. Facility Requirements

Several types of facilities are needed to support this training. First, a standard classroom is required. This should provide comfortable seating and adequate desk/table space for each student, and should be equipped with a large screen, overhead and 35mm slide projectors, chalkboards and/or flip-charts and video tape players and monitors. All visuals should be readily and fully visible from all seating locations. The classroom should also provide adequate unobstructed space to allow the instructors to demonstrate examination procedures. A "U"-shaped seating arrangement is preferable for the classroom.

A large, open area also is needed to support the hands-on practice sessions. A gymnasium or similar facility will serve this need very well. Ideally, it should be possible to control the lighting in this practice facility to the point of total darkness, to demonstrate and practice key elements of the drug evaluation and classification procedures that take place in a darkroom.

A separate room must be available, ideally adjacent to the gymnasium or practice facility. This room will serve as the "staging area" for the volunteer drinkers who will participate in the alcohol workshop (Session XII).

Another separate room must be provided to serve as the instructors' "office", i.e., the place where they can prepare for their teaching assignments, store materials, etc.

2. Special Instructional Equipment and Personnel.

For the alcohol workshop, volunteer drinkers must be available. The volunteer drinkers cannot be members of the class. There should be one volunteer for every three or four students. For example, if there are 25 students in the class, there should be 7-9 volunteer drinkers. Sufficient alcohol, mixers, cups, napkins, ice, etc. must be provided. Adequate breath testing devices must be available to provide for monitoring volunteers' blood alcohol concentrations. At least three people must be assigned to monitor and escort the volunteers; ideally, each volunteer should have his or her own monitor.

Note: Every volunteer must read and sign the "Statement of Informed Consent" prior to receiving any alcohol. Any person who refuses to sign the Statement cannot serve as a volunteer drinker.

For the hands-on practice sessions involving eye examinations, at least one pupillometer and one onset angle template should be provided for every two students. Ideally, each student should have his or her own pupillometer and template. The pupillometer should be capable of measuring pupil diameters across the range from 1.0 mm to 9.0 mm, in one-half millimeter increments. The template should display angles between 30 and 50 degrees, in 5 degree increments.

For the hands-on practice sessions involving vital signs examinations, a sphygmomanometer and stethoscope must be provided for every three students. Ideally, each student should have his or her own. Also, it is desirable that several <u>training</u> stethoscopes be available. These are stethoscopes that have two sets of earpieces, and allow an instructor to monitor exactly what the student is hearing.

Each student should be provided with a penlight suitable for conducting the various eye examinations.

At the beginning of DRE training, it is essential that every student have his or her own full complement of DRE equipment. In addition, every student must have access to a PDR, and ideally should own a PDR.

3. Instructor Qualifications.

The principal instructors for this course must be IACP-certified Drug Recognition Expert Instructors. That means that they (1) hold currently-valid certificates as DREs; (2) have completed the NHTSA DRE Instructor Training Course; and, (3) have completed the required delivery of both classroom and certification training, under the supervision of teacher-trainers. Only a certified DRE instructor can credibly teach:

- o Session IV (Overview of Drug Evaluation and Classification Procedures)
- o Session V (Eye Examinations)
- o Session VIII (Demonstrations of the Evaluation Sequence)
- o The segment entitled "Expected Results of the Evaluation" in Sessions IX, X, XIV, XVI, XVII, XIX XXI and XXIV (The sessions covering individual drug categories and combinations of categories)
- o The hands-on practice sessions (Sessions XI, XX, XVIII and XXIX)
- o The Test Interpretation Practice Sessions (Sessions XV, XVII and XXV)
- o Session XXVI (Narrative Drug Report)
- o Session XXIII (Resume Preparation and Maintenance)

The above-listed sessions and segments constitute approximately 75% of the course.

A qualified DRE <u>could</u> instruct the remaining 25% of the course, as well. However, some agencies may wish to enlist instructors with special credentials for certain blocks of instruction. For example, a physician would be well qualified to teach Session VII (Examination of Vital Signs), and a prosecutor might be a good choice as the instructor for Session XXVIII (Case Preparation and Testimony), and for Session XXVI (Preparing the Narrative Report).

In addition to their occupational competencies, all instructors must be qualified teachers. They need to understand, and be able to apply, fundamental principles of instruction. Perhaps most importantly, they need to be competent <u>coaches</u>. Much of this classroom training is devoted to hands-on practice. The quality of coaching will have a major impact on the success of those practice sessions. It is <u>highly</u> recommended that every instructor be a graduate of the NHTSA DRE Instructor Training School.

For the hands-on practice sessions, there should be at least one instructor for every three students, to permit adequate monitoring and coaching.

4. Class Size Considerations.

The recommended maximum class size for this course is 25 students. Larger classes make it difficult to devote sufficient attention to each student to ensure that he or she develops examination skills to a level sufficient to progress to the Certification Phase. The preferred class size is 15-20 students.

E. Course Planning and Preparation Requirements

The fundamental preparatory step for any law enforcement agency desiring this training is to ensure that the agency and its community or state satisfy the prerequisites outlined in Section B, part 1 of this Administrator's Guide.

The next step is to select a cadre of <u>appropriate</u> candidate DREs. Make sure that each candidate satisfies the student prerequisites outlined in Section B.

The third step is to provide <u>preliminary</u> training to the candidate DREs. The National Highway Traffic Safety Administration (NHTSA) has developed a curriculum to support preliminary training for potential DREs. This training enables the candidates to become familiar with, and to start to develop skills in, the vital signs examinations and other elements of the drug evaluation and classification procedures.

The next step will be to schedule the class. States with well-established DRE programs, including a cadre of experienced DRE instructors, are expected to plan and manage their own DRE Schools. However, they can receive the services of additional (in-State and out-of-State) instructors, at NHTSA's expense. And of course, NHTSA supplies Student Manuals and other standard instructional materials at no charge. For States whose DRE programs are new or developing, NHTSA assists with the planning and management of the Schools, and supplies most or all instructors.

In general, this classroom training course is conducted at facilities operated by the delivery agency or at other suitable locations. Departments are responsible for all costs associated with transporting their personnel to and from the training site, and for their lodging and subsistence during the training.

F. Examinations of Students' Knowledge and Proficiency

It is very important to test the students' knowledge and skill development. Testing in this course is conducted for two principal reasons: (1) to assess students' progress, and identify deficiencies that need correction; and, (2) as a learning activity for the students. Knowledge testing starts in the very first session of the course, when a PRE-Test is given. After the students have finished the PRE-Test, you will give them a new, blank copy of the test, so that they can use it as a study guide throughout the course. Five formal quizzes also will be given. The first of these is given at the start of the third day of the school. The second quiz is given at the start of the fifth day, and the third quiz at the start of the sixth day. The fourth quiz is given at the end of the sixth day. The fifth quiz is given during the Optional Review Session that occurs during the evening of the sixth day. In addition, a self-study quiz is provided in the Student's Manual.

The most important knowledge test, of course, is the Final Examination. It is given on the afternoon of the final day of the School. The student must achieve a grade of at least 80% in order to progress to certification training. If a student fails the examination, the National minimum Standards permit one additional attempt. The additional attempt must be based on an examination approved for that purpose by NHTSA and IACP, and cannot occur earlier than two weeks, nor later than four weeks, following completion of the DRE School.

A skill examination also occurs during the next-to-last session of the DRE School. That is the session in which the students will examine instructors who are "playing the roles" of drug-impaired person. A Proficiency Examination Checklist (found in Session XXX of this Manual) is used to evaluate the students' performance.

G. Follow-Up Requirements

Upon completion of the classroom training, students will commence the Certification Phase, i.e., the application of drug evaluation and classification procedures in an actual enforcement context. During certification training, the students are supervised by certified DRE instructors. Under the national minimum standards for certification established by the International Association of Chiefs of Police (IACP), each student must participate in conducting at least 12 drug examinations, at least six of which he or she must personally administer.

The student must also identify at least three of the seven drug categories in his or her examinations. And, toxicologic specimens must be submitted from at least nine of the examined subjects, and analysis of those specimens must corroborate the student's opinion for at least 75% of the specimens submitted. Most importantly, the numbers and percentages cited here are minimum requirements: no student can be certified as a DRE until two instructors attest that he or she qualifies for certification.

NHTSA requires that a formal assessment of the drug evaluation and classification program be conducted by each agency that receives this training. At a minimum, it is expected that departments will maintain records, for a period of at least 6 months following completion of the Certification Phase training, on each DRE's on the job performance, and that copies of those records will be submitted to NHTSA. The records should include the Standard Drug Evaluation and Classification form (or its equivalent); the arrest report; the Narrative Report; and, the results of the laboratory analysis of a blood or urine sample (if available) on each suspect examined by a DRE.

The training delivery agency will compile the information needed to support an assessment of the classroom training each time it is conducted. This assessment will be based primarily on the (anonymous) Student's Critique Form, which appears in Session XXX of the Instructor's Lesson Plans Manual. Guidelines for preparing a post-course evaluation report based on the Student's Critique Form are covered in Section H.

H. Guidelines for Preparing Post-Course Evaluation

A standard NHTSA/TSI participant's critique form is provided to document participant's initial ratings of course content and activities. The form is divided into eight parts:

- A. Workshop/Seminar Objectives
- B. Course Activities
- C. Course Design
- D. Topic Deletions
- E. Topic Additions
- F. Ability to Identify Drug Categories
- G. Overall Quality of the Course
- H. Quality of Instruction
- I. Final Comments or Suggestions

The following instructions are provided to guide review, analysis and interpretation of participant's comments:

Section A - Workshop/Seminar Objectives

Determine raw tabulation and percentages for each objective:

o If the "no"/"not sure" responses total 20% or more, some explanation should be provided. Assess the problem and explain or recommend changes as appropriate.

Section B - Course Activities

The rating choices are as follows:

- 1. Very Important
- 2. Somewhat Important
- 3. Un-Important
- 4. Not Sure

Analysis Procedures

- Step 1: Tabulate total number of responses in each category for each activity.
- Step 2: The following values should be applied:
 - o +2 for each "very important"
 - o 0 for each "somewhat important"
 - o -2 for each "un-important"
 - o -1 for each "not sure"
- Step 3: Determine total number of points for each activity.
- Step 4: Divide the totals by twice the number of votes (N).
- Step 5: The result is the final rating.

Any rating of +.5 or higher indicated the participant's consensus was that the activity (segment) was "very important".

If the rating is below +.2, some explanation should be provided...assess the reason(s) and explain or recommend changes as appropriate.

If the rating is below 0 there is a serious problem...assess the problem(s) and explain or recommend changes as appropriate.

Section C - Course Design

Determine raw tabulation and percentage for each statement.

Some comment or explanation should be provided if the inappropriate ("agree"/"disagree") or "not sure" responses exceed 20%.

Section D & E - Topic Deletion/Additions

Prepare a summary of responses for each section. Comment as appropriate.

Section F - Ability to Identify Drug Categories

Total the numerical ratings, and divide by the number of responding participants. That gives the average rating for the section, on the scale from 1 ("very confident") to 3 ("not confident"). Comment as appropriate.

Section G - Overall Quality of the Seminar

Total the numerical ratings, and divide by the number of responding participants. That gives the average rating for the seminar, on the scale from 1 ("poor") to 5 ("excellent"). Comment as appropriate.

Section H - Quality of Instruction

For each instructor, tabulate his or her numerical ratings, and divide by the number of responding participants. Comment as appropriate.

Section I - Final Comments

Prepare a summary of responses for each section. Comment as appropriate.

<u>NOTE</u>: A copy of the completed post course evaluation report should be forwarded to the appropriate State Highway Safety Office and/or NHTSA Region Office as they are completed. These reports will be used to assist in determining what revisions are needed to the course curriculum in the future when periodic course reviews are conducted by the NHTSA.

I. Requests for Information, Assistance or Materials

Departments interested in this program should contact their state's Office of Highway Safety. Formal requests for this training should come from the State Highway Safety Office, and should be directed to the cognizant NHTSA Regional Office.

One Hour and Fifty Minutes

SESSION I INTRODUCTION AND OVERVIEW

Upon successfully completing this session, the participant will be able to:

- o State the goals and objectives of the course.
- o Outline the major course content.
- o Outline the schedule of major course activities.
- o Outline the contents and arrangement of the student manual.

During this session, the participant will demonstrate his or her current knowledge of basic concepts and terminology relevant to the Drug Evaluation and Classification Process.

Content Segments		Learning Activities		
A.	Welcoming Remarks and Goal	0	Instructor Led Presentations	
B.	Participant Introductions	0	Participant Led Presentations	
C.	Objectives		Knowledge Examination	
D.	Overview of Content and Schedule	0	Reading Assignments	
E.	Overview of Student Manual			
F.	Administrative Matters			
G.	Glossary of Terms			

000043
Aides
I-0 (Session
Objectives)
10 Minutes
SF
I-1 (Goal)



(Tennessee)



(Maryland)

Lesson Plan

Instructor Notes

INTRODUCTION AND OVERVIEW

A. Welcoming Remarks and Goal

1. Welcome to the seven day DRE School.

The goal of this school is simple:

To help you prevent crashes, deaths and injuries caused by drug impaired drivers.

University of Tennessee study (1988)

> 40% of drivers treated at Trauma Center for crash injuries had drugs other than alcohol in them.

Maryland Shock Trauma Center study (1985-1986)

> 32% of drivers treated at the Shock Trauma Center had used marijuana prior to their crashes.

Total Lesson Time: Approximately 110 Minutes

Briefly review the content, objectives and activities of this session.

Course title on wall chart.

Brief welcoming remarks by the lead-off instructor (not longer than one minute).

The Tennessee study was conducted by Kirby, Jackie M. (RN, MSN) and Maull, Kimball I. (MD), Division of Trauma/ Critical Care, Department of Surgery, University of Tennessee Medical Center, Knoxville, Tennessee.

Emphasize that these studies clearly show that drug impaired driving is a major problem in this country.

Instructor note: Remind students that all studies published are subject to interpretation.

Aides	Lesson Plan	Instructor Notes
, .		For more information contact NHTSA, The National Traffic Law Center, or the IACP DEC Technical Advisory Panel.
	3. We can do something to remove drugged drivers from our roads.	
	a. The DRE Program is based on solid medical and scientific facts.	,
	b. The validity of the DRE Program has been tested in carefully controlled research in both the laboratory and the field.	Point out that the students will hear more about this research later today.
·	4. By enrolling in DRE training, you have become part of an elite International Program.	
	 a. Drug Recognition Experts form one of the tightest knit fraternities in law enforcement. 	·
	b. DREs from many agencies and from many parts of the country work closely together to share information and other resources, and to maintain the highest standards of quality.	Mention the various agencies represented among the instructors and the students in this school.
	c. Each of you was selected to receive this training because you were recognized by your department as a skilled and dedicated law enforcement professional.	

Aides	Lesson Plan	Instructor Notes
	d. Your instructors welcome you to this school. We're proud to have you here, and we're sure that you are proud to be here.	
	B. Introductions	
25 Minutes		
	Introduction of representatives of host agencies and other dignitaries.	The introductions of dignitaries, and their welcoming remarks, must be kept brief: no more than 10 minutes can be devoted to this.
	2. Introduction of faculty.	The lead-off instructor should mention the names and agency affiliations of all other instructors, asking each to stand as their name is called.
	3. Students' introductions.	Whenever possible, instructor should consider using creative and innovative icebreaking techniques. At a minimum, instruct each student to stand and give their name, agency affiliation and experience.
	C. Objectives	
10 Minutes		
I-3A (First Three Objectives)	 If you successfully complete this School, you will be able to: Describe the involvement of drugs in impaired driving incidents. Name the seven categories of drugs and recognize their effects. 	-

000046
Aides
I-3B (Next Two Objectives) I-3C (Last Three Objectives)
25 Minutes
A A A A A A A A A A A A A A A A A A A

Lesson Plan

Instructor Notes

- c. Describe and properly conduct the drug evaluation.
- d. Document the results of the drug evaluation.
- e. Properly interpret the results of the evaluation.
- f. Write narrative Drug Influence Reports.
- g. Testify clearly and convincingly in drug evaluation cases.
- h. Maintain an up to date resume.
- 2. Every DRE needs to be able to do these eight things.
- 3. Before you can be certified as a DRE, you will have to demonstrate that you can do each of these things.
- D. Overview of Content and Schedule
 - Major content topics
 - a. Drugs in society and in vehicle operation.
 - b. Development and effectiveness of the Drug Evaluation and Classification Program.
 - c. Overview of Drug Recognition Expert Procedures.

Solicit students' questions about the objectives.

Refer to wall charts in previewing the content topics.

Briefly overview the contents covered under each major topic.

Aides		Lesson Plan	Instructor Notes
· ·	d.	Eye Examinations (a major component of the Drug Evaluation Procedures).	
	e.	Physiology and Drugs.	
	f.	Vital signs examinations (a major component of the Drug Evaluation Procedures).	·
	g.	The seven categories of drugs.	,
	h.	The Physicians's Desk Reference (PDR) and other reference sources.	
	i.	Interviewing suspects (a major component of the Drug Evaluation Procedures).	
	j.	Resume preparation and maintenance.	
	k.	Case preparation and testimony.	
	1.	Classifying a suspect (interpreting and documenting the results of an examination)	Solicit students' questions concerning the content topics.
	2. Ha	ands-on practice sessions.	Emphasize that hands on practice is the principal learning activity of this course.
	a.	Eye Examinations practice (Nystagmus, Lack of Convergence, pupil size and reaction to light)	Refer to wallchart outlining practice sessions.

Aides	Lesson Plan	Instructor Notes
	b. Alcohol workshop (psychophysical testing practice)	Point out that volunteer drinkers from outside the class will be recruited for this session.
	c. Practicing interpretation of the examination results.	Point out that several sessions will be devoted to this. In each, students will review drug evaluation reports and identify the probable category or combinations of categories of drugs involved.
	d. Vital signs examinations practice (pulse, blood pressure)	
	e. Practicing administration of the DRE examination.	Point out that several sessions will be devoted to this. In each, students will practice administering the drug examinations to each other. No hands on practice with actual drugged subjects is included in the classroom portion of DRE training.
	f. Simulated drug impaired subjects examinations.	Point out that students will work in teams to conduct and document examinations of instructors who will be simulating the indicators of drug-impaired subjects.
		Solicit students' questions concerning the hands-on practice sessions.
	3. Course schedule.	Refer students to the schedule shown in their manuals.
		Briefly overview the schedule of sessions.
		Solicit students' questions concerning the schedule.

Aides	Lesson Plan	Instructor Notes
	E. Overview of Student Manual	
25 Minutes	·	
	Student Manual is the basic reference document for this course.	Make sure each student has a copy of the student manual.
	a. The Manual Contains a summary of presentations made by instructors throughout the classroom training.	
ı	b. The Manual includes a set of "class notes" for every session in the course.	Point out that the Student Manual has a separate chapter, or section, for each session of the course.
·		Instruct students to open their Manuals to Session I, and briefly review the content of that section of the Manual, to illustrate how the document is organized.
	2. Students are expected to use the Manual to review the material covered in class.	
	3. The Manual should also be used to <u>preview</u> the class sessions.	Encourage students to read the appropriate Student Manual Sections prior to each day's classes.
	4. By taking good notes, and by studying the Manual carefully, students should have no trouble in passing the course.	-
	5. At the conclusion of the classroom training, the student	

Remind students that there will be numerous quizzes during the class.

must pass the written test with a score of 80% or better in order

to progress to the certification

phase.

Aides	Lesson Plan	Instructor Notes
	F. Administrative Matters	
15 Minutes		
	Logistics. (Completion of registration forms, travel vouchers, etc.)	
	2. Mandatory attendance at all sessions of this school.	Emphasize that, if a student misses any portion of this school, he or she must make up
	3. Facilities. (Locations of restrooms, lunchrooms, etc.)	the deficiency via after hours tutoring before beginning certification training.
	4. Pre-test	Hand out pre-tests. Emphasize that the pre-test scores do not affect passage of this course, nor will the pre-test be a part of the student's permanent record. Allow 10 minutes for students to complete, then collect the pre-tests. Point out to the students that they will find a "clean" copy of the pre-test at the end of Section I of their Student's Manual. Inform students to use the pre-test as a study guide while they progress through the course.
,		

DRE 7-Day School

Session I

Introduction and Overview



Introduction and Overview

Upon successfully completing this session, the participant will be able to:

- · State the goals and objectives of the course
- · Outline the major course content
- Outline the schedule of major course activities
- Outline the contents and arrangement of the student manual

Drug Realization & Classification Training

140

Ultimate Goal of the Program:

To help you prevent crashes, deaths and injuries caused by drug-impaired drivers





Drug Evaluation & Classification Training

1.

University of Tennessee Study (1988):

40% of drivers receiving emergency treatment had used drugs prior to their crashes



Drug Evaluation & Classification Training

I-2A

Maryland Shock Trauma Center Study (1985-1986):

32% of drivers treated at the shock trauma center had used marijuana prior to their crashes

Drug Evaluation & Classification Training

1-28

Classroom Training Objectives

You will become better able to:

- Describe the involvement of drugs in impaired driving incidents
- 2. Name the seven drug categories and recognize their effects
- 3. Describe and properly conduct the drug evaluation

Drug Evaluation & Classification Trainin

I-3A

Classroom Training Objectives (continued)

- 4. Document the results of the drug evaluation and classification process
- 5. Properly interpret the results of the evaluation

Drug Evaluation & Classification Training

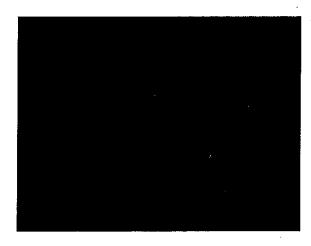
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Classroom Training Objectives (continued)

- 6. Prepare a narrative drug influence report
- 7. Discuss appropriate procedures for testifying in typical drug evaluation/classification cases
- 8. Maintain a relevant and up-to-date resume

Drug Evaluation & Classification Training

I-3C



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SCORE SHEET

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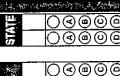
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FINAL WRITTEN

EXAMINATION











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DRUG EVALUATION AND

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DRUG EVALUATION AND CLASSIFICATION PROGRAM

GLOSSARY OF TERMS

ADDICTION

Habitual, psychological, and physiological dependence on a substance beyond one's voluntary control.

ADDITIVE EFFECT

One mechanism of polydrug interaction. For a particular indicator of impairment, two drugs produce an additive effect if they both affect the indicator in the same way. For example, cocaine elevates pulse rate and PCP also elevates pulse rate. The combination of cocaine and PCP produces an additive effect on pulse rate.

AFFERENT NERVES

See: "Sensory Nerves."

ALKALOID

A chemical that is found in, and can be physically extracted from, some substance. For example, morphine is a natural alkaloid of opium. It does not require a chemical reaction to produce morphine from opium.

ANALGESIC

A drug that relieves or allays pain.

ANALOG (of a drug)

An analog of a drug is a chemical that is very similar to the drug, both in terms of molecular structure and in terms of psychoactive effects. For example, the drug Ketamine is an analog of PCP.

ANESTHETIC

A drug that produces a general or local insensibility to pain and other sensation.

ANTAGONISTIC EFFECT

One mechanism of polydrug interaction. For a particular indicator of impairment, two drugs produce an antagonistic effect if they affect the indicator in opposite ways. For example, heroin constricts pupils while cocaine dilates pupils. The combination of heroin and cocaine produces an antagonistic effect on pupil size. Depending on how much of each drug was taken, and on when they were taken, the suspect's pupils could be constricted, or dilated, or within the normal range of size.

ARRHYTHMIA

An abnormal heart rhythm.

ARTERY

The strong, elastic blood vessels that carry blood from the heart to the body tissues.

ATAXIA

A blocked ability to coordinate movements. A staggering walk and poor balance may be caused by damage to the brain or spinal cord. This can be the result of trauma, birth defect, infection, tumor, or drug use.

AUTONOMIC NERVE

A motor nerve that carries messages to the muscles and organs that we do not consciously control. There are two kinds of autonomic nerves, the sympathetic nerves and parasympathetic nerves.

AXON

The part of a neuron (nerve cell) that sends out a neurotransmitter.

BAC

(Blood Alcohol Concentration) - The percentage of alcohol in a person's blood.

BrAC

(Breath Alcohol Concentration) - The percentage of alcohol in a person's blood as measured by a breath testing device.

BLOOD PRESSURE

The force exerted by blood on the walls of the arteries. Blood pressure changes continuously, as the heart cycles between contraction and expansion.

BRADYCARDIA

Abnormally slow heart rate; pulse rate below the normal range.

BRADYPNEA

Abnormally slow rate of breathing.

BRUXISM

Grinding the teeth. This behavior is often seen in persons who are under the influence of cocaine or other CNS stimulants.

CANNABIS

- 1. One of the seven drug categories. Cannabis includes marijuana, hashish, hash oil, and marinol.
- 2. Several species of plants from which marijuana and related products are made (e.g., Cannabis Sativa and Cannabis Indicia).

CARBOXY THC

A metabolite of THC (tetrahydrocannabinol).

CHEYNE

Stokes Respiration - Abnormal pattern of breathing. Marked by breathlessness and deep, fast breathing.

CNS (Central Nervous System)

A system within the body consisting of the brain, the brain stem, and the spinal cord.

CNS DEPRESSANTS

One of the seven drug categories. CNS depressants include alcohol, barbiturates, anti-anxiety tranquilizers, and numerous other drugs.

CNS STIMULANTS

One of the seven drug categories. CNS Stimulants include cocaine, the amphetamines, ritalin, preludin, and numerous other drugs.

CONJUNCTIVITIS

An inflammation of the mucous membrane that lines the inner surface of the eyelids caused by infection, allergy, or outside factors. May be bacterial or viral. Persons suffering from conjunctivitis may show symptoms in one eye only. This condition is commonly referred to as "pink eye", a condition that could be mistaken for the bloodshot eyes produced by alcohol or Cannabis.

CONVERGENCE

The "crossing" of the eyes that occurs when a person is able to focus on a stimulus as it is pushed slowly toward the bridge of his or her nose. (See, also, "Lack of Convergence".)

CRACK

A hard chunk form of cocaine that produces a very intense, but relatively short duration "high". (Rock is a different process.)

CYCLIC BEHAVIOR

A manifestation of impairment due to certain drugs, in which the suspect alternates between periods (or cycles) of intense agitation and relative calm. Cyclic behavior, for example, sometimes will be observed in persons under the influence of PCP.

DENDRITE

The part of a neuron (nerve cell) that receives a neurotransmitter.

DIACETYL MORPHINE

The chemical name for Heroin.

DIASTOLIC

The lowest value of blood pressure. The blood pressure reaches its diastolic value when the heart is fully expanded, or relaxed (Diastole).

DIPLOPIA

Double vision.

DISSOCIATIVE ANESTHETIC

A drug that inhibits pain by cutting off (or "disassociating") the brain's perception of the pain. PCP is usually described as a dissociative anesthetic.

DIVIDED ATTENTION

Concentrating on more than one thing at a time. The four psychophysical tests used by DREs require the suspect to divide attention.

DRUG

Any substance, which when taken into the human body, can impair the ability of the person to operate a vehicle safely.

DYSARTHIA

Slurred speech. Difficult, poorly articulated speech.

DYSPNEA et. al.

Shortness of breath.

DYSMETRIA

An abnormal condition that prevents the affected person from properly estimating distances linked to muscular movements.

DYSPHORIA

A disorder of mood. Feelings of depression and anguish.

EFFERENT NERVES

See: "Motor Nerves".

ENDOCRINE SYSTEM

The network of glands that do not have ducts and other structures. They secrete hormones into the blood stream to affect a number of functions in the body.

EXPERT WITNESS

A person skilled in some art, trade, science or profession, having knowledge of matters not within knowledge of persons of average education, learning and experience, may assist a jury in arriving at a verdict by expressing an opinion on a state of facts shown by the evidence and based upon his or her special knowledge. (NOTE: Only the court can determine whether a witness is qualified to testify as an expert.)

FLASHBACK

A vivid recollection of a portion of an hallucinogenic experience. Essentially, it is a very intense daydream. There are three types: (1) emotional -- feelings of panic, fear, etc.; (2) somatic -- altered body sensations, tremors, dizziness, etc.; and (3) perceptual -- distortions of vision, hearing, smell, etc.

GARRULITY

Chatter, rambling or pointless speech. Talkative.

HALLUCINATION

A sensory experience of something that does not exist outside the mind, e.g., seeing, hearing, smelling, or feeling something that isn't really there. Also, having a distorted sensory perception, so that things appear differently than they are.

HALLUCINOGENS

One of the seven drug categories. Hallucinogens include LSD, MDMA, peyote, psilocybin, and numerous other drugs.

HASHISH

A form of cannabis produced by boiling, compressing and drying the leaves of the female marijuana plant. Hashish has a higher concentration of THC (tetrahydrocannabinol) than does the marijuana from which it is produced.

HASH OIL

A liquid extracted from hashish, and containing a relatively high concentration of THC.

HEROIN

A powerful and widely-abused narcotic analgesic that is chemically derived from morphine. The chemical, or generic name of heroin is "diacetyl morphine".

HIPPUS

A rhythmic pulsating of the pupils of the eyes, as they dilate and constrict within fixed limits.

HOMEOSTASIS

The dynamic balance, or steady state, involving levels of salts, water, sugars, and other materials in the body's fluids.

HORIZONTAL GAZE NYSTAGMUS (HGN)

Involuntary jerking of the eyes occurring as the eyes gaze to the side.

HORMONES

Chemicals produced by the body's endocrine system that are carried through the blood stream to the target organ. They exert great influence on the growth and development of the individual, and that aid in the regulation of numerous body processes.

HYDROXY THC

A metabolite of THC (tetrahydrocannabinol).

HYPERFLEXIA

Exaggerated or over extended motions.

HYPERGLYCEMIA

Excess sugar in the blood.

HYPERPNEA

A deep, rapid or labored breathing.

HYPERPYREXIA

Extremely high blood temperature.

HYPERREFLEXIA

A neurological condition marked by increased reflex reactions.

HYPERTENSION

Abnormally high blood pressure. Do not confuse this with hypotension.

HYPOGLYCEMIA

An abnormal decrease of blood sugar levels.

HYPOPNEA

Shallow or slow breathing.

HYPOTENSION

Abnormally low blood pressure. Do not confuse this with hypertension.

HYPOTHERMIA

Decreased body temperature.

ICE

A crystalline form of methamphetamine that produces a very intense and fairly long-lasting "high".

INHALANTS

One of the seven drug categories. The inhalants include volatile solvents (such as glue and gasoline), aerosols (such as hair spray and insecticides) and anesthetic gases (such as nitrous oxide).

INSUFFLATION

See "snorting".

INTEGUMENTARY SYSTEM

The skin and accessory structures, hair and nails. Functions include protection, maintenance of body temperature, excretion of waste, and sensory perceptions.

INTRAOCULAR

"Within the eyeball".

KOROTKOFF SOUNDS

A series of distinct sounds produced by blood passing through an artery, as the external pressure on the artery drops from the systolic value to the diastolic value.

LACK OF CONVERGENCE

The inability of a person's eyes to converge, or "cross" as the person attempts to focus on a stimulus as it is pushed slowly toward the bridge of his or her nose.

MARIJUANA

Common term for the Cannabis Sativa plant. Usually refers to the dried leaves of the plant. This is the most common form of the cannabis category.

MARINOL

A drug containing a synthetic form of THC (tetrahydrocannabinol). Marinol belongs to the cannabis category of drugs, but marinol is not produced from any species of cannabis plant.

METABOLISM

The sum of all chemical processes that take place in the body as they relate to the movements of nutrients in the blood after digestion, resulting in growth, energy, release of wastes, and other body functions. The process by which the body, using oxygen, enzymes and other internal chemicals, breaks down ingested substances such as food and drugs so they may be consumed and eliminated. Metabolism takes place in two phases. The first step is the constructive phase (anabolism) where smaller molecules are converted to larger molecules. The second steps is the destructive phase (catabolism) where large molecules are broken down into smaller molecules.

METABOLITE

A chemical product, formed by the reaction of a drug with oxygen and/or other substances in the body.

MIOSIS

Abnormally constricted pupils.

MOTOR NERVES

Nerves that carry messages away from the brain, to be body's muscles, tissues, and organs. Motor nerves are also known as efferent nerves.

MUSCULAR HYPERTONICITY

Rigid muscle tone.

MYDRIASIS

Abnormally dilated pupils.

NARCOTIC ANALGESICS

One of the seven drug categories. Narcotic analgesics include opium, the natural alkaloids of opium (such as morphine, codeine, and thebaine), the derivatives of opium (such as heroin, dilaudid, metopon, percodan and hycodan), and the synthetic narcotics (such as demerol and numorphan).

NERVE

A cord-like fiber that carries messages either to or from the brain. For drug evaluation and classification purposes, a nerve can be pictured as a series of "wire-like" segments, with small spaces or gaps between the segments.

NEURON

A nerve cell. The basic functional unit of a nerve. It contains a nucleus within a cell body with one or more axons and dendrites.

NEUROTRANSMITTER

Chemicals that pass from the axon of one nerve cell to the dendrite of the next cell, and that carry messages across the gap between the two nerve cells.

NULL EFFECT

One mechanism of polydrug interaction. For a particular indicator of impairment, two drugs produce a null effect if <u>neither</u> of them affects that indicator. For example, PCP does not affect pupil size, and alcohol does not affect pupil size. The combination of PCP and alcohol produces a null effect on pupil size.

NYSTAGMUS

An involuntary jerking of the eyes.

"ON THE NOD"

A state of deep relaxation, induced by impairment due to heroin or other narcotic analgesic. The suspect's eyelids droop, and chin rests on the chest. Suspect may appear to be asleep, but can be easily aroused and will respond to questions.

OVERLAPPING EFFECT

One mechanism of polydrug interaction. For a particular indicator of impairment, two drugs produce an overlapping effect if one of them affects the indicator but the other doesn't. For example, cocaine dilates pupils while alcohol doesn't affect pupil size. The combination of cocaine and alcohol produces an overlapping effect on pupil size: the combination will cause the pupils to dilate.

PALLOR

An abnormal paleness or lack of color in the skin.

PARANOIA

Mental disorder characterized delusions and the projection of personal conflicts, that are ascribed to the supposed hostility of others.

PARAPHERNALIA

Drug paraphernalia are the various kinds of tools and other equipment used to store, transport or ingest a drug. Hypodermic needles, small pipes, bent spoons, etc., are examples of drug paraphernalia. The singular form of the word is "paraphernalium". For example, one hypodermic needle would be called a "drug paraphernalium".

PARASYMPATHETIC NERVE

An autonomic nerve that commands the body to relax and to carry out tranquil activities. The brain uses parasympathetic nerves to send "at ease" commands to the muscles, tissues, and organs.

PARASYMPATHOMIMETIC DRUGS

Drugs that mimic neurotransmitter associated with the parasympathetic nerves. These drugs artificially cause the transmission of messages that produce lower blood pressure, drowsiness, etc.

PDR (Physician's Desk Reference)

A basic reference source for drug recognition technicians. The PDR provides detailed information on the physical appearance and psychoactive effects of all licitly-manufactured drugs.

PHENCYCLIDINE

A contraction of PHENYL CYCLOHEXYL PIPERIDINE, or PCP.

Phencyclidine is the name of one of the seven drug categories, and is also the name of the major drug in that category.

PHENYL CYCLOHEXYL PIPERIDINE (PCP)

- 1. One of the seven drug categories, often called "phencyclidine".
- 2. A specific drug belonging to the phencyclidine category.

PHYSIOLOGY

The study of living organisms and the changes that occur during activity.

PILOERECTION

Literally, "hair standing up", or goose bumps. This condition of the skin is often observed in persons who are under the influence of LSD.

PSYCHEDELIC

A mental state characterized by a profound sense of intensified or altered sensory perception sometimes accompanied by hallucinations.

PSYCHOPHYSICAL TESTS

Methods of investigating the mental (psycho-) and physical characteristics of a person suspected of alcohol or drug impairment. Most psychophysical tests employ the concept of divided attention to assess a suspect's impairment.

PSYCHOTOGENETIC

Literally, "creating psychosis" or "giving birth to insanity". A drug is considered to be psychotogenetic if persons who are under the influence of the drug become insane, and remain so after the drug wears off.

PSYCHOTOMIMETIC

Literally, "mimicking psychosis" or "impersonating insanity". A drug is considered to be psychotomimetic if persons who are under the influence of the drug look and act insane <u>while</u> they are under the influence.

PTOSIS

Droopy eyelids.

PULSE

The expansion and relaxation of the walls of an artery, caused by the surging flow of blood.

PULSE RATE

The number of expansions of an artery per minute.

REBOUND DILATION

A period of constriction fol-lowed by dilation with a change equal to or greater than 2 mm the final size determination being estima-ted at the end of a 15-second time period in which the light from the penlight is directed into the eye.

RESTING NYSTAGMUS

Jerking of the eyes as they look straight ahead.

RESUME

A written summary of a person's education, training, experience, noteworthy achievements and other relevant information about a particular topic.

SCLERA

A dense white fibrous membrane that, with the cornea, forms the external covering of the eyeball (i.e., the white part of the eye).

SENSORY NERVES

Nerves that carry messages to the brain, from the various parts of the body, including notably the sense organs(eyes, ears, etc.). Sensory nerves are also known as afferent nerves.

SINSEMILLA

The unpollenated female cannabis plant, having a relatively high concentration of THC.

SFST

Standardized Field Sobriety Testing. There are three SFSTs, namely Horizontal Gaze Nystagmus (HGN), Walk and Turn, and One Leg Stand. Based on a series of controlled laboratory studies, scientifically validated clues of alcohol impairment have been identified for each of these three tests. They are the <u>only</u> Standardized Field Sobriety Tests for which validated clues have been identified.

SNORTING

One method of ingesting certain drugs. Snorting requires that the drug be in powdered form. The user rapidly draws the drug up into the nostril, usually via a paper or glass tube. Snorting is also known as insufflation.

SPHYGMOMANOMETER

A medical device used to measure blood pressure. It consists of an arm or leg cuff with an air bag attached to a tube and a bulb for pumping air into the bag, and a gauge for showing the amount of air pressure being pressed against the artery.

STETHOSCOPE

A medical instrument used, for drug evaluation and classification purposes, to listen to the sounds produced by blood passing through an artery.

SYMPATHETIC NERVE

An autonomic nerve that commands the body to react in response to excitement, stress, fear, etc. The brain uses sympathetic nerves to send "wake up calls" and "fire alarms" to the muscles, tissues and organs.

SYMPATHOMIMETIC DRUGS

Drugs that mimic the neurotransmitter associated with the sympathetic nerves. These drugs artificially cause the transmission of messages that produce elevated blood pressure, dilated pupils, etc.

SYNAPSE (or Synaptic Gap)

The gap or space between two neurons (nerve cells).

SYNESTHESIA

A sensory perception disorder, in which an input via one sense is perceived by the brain as an input via another sense. An example of this would be a person "hearing" a phone ring and "seeing" the sound as a flash of light. Synesthesia sometimes occurs with persons under the influence of hallucinogens.

SYSTOLIC

The highest value of blood pressure. The blood pressure reaches its systolic value when the heart is fully contracted (systole), and blood is sent surging into the arteries.

TACHYCARDIA

Abnormally rapid heart rate; pulse rate above the normal range.

TACHYPNEA

Abnormally rapid rate of breathing.

THC (Tetrahydrocannabinol)

The principal psychoactive ingredient in drugs belonging to the cannabis category.

TOLERANCE

An adjustment of the drug user's body and brain to the repeated presence of the drug. As tolerance develops, the user will experience diminishing psychoactive effects from the same dose of the drug. As a result, the user typically will steadily increase the dose he or she takes, in an effort to achieve the same psychoactive effect.

TRACKS

Scar tissue usually produced by repeated injection of drugs, via hypodermic needle, along a segment of a vein.

VERTICAL GAZE NYSTAGMUS

An up-and-down jerking of the eyeball that occurs as the eyes gaze upward in the vertical plane.

VOIR DIRE

A french expression literally meaning "to see, to say". Loosely, this would be rendered in English as "To seek the truth", or "to call it as you see it". In a law or court context, one application of voir dire is to question a witness to assess his or her qualifications to be considered an expert in some matter pending before the court.

VOLUNTARY NERVE

A motor nerve that carries messages to a muscle that we consciously control.

WITHDRAWAL

This occurs in someone who is physically addicted to a drug when he or she is deprived of the drug. If the craving is sufficiently intense, the person may become extremely agitated, and even physically ill. Withdrawal from heroin is reported to be an especially unpleasant experience.

SESSION II

DRUGS IN SOCIETY AND IN VEHICLE OPERATION

SESSION II DRUGS IN SOCIETY AND IN VEHICLE OPERATION

Upon successfully completing this session, the participant will be able to:

- o Define the term "drug" in the context of this course.
- o Name the seven major categories of drugs that are relevant to the Drug Evaluation and Classification Process.
- o State in approximate, quantitative terms the incidence of drug use among various segments of the American public.
- o State in approximate, quantitative terms the incidence of drug involvement in motor vehicle crashes and other driving incidents.

Content Segments

A. Definition and Categories of Drugs

B. Drugs and Driving

Learning Activities

- o Instructor Led Presentations
- o Reading Assignments



II-0A&B (Session Objectives)





35 Minutes

DRUGS IN SOCIETY AND IN VEHICLE OPERATION

- A. Definition and Categories of Drugs
 - 1. What do we mean by the word "drug"?
 - a. Medicines? Are all drugs medicines? Are all medicines drugs?
 - b. Narcotics? Are all drugs narcotics?
 - c. Habit forming substances?
 Are all drugs habit forming?
 Are all habit forming
 substances drugs?
 - 2. A simple, law enforcement oriented definition.

"Any substance, which, when taken into the human body, can impair the ability of the person to operate a vehicle safely." Total Lesson Time: Approximately 50 Minutes

Briefly review the objectives, content and activities of this session.

Session title on wallchart.

Instructor: If this has been covered in the Pre-School, pose this question "What is our working definition of the word 'drug'?" and proceed to number 2.

<u>Pose</u> this question to the students.

Solicit several responses.

This definition is derived from the California Vehicle Code, Section 312.

Point out that this definition excludes many substances that physicians, chemists, etc. might consider to be "drugs", e.g., antibiotics, Novocain, vitamins, etc. It also includes some substances that aren't normally thought of as "drugs", such as model airplane glue, insecticides, etc.



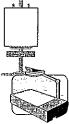
II-1 (Definition of "Drugs")

Instructor Notes

- 3. Within this simple, law enforcement oriented definition, there are seven categories of drugs.
 - a. Each category consists of substances that impair a person's ability to drive.
 - b. The categories differ from one another in terms of <u>how</u> they impair driving ability and in terms of the <u>kinds</u> of impairment they cause.
 - c. Because the categories produce different types of impairment, they generate different signs and symptoms.
 - d. With training and practice, you will be able to recognize the different signs of drug influence and determine which category is causing the impairment you observe in a suspect.

<u>Ask</u> students: "What are the seven categories of drugs?"

Write the names of the categories on the chalkboard or flip-chart as they are mentioned by the students.



11-2

- 4. Central Nervous System Depressants.
 - a. The category of CNS
 Depressants includes some
 of the most commonly
 abused drugs.

<u>Point out</u> that tens of millions of prescriptions for such drugs are written in this country each year.

Aides

	o Alcohol, the most familiar drug of all is abused by an estimated 40-50 million Americans.	
	o In 1998, more than 75 million prescriptions were written for Valium and similar tranquili- zers and sedatives.	Some examples of prescription drugs are Xanax, Prozac, and muscle relaxants.
	b. Depressants slow down the operation of the Central Nervous System (i.e., the brain, brain stem and spinal cord).	·
	o cause the user to react more slowly.	
, , , , , , , , , , , , , , , , , , ,	o cause the user to process information more slowly.	
	o relieve anxiety and tension.	
	o induce sedation, drowsiness and sleep.	
	o in high enough doses, CNS Depressants will produce general anesthesia.	i.e., depress the brain's ability to sense pain.
	o in very high doses, in- duce coma and death.	
II-3	5. Central Nervous System Stimulants	
	 a. CNS Stimulants constitute another widely abused category of drugs. 	
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Lesson Plan

Instructor Notes

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Instructor Notes



- o There appear to be more than 20 million Americans who have used Cocaine.
- o Cocaine is one of the most frequently reported drugs in overdose cases treated at hospital emergency rooms.
- o Several million Americans appear to use Amphetamines.
- o Methamphetamine is becoming increasingly popular.
- b. CNS Stimulants speed up the operation of the central nervous system, and of the various bodily functions controlled by the Central Nervous System.
 - o cause the user to become hyperactive, extremely talkative.
 - o speech may become rapid and repetitive.
 - o heart rate increases.
 - o blood pressure increases.
 - o body temperature rises, user may become excessively sweaty.

Source: National Institute on Drug Abuse (1988).

NOTE: Estimates of drug use vary widely, especially for illicit drugs such as Cocaine, Methamphetamine, etc.

In February 1989, the **Washington Post** reported an alarming increase, nationally, in use of Methamphetamine.

Aides	Aides Lesson Plan		Instructor Notes
		o induce emotional excitement, restlessness, irritability. o can induce cardiac arrhythmia (abnormal beating of the heart), cardiac seizures and death.	Remind students of well-known athletes and others who have died because of Cocaine abuse.
II-4	6. Ha	llucinogens	
	a.	Hallucinogens are also widely abused.	Point out that LSD and Peyote are only two examples of Hallucinogens. There are many other Hallucinogens.
	b.	In recent years, significant increases in the abuse of both LSD and "Ecstasy" (MDMA) have been reported.	In many communities, LSD has become the drug of choice among high school students.
	c.	Hallucinogens create perceptions that differ from reality.	
	d.	These perceptions are often very distorted, so that the user sees, hears and smells things in a way quite different from how they really look, sound and smell.	
	e.	Hallucinogens cause the nervous system to send strange or false signals to the brain.	Clarification: Hallucinogens confuse the Central Nervous System (as well as speeding it up, like CNS Stimulants).
		o Produce sights, sounds, odors, feelings and tastes that aren't real.	

Aides	Lesson Plan	Instructor Notes	
	o Induce a temporary condition very much like psychosis or insanity.		
	o Can create a "mixing" of sensory modalities, so that the user "hears colors", "sees music".	Point out that this mixing of the senses is called Synesthesia. Point out that, with all of these false, and distorted perceptions, a person under the influence of hallucinogens would be a very unsafe driver.	
II-5	7. Phencyclidine (PCP)		
	a. PCP is considered by the medical community to be a Hallucinogen. However, because of the symptomology it presents, it is in a separate category.	Point out that people under the influence of PCP may exhibit a combination of the signs associated with Hallucinogens, CNS Stimulants and Depressants.	
		Phencyclidine is a short form of the chemical name Phenyl Cyclohexyl Piperidine, from which we get the abbreviation "PCP".	
	b. PCP is a synthetic drug, i.e., it does not occur naturally but must be produced in a laboratory-like setting.	Point out that PCP has many analogs, or "chemical cousins" that are very similar to PCP in chemical structure, and that produce essentially the same effects.	
	c. PCP has some effects that resemble the effects of other categories.	The category "Phencyclidine" consists of PCP and its various analogs.	
	d. PCP is similar to CNS		

Depressants in that it depresses brain wave activity.

- o slows down thought
- o slows reaction time
- o slows verbal responses
- e. But PCP is similar to CNS
 Stimulants in that it
 activates the parts of the
 brain that control emotions,
 the heart and the other
 autonomic systems.
 - o heart rate increases
 - o blood pressure increases
 - o adrenalin production increases
 - o body temperature rises
 - o muscles become rigid
- f. And PCP is similar to Hallucinogens in that it distorts or "scrambles" signals received by the brain.
 - o sight, hearing, taste, smell and touch may all be distorted
 - o user's perception of time and space may be distorted
 - o user may become paranoid, feel isolated and depressed
 - o user may develop a strong fear of and preoccupation with death

Aides	Lesson Plan	Instructor Notes	
	o user may become unpredictably violent		
	g. PCP is also a very powerful pain killer, or anesthetic.	Point out that PCP is known as a "dissociative anesthetic"; it "separates" the user from any sensation of pain without making him or her unconscious.	
II-6	8. Narcotic Analgesics		
3.0	a. There are two subcategories of Narcotic Analgesics.		
	o Opiates are derivatives of Opium.	Point out that Morphine and Codeine are examples of Opiates.	
	o Synthetics are produced chemically in the laboratory. The synthetics are not derived in any way from Opium, but produce similar effects.	Point out that Methadone and Numorphan are examples of Synthetic Narcotics.	
	b. The word "Analgesic" means pain killer. All of the drugs in this category reduce the person's reaction to pain.		
	c. Heroin is one of the most commonly abused of the Narcotic Analgesics. Its use is on the rise. There are an estimated one-half million		

Heroin addicts in America.

d. Heroin is highly addictive.

Aides	Lesson Plan		Instructor Notes
		o many addicts support their habit by stealing property and converting it to cash. o America's narcotic	Note: That is \$8,000 worth of
		addicts annually steal property estimated to have a value of \$4 billion.	stolen property annually per Heroin addict.
	e.	In addition to reducing pain, Narcotic Analgesics produce euphoria, drowsiness, apathy, lessened physical activity and sometimes impaired vision.	
	f.	Persons under the influence of Narcotic Analgesics often pass into a semi-conscious type of sleep or near-sleep. o they often are sufficiently alert to respond to questions effectively.	<u>Point out</u> that this condition is often called being "on the nod".
	g.	Higher doses of Narcotic Analgesics can induce coma, respiratory failure and death.	
	9. In	halants	
II-7	a.	Inhalants are the fumes of certain substances. Inhalant abuse is on the rise.	
	b.	These substances are found in many common products.	
		o gasoline o oil-based paints o glue o aerosol cans	
	•		

Aides	Lesson Plan	Instructor Notes	
	o varnish remover o cleaning fluids o etc.		
	c. Different Inhalants produce different effects.		
	o many produce effects similar to those of CNS Depressants.		
	o a few produce Stimulant-like effects.		
	o some produce Hallucinogenic effects.		
	d. The Inhalant abuser's attitude and demeanor can vary from inattentive, stuporous and passive to irritable, violent and dangerous.	,	
B	e. The abuser's speech will often be slow, thick and slurred.		
II-8	10. Cannabis		
The state of the s	a. The category "Cannabis" includes the various forms and products of the Cannabis Sativa plant and other species of Cannabis plants.	Write "Cannabis Sativa" on the chalkboard or flip chart.	
	b. The primary active ingredient in Cannabis products is the substance known as "Delta-9 Tetrahydrocannabinol", or "THC".	Write "Δ-9 THC" on the chalkboard or flip-chart.	

Aides		Lesson Plan	Instructor Notes
	M co	part from alcohol, arijuana is the most mmonly abused drug in is country.	
	19 A re th (4 us	n a household survey in 296, almost 27% of mericans age 18-25 eported using Marijuana in ne past year, nearly half 18%) indicated they had sed Marijuana during their fetimes.	Source: White House Office of National Drug Control Policy; 1996
	ir p: ir d	annabis appears to nterfere with the attention rocess. Drivers under the nfluence of Marijuana often o not pay attention to their riving.	Point out that divided attention Standardized Field Sobriety Tests usually disclose some of the best evidence of Cannabis impairment.
	d p ii o	Cannabis also produces a istortion of the user's erception of time, an acreased heart rate (often ver 100 beats per minute) and a reddening of the eyes.	
II-9 (Drug Combina- tions)	11. Drug	; Combinations	
	k n	Many drug users appear to be "chemical gluttons". They often ingest more than one drug at a time.	
[1] H	1	The term for this is polydrug use"	Note: "poly" is the Greek prefix for "many".
CONTROL OF THE PARTY OF THE PAR		Some very common examples of polydrug use include:	Write "polydrug use" on the chalkboard or flip-chart.

Aides		Lesson Plan	Instructor Notes
		o Alcohol with virtually any other drug.	
		o Marijuana and PCP	Point out that a common way to ingest PCP is to sprinkle it on a Marijuana "joint" and smoke it.
		o Cocaine and Heroin	Sometimes called a "speedball".
		o Heroin and Amphetamine	Sometimes called a," poor man's speedball".
		o Heroin and PCP	Sometimes called a "fireball".
•		o "Crack" Cocaine and PCP	This is sometimes called a "space base".
		o "Crack" Cocaine and Marijuana	Sometimes called a "primo".
		o "Crack" and Methamphetamine	Sometimes called "croak".
	d.	Sometimes, people take two different drugs (such as Heroin and Cocaine) that	Example: o Heroin tends to lower blood pressure.
		produce some opposite effects.	o Cocaine tends to elevate blood pressure
	e.	Different drug combinations may produce unique, interactive effects.	Write on chalkboard or flipchart: "Polydrug use unique, interactive effects."
	f.	When a person has ingested multiple drugs, that person will experience multiple drug effects.	Note, however, that under proper medical supervision, specific drugs often are used to reverse overdose conditions.



15 Minutes







- g. However, it is important to bear in mind that, in a polydrug situation, some of the signs of a particular drug may not be evident even though the person is under the influence of that drug.
- B. Drugs and Driving
 - 1. All available information shows that drug use and abuse are widespread among large segments of the American public.
 - a. Fact: 14% of 600 drivers killed in single vehicle crashes in 78-81 in North Carolina had drugs other than alcohol in them at the time of the crash.
 - b. <u>Fact</u>: 1997 Monitoring the Future Study: Drug use among high school seniors:

Drug	Ever used	Past year	Past month
Marijuana	49.6%	38.5%	23.7%
Cocaine	8.7	5.5	2.3
Crack	3.9	2.4	0.9
CNS Stimulants	16.5	10.2	4.8
LSD	13.6	8.4	3.1
PCP	3.9	2.3	0.7
Heroin	2.1	1.2	0.5

Source: North Carolina, 1981

Source: National Institute on Drug Abuse; 1997.

		-	
A	1	d	es

Instructor Notes



c. <u>Fact</u>: More than 75 million prescriptions for Valium, Librium and similar tranquilizers are written in America annually.

Source: Washington Post, February 17, 1987.

- d. Fact: An estimated 1.6 million Americans age 25 and under reported using cocaine in the past year. An estimated 9.4 million Americans in the same age group reported using Marijuana in the past year.

Source: Substance Abuse and Mental Health Services Administration; 1996

e. Fact: Nearly less than half (49%) of inmates surveyed in state prisons reported being under the influence of drugs or alcohol while committing the offense they were incarcerated for. 17% reported committing the offense for money to buy drugs.

Source: Bureau of Justice Statistics; 1991

- 2. Evidence of drug use frequently shows up in people killed or injured in motor vehicle crashes.
 - a. <u>Fact</u>: University of Tennessee (1988) found 40% of crash injured drivers had drugs other than alcohol in them.
 - b. <u>Fact</u>: The Maryland Shock Trauma Center (1986) found nearly one-third of crash injured drivers had recently used Marijuana.

Instructor Notes



II-10





- c. <u>Fact</u>: A study of young male drivers fatally injured in California crashes found that 51% had used drugs other than alcohol.
- d. Fact: A study completed in 2000, of 880 crash-injured drivers in Rochester, New York, found that 33% had used drugs.
- 3. The facts are unmistakable:
 - a. Drug use is common among many Americans.
 - b. So is drug impaired driving.

Source: Compton, R. and Anderson, T., The Incidence of Driving Under the Influence of Drugs: 1985. National Highway Traffic Safety Administration, 1985.

Research Accident Investigation Team, Department of Community and Preventative Medicine, University of Rochester

INSTRUCTOR PLEASE
NOTE: You should consult the "DRE" newsletter and other sources for updated statistics on drugs and driving.
Solicit students' comments and questions about drugs in society and vehicle operation

Session II

Drugs in Society and in Vehicle Operation



Drugs in Society and in Vehicle Operation

Upon successfully completing this session, the participant will be able to:

- Define the term "drug" in the context of this course
- Name the seven major categories of drugs that are relevant to the Drug Evaluation and Classification process

Drug Evaluation & Classification Training

II-OA

Drugs in Society and in Vehicle Operation (continued)

- State in approximate, quantitative terms the incidence of drug use among various segments of the American public
- State in approximate, quantitative terms the incidence of drug involvement in motor vehicle crashes and other driving incidents

Drug Evaluation & Classification Trainin

II-GE

Working Definition of "Drug":

Any substance which, when taken into the human body, can impair the ability of the person to operate a vehicle safely

Drug Evaluation & Classification Training

11.4

Central Nervous System Depressants



Examples:

- Alcohol
- Barbiturates
- Tranquilizers
- Anti-Anxiety



Drug Evaluation & Classification Training

11-2

Central Nervous System Stimulants

Examples:

- Cocaine
- Amphetamine
- Methamphetamine
- Ritalin



Drug Evaluation & Classification Training

11-3

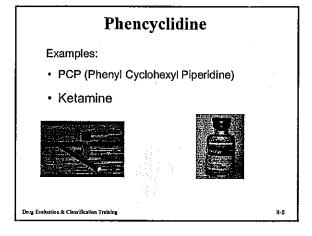
Examples:

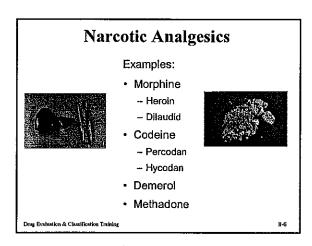
• LSD

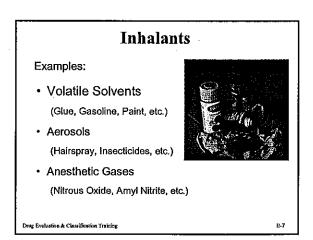
Peyote

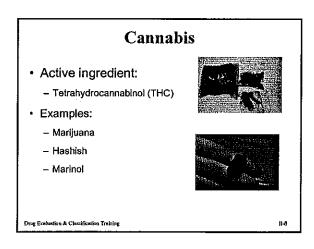
Drug Evaluation & Classification Training

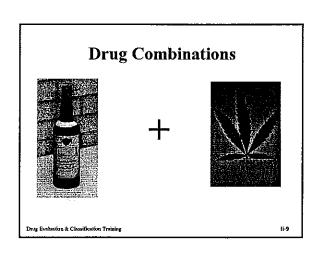
Hallucinogens · MDMA (Ecstasy) Psilocybin



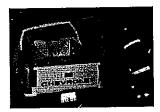






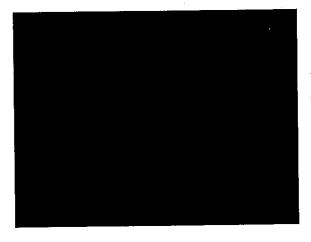


A study of young male drivers fatally injured in California crashes found that 51% had used drugs other than alcohol.



Drug Evaluation & Classification Training

II-10



Fifty Minutes

SESSION III

DEVELOPMENT AND EFFECTIVENESS OF THE DRUG EVALUATION AND CLASSIFICATION PROGRAM

SESSION III DEVELOPMENT AND EFFECTIVENESS OF THE DRUG EVALUATION AND CLASSIFICATION PROGRAM

Upon successfully completing this session, the participant will be able to:

- o State the origin and evolution of the drug evaluation and classification program.
- o Describe research and demonstration project results that validate the effectiveness of the program.
- o State the impact of legal precedents established by case law.
- o Correctly answer the "topics for study" questions at the end of this Section.

Content Segments

Learning Activities

- A. Origin and Evolution of Drug Evaluation & Classification Program
- o Instructor Led Presentations
- B. Evidence of Effectiveness
- o Reading Assignments
- C. Case Law Review

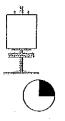
Aides

Lesson Plan

Instructor Notes



III-0A&B (Session Objectives)



15 Minutes



DEVELOPMENT AND
EFFECTIVENESS OF THE DRUG
EVALUATION AND
CLASSIFICATION PROGRAM

- A. Origin and Evolution of the Drug Evaluation and Classification (DEC) Program
 - The Drug Evaluation and Classification Program was developed by personnel of the Los Angeles Police Department.
 - 2. Development of the DEC program began in the early 1970's, in response to a growing awareness that many people apprehended for impaired driving were under the influence of drugs other than alcohol.
 - 3. Individuals principally responsible for initiation and development of the program.
 - a. Dick Studdard (A Traffic Officer)
 - o encountered many impaired drivers whose BACs were zero or very low.
 - o occasionally succeeded in having physicians examine some of these low BAC subjects, resulting in diagnosis of drug influence.

Total lesson time: Approximately 50 Minutes

Briefly review the content, objectives and activities of this session.

Session title on wall chart.

Write: "LAPD" on chalkboard or flip chart.

Sergeant Studdard retired from the LAPD in June, 1990.

<u>Note</u>: examining physicians subsequently would be subpoenaed to testify in contested cases.

- for various reasons,
 physicians were often
 reluctant or unwilling to
 conduct these examinations and offer opinions.
- Some reasons why doctors may be reluctant:
- (1) They typically receive little training in the recognition of specific signs of drug impairment, particularly at street level doses.
- (2) They may not see the suspect until hours after the drugs were used, by which time the signs and symptoms often have changed.
- o as a result, some drivers whom Studdard and other officers were certain were impaired were not prosecuted or convicted for DWI.
- o Studdard concluded that it was essential to develop diagnostic procedures that officers could use when confronted with persons suspected of drug impairment.
- b. Len Leeds (A Narcotics Officer)
 - o was approached by Studdard and asked to collaborate in the development of a program.

Deceased in 1995.

- o initiated some independent research by consulting with physicians, enrolling in relevant classes, studying text books, technical articles, etc.
- secured management level support within the department to continue research and program development.
- c. As time went on, many other key persons both within and outside LAPD contributed to the development and refinement of the program.
- 4. Around 1979, the DRE program was officially recognized by LAPD.
- 5. The DRE program evolved into what is essentially a three-step process.
 - a. First, establish that the suspect is impaired and verify that his or her alcohol level is not consistent with the degree of impairment that is evident.
 - b. Second, use some simple diagnostic procedures to determine whether the impairment may stem from illness or injury, requiring prompt medical attention.

Clarification: the first portion of the drug evaluation examination is devoted principally to Standardized Field Sobriety Testing of the suspect, and to the administration of a breath test. Inconsistency between the observed impairment and the BAC suggests the presence of some other drug(s), or some other complicating factor such as an illness or injury.



III-1 ("Three-Step Process")

- Aides
- c. Third, use diagnostic procedures to determine what category (or categories) of drugs is the likely cause of the impairment.
- 6. Key point: the entire examination is standardized.
 - a. Administered exactly the same was to all suspects.
 - b. Administered exactly the same way by all officers.
- 7. The need for diagnostic procedures.

Pose this question: "Why is it necessary for an officer to use diagnostic procedures to determine the category of drugs causing the impairment?"

Follow-up question: "If we see that a suspect is impaired, and the BAC is too low to account for that impairment, why don't we simply obtain a blood sample and ask the laboratory to analyze the sample for all drugs?"

Solicit responses from students.

a. One reason for needing the diagnostic procedures is that we may be called upon to submit evidence of an articulable suspicion of drug influence to support our request for a chemical test of the suspect.

Some courts or motor vehicle hearings officers may find that a low BAC result, by itself, does not provide adequate basis for requesting the suspect to submit to a second chemical test.

- b. Another reason is that the suspect may refuse to submit to the chemical test, denying us of scientific evidence of drug influence. In that case, conviction or acquittal may hinge on the officer's observations and expertise as a drug examiner.
- c. A third reason is that chemical tests usually disclose only that the suspect has used a particular drug recently. The chemical test usually does not indicate whether the drug is psychoactive at the present time.

Thus, the DRE procedures are needed to establish that the suspect not only has used the drug, but also that he or she is <u>under the</u> influence at this time.

d. A fourth reason is that it can be expensive, and require a large sample of blood or urine, to perform a broad analysis for any or all drugs. Practical constraints require that we be able to point the laboratory technician toward those types of drugs most likely to be found in the sample.

<u>Pose this question</u>: "Are there other toxicological samples that can be obtained for drug analysis by the lab?"

Solicit responses on hair and saliva sampling.



e. It is always possible that a person suspected of drug impairment is actually suffering from some medical problem. If a sample is collected, and the suspect are not examined by someone who is qualified, evidence of medical problems may not come to light until it is too late.

Solicit students' questions and comments concerning the origin, evolution and need for the Drug Evaluation and Classification Program.

B. Evidence of Effectiveness



20 Minutes

- LAPD began to work with the National Highway Traffic Safety Administration (NHTSA) on issues relating to this program in the early 1970's.
 - a. The first step was to develop and validate a battery of Standardized Field Sobriety Tests for investigating alcohol impaired driving.
 - b. LAPD personnel played a major role in the research that led to the wide spread use of Horizontal Gaze Nystagmus, the Walk and Turn test, and the One Leg Stand test.
 - c. By the early 1980's, NHTSA completed its validation of the standardized tests for alcohol enforcement.
 - d. At that time, NHTSA began to assist LAPD in validating the drug enforcement program.



III-2 ("Two Stages of Validation") 2. NHTSA assisted LAPD in a two-phased validation study.

- a. Laboratory Validation, using volunteers who ingested selected drugs.
- Field Validation, using persons actually arrested in Los Angeles on suspicion of drug influence.
- The Laboratory Validation took place at Johns Hopkins University in Maryland.
 - a. The drug examiners were senior DREs from LAPD.
 - The laboratory experiments were planned and conducted by researchers from Johns Hopkins.
 - Volunteers each took a "pill" and smoked a "cigarette".
 - d. The "pill" contained either no drug (placebo) or one of the following drugs:
 - o Secobarbital (CNS Depressant)
 - o Valium (i.e. Diazepam -CNS Depressant)
 - o Desoxyn (i.e. Methamphetamine Sulfate - CNS Stimulant)

The LAPD participants:
Dick Studdard
Jerry Powell
Pat Russell
Doug Laird

Aides		Lesson Plan	Instructor Notes
	e.	The "cigarette" contained either Marijuana or no drug (placebo).	
	f.	Neither the volunteers nor the LAPD officers knew what the volunteers had taken.	Note: this condition is known as a "double blind" experiment. The people being tested and the people doing the testing are kept uninformed of the test condition.
	g.	Two different dose levels of Marijuana, Diazepam and Methamphetamine Sulfate were used.	Clarification: some of the Diazepam and Methamphetamine Sulfate pills were "weak", some were "strong". Similarly, some of the Marijuana cigarettes were "weak", some "strong". All of the Secobarbital pills were "strong".
			Instructor: The following is given for your information.
			Normal daily doses for therapeutic purposes:
		:	• Secobarbital: approx 100mgs
:			• Diazepam: 4-40mgs
			• Desoxyn (methamphetamine sulfate): 15mgs
			Doses administered for this study:
			• Secobarbital: 300 mgs
		-	• Diazepam: weak - 15mgs; strong - 30mgs

• Desoxyn: weak - 15mgs strong - 30mgs



- 4. Results of the Johns Hopkins study.
 - a. The DREs were excellent in identifying subjects who received only placebo doses: they classified 95% of the drug free subjects as "not impaired".
 - b. Similarly, they were excellent in identifying the high dose subjects.
 - o they classified as
 "impaired" 98.7% of the
 subjects who received
 Secobarbital or high
 doses of Marijuana,
 Diazepam or
 Methamphetamine
 Sulfate.
 - o they correctly identified the category of drug for 91.7% of those high dose subjects.
 - c. The DREs were less successful in identifying the low dose subjects.
 - o only 17.5% of the subjects who received the low dose of Methamphetamine Sulfate were classified as "impaired".

• Marijuana:

weak - 12 puffs of 1.3% THC cigarettes strong - 12 puffs of 2.8% THC cigarettes

Instructor Notes

- o only 32.5% of the subjects who smoked the "weak" Marijuana cigarettes were classified as "impaired".
- d. The results of the laboratory validation study were considered to be extremely positive.
 - o the DRE procedures correctly identified the category of drugs in more than 90% of the subjects who were impaired.
 - o the procedures only rarely indicated that unimpaired subjects were under the influence of drugs.
- 5. The field validation study was based on 173 people actually arrested on suspicion of driving under the influence of drugs.
 - a. None of the cases involved a crash.
 - b. In all of the cases, the arrested suspects agreed to submit to a blood test.

Emphasize that these low dose subjects probably would never have been stopped by police officers, if they had been driving.

<u>Point out</u> that, during the study period, many other drugged driving arrests were made by LAPD officers.

Aides	Lesson Plan	Instructor Notes
		But the researchers excluded all cases where the suspects refused to give blood, since it would have been impossible to check the DREs accuracy in those cases. Similarly, they excluded all cases that involved crashes, since the suspects' injuries could have confounded the drug examination.
	c. Twenty-eight different DREs from LAPD participated in the examinations of these 173 suspects.	,
	6. Results of the Field Study.	
III-4 (The Los Angeles Field Validation Study)	a. Based on the independent blood tests, only one of the 173 suspects was found to have no alcohol or other drugs.	
	b. Another 10 suspects were found to have only alcohol in them.	POINT OUT that it is possible that these 11 so-called "drug free" suspects may have used drugs that the independent laboratory could not identify, for various reasons.
	·	Even if we assume that these 11 people really had not used any drug other than alcohol, 11 out of 173 is a very small "false positive" rate.
	c. 37 (21%) of the suspects were found to have only one drug other than alcohol.	
	d. 82 had two drugs other than alcohol (47%), and 43 (25%) had three or more drugs other than alcohol.	

Instructor Notes

- e. This means that 125 of the 173 suspects had ingested two or more drugs other than alcohol: That is more than 70% of the suspects.
- f. PCP was the drug most often found among these 173 suspects: more than half of them (56%) had used PCP.
- 7. The Key Finding of this study was the following:

For more than nine out of ten of the suspects (92.5%), the blood test confirmed the presence of at least one drug category "predicted" by the DREs.

- 8. The confirmation rates for specific categories:
 - a. PCP: blood tests con-firmed DREs' predictions in 92% of the cases.
 - b. Narcotic Analgesics: blood tests confirmed 85% of the DREs' predictions.
 - c. Cannabis: blood tests confirmed 78% of DREs' predictions.

Emphasize: Polydrug use is very common.

Write on chalkboard "70% two or more drugs other than alcohol".





III-6 (Confirmation ...)

POINT OUT that in the other 8% it is possible that <u>a PCP</u> analog might have been used.

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3				

Instructor Notes

- d. CNS Depressants: blood tests confirmed 50% of DREs' opinions.
- e. CNS Stimulants: blood tests confirmed 33% of DREs' opinions.

- 9. Numerous states have conducted comparisons of laboratory analysis and DRE opinions. The correlation rates exceeded 80% in those studies.
- 10. The overall conclusion of the laboratory and field studies is that the DRE program is an effective tool for law enforcement.

C. Case Law Review

- Favorable Court Rulings on DRE Procedures
 - Courts in various states have ruled favorably on the DEC Program. Some judges have held that the drug influence examination procedures need to meet the Frye standard for admissibility of "new" scientific evidence, while others have ruled that Frye need not apply.

POINT OUT that there are literally hundreds of different CNS Depressants, many of which may not have been identifiable by the independent laboratory.

EMPHASIZE that, in this study, the blood samples were not frozen after collection. Unfortunately, cocaine continues to degenerate in a blood sample if the sample isn't frozen. It is quite possible that the cocaine had metabolized from some samples before the lab analyzed them.

EMPHASIZE: Simply because a lab cannot find "drugs" in a sample does not guarantee that no drug is present. All labs have some blind spots

Solicit students' questions about the laboratory and field studies.



15 Minutes



b. The <u>Frye</u> standard: "is the procedure or principle espoused accepted by the relevant scientific community?"

NOTE: <u>Frye</u> standard was set by the US Supreme Court in 1923.

Print "Frye Standard" on the chalkboard or flip-chart.

State of Arizona v. Dayton Johnson and Samuel Rodriquez, et al, NOS 90056865 and 90035883, (1990).

State of Minnesota, City of Minneapolis v. Larry Michael Klawitter, 518 N.W.2d 577, (1993).

State of Colorado v. Daniel Hernandez, 92M 181, (1992).

- c. An Arizona court (Tuscon Municipal Court) ruled that the Frye Standard was met. However, upon appeal, The Arizona State Supreme Court ruled that the Frye Standard did not apply to the DEC Program.
- d. A Minnesota Court (City of Minneapolis) ruled that outside of nystagmus, the DEC Program is not subject to the Frye Standard.
- e. A Colorado Court (Boulder County Court) ruled that the procedures used by DREs are not new or novel and the Frye Standard did not apply.

- f. In many jurisdictions, it will not be necessary to have expert scientific testimony to secure admissibility of a DRE's examination of a suspect.
- 2. The DEC program is gaining acceptance in many courts.
- 3. One key element of DEC -namely, Horizontal Gaze
 Nystagmus -- has been
 recognized as meeting the <u>Frye</u>
 standard by several State
 Supreme Courts.
 - a. First to do so was Arizona, in the case known as <u>State</u> vs. Blake.
 - b. Many more State Supreme Courts are expected to rule favorably on HGN in the near future.

Expert testimony regarding drug influence has long been accepted by numerous courts. The components of DRE evaluation are generally accepted in the scientific community. The DEC program simply combined those components into a systematic and standardized procedure. Thus many prosecutors believe that FRYE standards do not apply to DRE evaluations and testimony.

In fact, testimony based on DRE investigation have been accepted by courts for many years.

Print "State vs. Blake" on the chalkboard or flip-chart.

Point out that additional court rulings on HGN are summarized in the Student's Manual.

Emphasize that students should familiarize themselves with the case law on HGN to ensure they avoid the errors that kept that evidence from being admitted in the past.

If there are significant cases concerning DEC or HGN <u>from</u> <u>the students' State</u>, review them at this



III-15

Instructor Notes

- 4. Summary of HGN Case Law.
 - a. The prevailing trend, in recent years, is for courts to admit HGN as evidence of impairment, provided the proper scientific foundation has been laid.
 - b. But courts consistently reject all attempts to introduce HGN as evidence of a quantitative BAC.
 - (1) The court ruled that in cases where there is no chemical test to determine a BAC level, HGN test results can be admitted the same as of Standardized Field Sobriety Tests to show a "neurological dysfunction", one cause of which could be the ingestion of alcohol.

Solicit students' questions and comments about case law.

Write "No Chemical Test - HGN Admissible".

Write on chalkboard or flip chart - "Cannot be used as evidence of specific BAC level".



Session III

Development and Effectiveness of the Drug Evaluation and Classification Program



Development and Effectiveness of the Drug Evaluation and Classification Program

Upon successfully completing this session, the participant will be able to:

- State the origin and evolution of the drug evaluation and classification program
- Describe research and demonstration project results that validate the effectiveness of the program

Drug Evaluation & Classification Training

III-O

Development and Effectiveness of the Drug Evaluation and Classification Program (continued)

- State the impact of legal precedents established by case law
- Correctly answer the "topics for study" questions at the end of this section

Drug Evaluation & Classification Training

III-OB

The Three-Step Process of Drug Evaluation

Step One

Establish that the suspect is impaired

Step Two

Rule out medical impairment

Step Three

Determine the category of drugs involved

rug Evaluation & Classification Training

DJ-1

Two Stages of Validation

Stage One: Laboratory Validation Study Johns Hopkins University

Stage Two: Field Validation Study Los Angeles

Drug Evaluation & Classification Training

111-2

Laboratory Study Results

- DRE officers correctly identified 95% of drugfree subjects as "unimpaired"
- DRE officers classified 98.7% of high-dose subjects as "impaired"
- 3. Correctly identified the category of drugs for 91.7% of high-dose subjects
- 4. DRE officers were less successful in classifying low-dose subjects

rug Evaluation & Classification Training

H-3

The Los Angeles Field Validation Study

- · 173 drivers accused of drug impairment
- · Blood tests "confirmed":
 - One suspect had no drugs or alcohol
 - 10 had alcohol only
 - 37 (21%) had one other drug
 - 82 (47%) had two other drugs
 - 43 (25%) had three or more other drugs

Drug Evaluation & Classification Training

III-4

The Los Angeles Field Validation Study (continued)

 Blood tests confirmed the presence of at least one "predicted" category of drugs for more than 90% of suspects

Drug Evaluation & Classification Training

...

Confirmation Rates for Specific Categories

• 92%: Phencyclidine (PCP)

· 85%: Narcotic analgesics

• 78%: Cannabis

• 50%: CNS depressants

· 33%: CNS stimulants

Drug Evaluation & Classification Trainin



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"Frye" Decisions Regarding Admissibility of Drug Recognition Expert Testimony

"Frye" refers to a United States Federal Court opinion dealing with the admissibility of scientific evidence. The court established that new or novel scientific evidence, or the novel application of scientific principles, must be shown to have met with general acceptance in the relevant scientific community before it can be admitted.

1990

State of Arizona v. Dayton Johnson and Samuel Rodriguez, et al. Defendants Nos 90056865 & 90035883 (Unpublished Opinion).

The Municipal Court of the City of Tucson, County of Pima, State of Arizona

"Virtually all the witnesses agreed that the scientific procedures utilized by trained drug recognition experts are reliable and are generally accepted in the scientific community. The methodology in place, used by trained law enforcement personnel in the field, has been shown to produce reasonably reliable and uniform results that will contribute materially to the ascertainment of the truth."

On May 7, 1992, the Arizona Supreme Court heard oral arguments in a special proceeding regarding this case. The Justices uniformly rejected the application of "Frye" to the DRE procedures. The Chief Justice observed that the component examination procedures had been established for fifty years.

The prosecutors in this case were Tom Rankin (Tucson) and Cliff Vanell (Phoenix). Expert witnesses for the prosecution included: Sgt. Richard Studdard, LAPD, Marcelline Burns, Ph.D., Sgt. Thomas Page, LAPD, Zenon Zuk, M.D., and Eugene Adler, toxicologist.

1991

The people of the State of New York v. Mary Quinn, Defendant, Docket No. 3130122, District Court, Suffolk County, October 24, 1991, 580 N.Y.S. 2d 818, Misc.2d 139 (N.Y.D.C. 1991).

"The Court found the People's evidence to be persuasive. The protocol is relatively simple. Jurors should have no trouble understanding the testimony of the DRE witness."

"Further, nothing contained in the protocol is a new invention. It is rather a compilation of tried and true procedures utilized by medical science and the law enforcement community in similar contexts for many years."

"The Court believes that the protocol's underlying principles are not so hypertechnical nor the skills required so specialized as to require professional medical training."

"The Court holds that the people have successfully established that both the HGN test and the DRE protocol meet the standards enunciated by "Frye" and "Middleton."

The prosecutors in this case were Joe Lombardo and Richard Frankel (Suffolk County). Expert witnesses for the prosecution included: Richard Studdard, retired LAPD Sergeant, Marcelline Burns, Ph.D., Sergeant Thomas Page, LAPD, Technical Sgt. Douglas Paquette, New York State Police, Zenon Zuk, M.D., David Peed, O.D., and Edward Briglia, Ph.D.

1992

County Court, Boulder, Colorado Case No. 92M181 (Unpublished Opinion) People of the State of Colorado v. Daniel Hernandez

"The DRE methods are accepted within the scientific community because they have found to be reliable."

"The Court finds that the expert does have sufficient specialized knowledge to assist the jurors in better deciding whether the defendant drove his car when under the influence of a specific drug. The DRE testimony can be used at trial provided a sufficient foundation is laid." Overall, this court ruled that the procedures used by DRE's are not new or novel scientific techniques that must meet the "Frye" standard.

The prosecutor in this case was David Archeluta (Boulder County). Expert witnesses for the prosecution include: Sergeant Thomas Page, LAPD, Zenon Zuk, M.D., Marcelline Burns, Ph.D., Rick Abbott, M.D., and Laurel Farrell (chemist).

1993

State of Minnesota in Supreme Court, C6-93-2092, filed June 30, 1994. (Unpublished Opinion)

State of Minnesota, City of Minneapolis vs. Larry Michael Klawitter, 518 N.W.2d 577 (1994)

"Given proper foundation and subject to other qualifications, opinion testimony by experienced police officers trained in use of so-called drug recognition protocol is generally admissible in evidence in a trial of a defendant for driving while under the influence of a controlled substance."

The Court determined that the gaze nystagmus test satisfies the requirements of "Frye".

"We agree with the trial court that the officer should be allowed to give an opinion based on the officer's training and experience and his or her observations following the 12-step drug recognition protocol, as long as (a) there is sufficient foundation for the specific opinion expressed, (b) the state does not attempt to exaggerate the officer's credentials by referring to the officer as a "Drug Recognition Expert" or to unfairly suggest that the officer's opinion is entitled to greater weight than it deserves, and..." "We add only that it should be obvious that the mere fact that such opinion testimony by itself will be sufficient to support a guilty verdict."

The court also determined that, outside of nystagmus, the components of a DRE examination are not scientifically new and are not subject to the "Frye" test.

The trial court stated, "...there is nothing scientifically new, novel, or controversial about any component of the DRE protocol itself. The symptomatology matrix used by DRE's to reach their conclusions is not new and is generally accepted in the medical community as an accurate compilation of signs and symptoms or impairment by the various drug categories."

The prosecutor in this case was Karen Herland (City of Minneapolis). Expert witnesses for the prosecution included: Sergeant Thomas Page, LAPD, Dr. Marcelline Burns (psychologist), Dr. David Peed (optometrist), Dr. Zenon Zuk (medical doctor), Eugene Adler (criminalist), Dr. S.J. Jejurikar (Minnesota Bureau of Criminal Apprehension), and Robert Meyer (toxicologist).

1994
11th Judicial Circuit in and for Dade County, Florida
Case No. 256998,9-I (Unpublished Opinion)
State of Florida v. Frederick Williams
Judge Maxine Cohen Lando
Original filed January 19, 1995

"Given proper foundation and subject to other qualifications, opinion testimony by an experienced police officer trained in the use of the drug recognition protocol is generally admissible in evidence in a trial of a defendant charged with driving under the influence of a controlled or chemical substance. Furthermore, Horizontal Gaze Nystagmus (HGN) test results are generally admissible to establish (1) that the defendant was impaired; and/or (2) that the defendant was over the legal limit; and/or (3) the defendant's specific breath or blood alcohol level at the time he performed the test."

This court found that the "Frye" standard is inapplicable to the DRE Protocol because neither the protocol nor any of its subsets (including HGN, VGN, and Lack of Convergence) are "scientific".

Further, these tests are neither new nor novel. The Court also state that "Frye" is inapplicable to HGN, VGN, and LOC because none of them are new or novel. "None of these tests or the theories and procedures they encompass, are new, novel, or emerging scientific techniques. The medical and psychological professions have acknowledged the tests' underlying theories and procedures for decades."

The Court concluded:

"Drug recognition training is not designed to qualify police officers as scientists, but to train them as observers. The training is intended to refine and enhance the skill of acute observation...and to focus that power...in a particular situation."

This court followed the Klawitter (Minnesota) decision, that it requires the state to "lay a proper predicate before referring to a DRE as anything other than a DRE or Drug Recognition Evaluator or Examiner."

"The real issue is not the admissibility of the evidence, but the weight it should receive. That is a matter for the jury to decide."

The prosecutor in this case was Steve Talpins (Dade County). Expert witnesses for the prosecution in this case included: Marcelline Burns, Ph.D., Zenon Zuk, M.D., Robert Dobie, M.D., Sergeant Thomas Page, LAPD, and others.

American Prosecutors Research Institute National Traffic Law Center

HORIZONTAL GAZE NYSTAGMUS STATE CASE LAW SUMMARY

INTRODUCTION

The following state case law summary contains the seminal cases for each state, the District of Columbia and the Federal courts on the admissibility of HGN. Three main issues regarding the admissibility of the HGN test are set out under each state: evidentiary admissibility, police officer testimony, and purpose and limits of the HGN test results. The case or cases that address each issue are then briefly summarized and cited.

Alabama

I. Evidentiary Admissibility

HGN is a scientific test that must satisfy the *Frye* standard of admissibility. The Supreme Court of Alabama found that the State had not presented "sufficient evidence regarding the HGN test's reliability or its acceptance by the scientific community to determine if the Court of Criminal Appeals correctly determined that the test meets the Frye standards."

Malone v. City of Silverhill, 575 So.2d 106 (Ala. 1990).

II. Police Officer Testimony Needed to Admit HGN Test Result

The Court did not address this issue.

III. Purpose and Limits of HGN

The Court did not address this issue.

Alaska

I. Evidentiary Admissibility

HGN is a scientific test. It is generally accepted within the relevant scientific community.

Ballard v. Alaska, 955 P.2d 931, 939 (Alaska Ct. App. 1998).

A police officer may testify to the results of HGN testing as long as the government establishes a foundation that the officer has been adequately trained in the test. *Ballard*, 955 P.2d at 941.

III. Purpose and Limits of HGN

HGN testing is "a reliable indicator of a person's alcohol consumption and, to that extent, HGN results are relevant." The court cautioned that the HGN test could not be used to correlate the results with any particular blood-alcohol level, range of blood-alcohol levels, or level of impairment.

Ballard, 955 P.2d at 940.

Arizona

I. Evidentiary Admissibility

HGN is a scientific test that needs to satisfy the *Frye* standard of admissibility. State has shown that HGN satisfies the *Frye* standard. State v. Superior Court (Blake), 718 P.2d 171, 181 (Ariz. 1986) (seminal case on the admissibility of HGN).

II. Police Officer Testimony Needed to Admit HGN Test Result

"The proper foundation for [admitting HGN test results]... includes a description of the officer's training, education, and experience in administering the test and showing that proper procedures were followed."

Arizona ex. rel. Hamilton v. City Court of Mesa, 799 P.2d 855, 860 (Ariz. 1990).

See also Arizona ex. Rel. McDougall v. Ricke, 778 P.2d 1358, 1361 (Ariz. Ct. App. 1989).

III. Purpose and Limits of HGN

HGN test results are admissible to establish probable cause to arrest in a criminal hearing.

State v. Superior Court (Blake), 718 P.2d at 182.

"Where a chemical analysis has been conducted, the parties may introduce HGN test results in the form of estimates of BAC over .10% to challenge or corroborate that chemical analysis."

Ricke, 778 P.2d at 1361.

When no chemical analysis is conducted, the use of HGN test results "is to be limited to showing a symptom or clue of impairment." *Hamilton*, 799 P.2d at 858.

Arkansas

I. Evidentiary Admissibility

Novel scientific evidence must meet the *Prater* (relevancy) standard for admissibility. Because law enforcement has used HGN for over thirty-five years, a *Prater* inquiry is not necessary as the test is not "novel" scientific evidence. *Whitson v. Arkansas*, 863 S.W.2d 794, 798 (Ark. 1993).

II. Police Officer Testimony Needed to Admit HGN Test Result

The Court did not address this issue.

III. Purpose and Limits of HGN

HGN may be admitted as evidence of impairment, but is not admissible to prove a specific BAC.

Whitson, 863 S.W.2d at 798.

California

I. Evidentiary Admissibility

HGN is a scientific test and the *Kelly/Frye* "general acceptance" standard must be applied.

California v. Leahy, 882 P.2d 321 (Cal. 1994).

 $California\ v.\ Joehnk,\ 35\ Cal.\ App.\ 4^{th}$ 1488, 1493, 42 Cal. Rptr. 2d 6, 8 (Cal. Ct. App. 1995).

"...[A] consensus drawn from a typical cross-section of the relevant, qualified scientific community accepts the HGN testing procedures...."

Joehnk, 35 Cal. App. 4th at 1507, 42 Cal. Rptr. 2d at 17.

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officer testimony is insufficient to establish "general acceptance in the relevant scientific community." Leahy, 882 P2d. at 609. Police officer can give opinion, based on HGN and other test results, that defendant was intoxicated. Furthermore, police officer must testify as to the administration and result of the test.

Joehnk, 35 Cal. App. 4th at 1508, 42 Cal. Rptr. 2d at 18.

III. Purpose and Limits of HGN

HGN may be used, along with other scientific tests, as some evidence that defendant was impaired.

Joehnk, 35 Cal. App. 4th at 1508, 42 Cal. Rptr. 2d at 17.

HGN test results may not be used to quantify the BAC level of the defendant. *California v. Loomis*, 156 Cal. App. 3d Supp. 1, 5-6, 203 Cal. Rptr. 767, 769-70 (1984).

Connecticut

I. Evidentiary Admissibility

HGN must meet the *Frye* test of admissibility. In this case, the state presented no evidence to meet its burden under the *Frye* test.

Connecticut v. Merritt, 647 A.2d 1021, 1028 (Conn. App. Ct. 1994).

HGN satisfies the *Porter* standards and is admissible. (In *State v. Porter*, 698 A.2d 739 (1997), the Connecticut Supreme Court held the *Daubert* approach should govern the admissibility of scientific evidence and expressed factors to be considered in assessing evidence.)

Connecticut v. Carlson, 720 A.2d 886 (Conn. Super. Ct. 1998).

II. Police Officer Testimony Needed to Admit HGN Test Result

Must lay a proper foundation with a showing that the officer administering the test had the necessary qualifications and followed proper procedures.

Connecticut v. Merritt, 647 A.2d 1021, 1028 (Conn. App. Ct. 1994).

III. Purpose and Limits of HGN

HGN test results can be used to establish probable cause to arrest in a criminal hearing.

Connecticut v. Royce, 616 A.2d 284, 287 (Conn. App. Ct. 1992).

Delaware

I. Evidentiary Admissibility

HGN evidence is scientific and must satisfy the Delaware Rules of Evidence standard.

Delaware v. Ruthardt, 680 A.2d 349, 356 (Del. Super. Ct. 1996).

HGN evidence is acceptable scientific testimony under the Delaware Rules of Evidence.

Ruthardt, 680 A.2d at 362.

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officer may be qualified as an expert to testify about the underlying scientific principles that correlate HGN and alcohol. Delaware police receiving three-day (twenty-four hour) instruction on HGN test administration are not qualified to do this.

Ruthardt, 680 A.2d at 361-62.

Police officer testimony about training and experience alone, without expert testimony, is not enough foundation to admit HGN test results. *Zimmerman v. Delaware*, 693 A.2d 311, 314 (Del. 1997).

III. Purpose and Limits of HGN

HGN test results admissible to show probable cause in a criminal hearing. *Ruthardt*, 680 A.2d at 355.

HGN test results admissible to show probable cause in a civil hearing. *Cantrell v. Division of Motor Vehicles*, 1996 Del. Super. LEXIS 265 (Del. Super. Ct. Apr. 9, 1996).

HGN test results cannot be used to quantify the defendant's BAC. However, they can be used as substantive evidence that the defendant was "under the influence of intoxicating liquor."

Ruthardt, 680 A.2d at 361-62.

Florida

I. Evidentiary Admissibility

The 3rd District Court found HGN to be a "quasi-scientific" test. Its application is dependent on a scientific proposition and requires a particular expertise outside the realm of common knowledge of the average person. It does not have to meet the *Frye* standard because HGN has been established and generally accepted in the relevant scientific community, and has been *Frye* tested in the legal community. The court took judicial notice that HGN is reliable based on supportive case law from other jurisdictions, numerous testifying witnesses and studies submitted. It is "no longer 'new or novel' and there is simply no need to reapply a *Frye* analysis." Williams v. Florida, 710 So. 2d 24 (Fla. Dist. Ct. App. 1998).

The 4th District Court found HGN to be a scientific test. However, because it is not novel, the *Frye* standard is not applicable. However, "[e]ven if not involving a new scientific technique, evidence of scientific tests is admissible only after demonstration of the traditional predicates for scientific evidence including the test's general reliability, the qualifications of test administrators and technicians, and the meaning of the results." Without this predicate, "the danger of unfair prejudice, confusion of issues or misleading the jury from admitting HGN test results outweighs any probative value." The state did not establish the appropriate foundation for the admissibility of HGN test results.

Florida v. Meador, 674 So. 2d 826, 835 (Fla. Dist. Ct. App. 1996), review denied, 686 So. 2d 580 (Fla. 1996).

II. Police Officer Testimony Needed to Admit HGN Test Result

"We take judicial notice that HGN test results are generally accepted as reliable and thus are admissible into evidence once a proper foundation has been laid that the test was correctly administered by a qualified DRE [Drug Recognition Expert]." Williams, 710 So. 2d at 32.

No evidence presented as to the police officer's qualifications nor administration of the HGN test in this case.

Meador, 674 So. 2d at 835.

III. Purpose and Limits of HGN

The HGN test results alone, in the absence of a chemical analysis of blood, breath, or urine, are inadmissible to trigger the presumption provided by the DUI statute, and may not be used to establish a BAC of .08 percent or more. Williams, 710 So. 2d at 36.

Georgia

I. Evidentiary Admissibility

The HGN test is admissible as a "scientifically reliable field sobriety evaluation" under the *Harper* "verifiable certainty" standard.

Manley v. Georgia, 424 S.E.2d 818, 819-20 (Ga. Ct. App. 1992).

HGN testing is judicially noticed as a scientifically reliable test and therefore expert testimony is no longer required before the test results can be admitted. *Hawkins v. Georgia*, 476 S.E.2d 803, 808-09 (Ga. Ct. App. 1996).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officer, who received specialized training in DUI detection and worked with a DUI task force for two years, was permitted to testify that, in his opinion, defendant was under the influence.

Sieveking v. Georgia, 469 S.E.2d 235, 219-20 (Ga. Ct. App. 1996).

III. Purpose and Limits of HGN

HGN test can be admitted to show that the defendant "was under the influence of alcohol to the extent that it was less safe for him to drive." Sieveking, 469 S.E.2d at 219.

Hawaii

I. Evidentiary Admissibility

HGN is a scientific test. The HGN test is reliable under the Hawaii Rules of Evidence and admissible as "evidence that police had probable cause to believe that a defendant was DUI." Judicial notice of the "validity of the principles underlying HGN testing and the reliability of HGN test results" is appropriate. HGN test results can be admitted into evidence if the officer administering the test was duly qualified to conduct the test and the test was performed properly. Hawaii v. Ito, 978 P.2d 191 (Haw. Ct. App. 1999).

II. Police Officer Testimony Needed to Admit HGN Test Result

Before HGN test results can be admitted into evidence in a particular case, however, it must be shown that (1) the officer administering the test was duly qualified to conduct and grade the test; and (2) the test was performed properly in the instant case.

Hawaii v. Ito, 978 P.2d 191 (Haw. Ct. App. 1999), See also Hawaii v. Toyomura, 904 P.2d

893, 911 (Haw. 1992) and *Hawaii v. Montalbo*, 828 P2d. 1274, 1281 (Haw. 1992).

III. Purpose and Limits of HGN

 $HGN\ test\ can be\ admitted\ as\ "evidence\ that\ police\ had\ probable\ cause\ to\ believe\ that\ a\ defendant\ was\ DUI."$

Hawaii v. Ito, 978 P.2d 191 (Haw. Ct. App. 1999).

Idaho

I. Evidentiary Admissibility

HGN test results admitted under the Idaho Rules of Evidence. Rule 702 is the correct test in determining the admissibility of HGN. *State v. Gleason*, 844 P.2d 691, 694 (Idaho 1992).

II. Police Officer Testimony Needed to Admit HGN Test Result

Officer may testify as to administration of HGN test, but not correlation of HGN and BAC.

State v. Garrett, 811 P.2d 488, 493 (Idaho 1991).

III. Purpose and Limits of HGN

"HGN test results may not be used at trial to establish the defendant's blood alcohol level . . . Although we note that in conjunction with other field sobriety tests, a positive HGN test result does supply probable cause for arrest, standing alone that result does not provide proof positive of DUI...."

Garrett, 811 P.2d at 493.

HGN may be "admitted for the same purpose as other field sobriety test evidence -- a physical act on the part of [defendant] observed by the officer contributing to the cumulative portrait of [defendant] intimating intoxication in the officer's opinion." *Gleason*, 844 P.2d at 695.

Illinois

I. Evidentiary Admissibility

HGN meets Frye standard of admissibility. People v. Buening, 592 N.E.2d 1222, 1227 (Ill. App. Ct. 1992). Despite the ruling of the *Buening* appellate court, the Fourth District Court of Appeals declined to recognize HGN's general acceptance without a *Frye* hearing. The court criticized the *Buening* court for taking judicial notice of HGN's reliability based on the decisions of other jurisdictions.

People v. Kirk, 681 N.E.2d 1073, 1077 (Ill. App. Ct. 1997).

The state supreme court held that the state was <u>no longer required to show than an HGN test satisfied the Frye standard</u> before introducing the results of the test into evidence. Absent <u>proof</u> by the defense that the HGN test was unsound, the State only had to show that the officer who gave the test was trained in the procedure and that the test was properly administered. *The People of the State of Illinois v. Linda Basler*, 2000 Ill. LEXIS 1698 (Ill. 2000)

II. Police Officer Testimony Needed to Admit HGN Test Result

"A proper foundation should consist of describing the officer's education and experience in administering the test and showing that the procedure was properly administered."

Buening, 592 N.E.2d at 1227.

III. Purpose and Limits of HGN

HGN test results may be used to establish probable cause in a criminal hearing. *People v. Furness*, 526 N.E.2d 947, 949 (Ill. App. Ct. 1988).

HGN test results admissible to show probable cause in a civil hearing. *People v. Hood*, 638 N.E.2d 264, 274 (Ill. App. Ct. 1994).

HGN test results may be used "to prove that the defendant is under the influence of alcohol."

Buening, 592 N.E.2d at 1228.

Iowa

I. Evidentiary Admissibility

HGN admissible as a field test under the Iowa Rules of Evidence. "[T]estimony by a properly trained police officer with respect to the administration and results of the horizontal gaze nystagmus test are admissible without need for further scientific evidence."

State v. Murphy, 451 N.W.2d 154, 158 (Iowa 1990).

Police officer may testify about HGN test results under Rule 702 if the officer is properly trained to administer the test and objectively records the results. *Murphy*, 451 N.W.2d at 158.

III. Purpose and Limits of HGN

HGN test results may be used as an indicator of intoxication. *Murphy*, 451 N.W.2d at 158.

Kansas

I. Evidentiary Admissibility

HGN must meet *Frye* standard of admissibility and a *Frye* hearing is required at the trial level. There was no *Frye* hearing conducted and the appellate court refused to make a determination based on the record it had.

State v. Witte, 836 P.2d 1110, 1121 (Kan. 1992).

II. Police Officer Testimony Needed to Admit HGN Test Result

The Court did not address this issue.

III. Purpose and Limits of HGN

The Court did not address this issue.

Kentucky

I. Evidentiary Admissibility

HGN test results admitted due to defendant's failure to object. Commonwealth v. Rhodes, 949 S.W.2d 621, 623 (Ky. Ct. App. 1996).

 Π . Police Officer Testimony Needed to Admit HGN Test Result

The Court did not address this issue.

III. Purpose and Limits of HGN

The Court did not address this issue.

Louisiana

I. Evidentiary Admissibility

HGN meets Frye standard of admissibility.

State v. Breitung, 623 So. 2d 23, 25-6 (La. Ct. App. 1993).

State v. Regan, 601 So. 2d 5, 8 (La. Ct. App. 1992).

State v. Armstrong, 561 So. 2d 883, 887 (La. Ct. App. 1990).

The standard of admissibility for scientific evidence is currently the Louisiana Rules of Evidence.

State v. Foret, 628 So. 2d 1116 (La. 1993).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officer may testify as to training in HGN procedure, certification in the administration of HGN test and that the HGN test was properly administered. *Armstrong*, 561 So. 2d at 887.

III. Purpose and Limits of HGN

The HGN test may be used by the officer "to determine whether or not he [needs] to 'go any further' and proceed with other field tests." *Breitung*, 623 So. 2d at 25.

HGN test results may be admitted as evidence of intoxication. *Armstrong*, 561 So. 2d at 887.

Maine

I. Evidentiary Admissibility

Because the HGN test relies on greater scientific principles than other field sobriety tests, the reliability of the test must first be established.

State v. Taylor, 694 A.2d 907, 912 (Me. 1997).

The Maine Supreme Court took judicial notice of the reliability of the HGN test to detect impaired drivers.

Taylor, 694 A.2d at 910.

"A proper foundation shall consist of evidence that the officer or administrator of the HGN test is trained in the procedure and the [HGN] test was properly administered."

Taylor, 694 A.2d at 912.

III. Purpose and Limits of HGN

HGN test results may only be used as "evidence of probable cause to arrest without a warrant or as circumstantial evidence of intoxication. The HGN test may not be used by an officer to quantify a particular blood alcohol level in an individual case." *Taylor*, 694 A.2d at 912.

Maryland

I. Evidentiary Admissibility

HGN is scientific and must satisfy the *Frye/Reed* standard of admissibility. The Court of Appeals took judicial notice of HGN's reliability and its acceptance in the relevant scientific communities.

Schultz v. State, 664 A.2d 60, 74 (Md. Ct. Spec. App. 1995).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officer must be properly trained or certified to administer the HGN test. [NOTE: In *Schultz*, the police officer failed to articulate the training he received in HGN testing and the evidence was excluded.] *Schultz*, 664 A.2d at 77.

III. Purpose and Limits of HGN

HGN testing may not be used to establish a specific blood alcohol level. *Wilson v. State*, 723 A.2d 494 (Md. Ct. Spec. App. 1999).

Massachusetts

I. Evidentiary Admissibility

HGN is scientific and is admissible on a showing of <u>either</u> general acceptance in the scientific community or reliability of the scientific theory. See Commonwealth v. Lanigan, 641 N.E.2d 1342 (Mass. 1994). HGN test results are inadmissible until the Commonwealth introduces expert testimony to establish that the HGN test satisfies one of these two standards.

Commonwealth v. Sands, 675 N.E.2d 370, 373 (Mass. 1997).

"[T]here must be a determination as to the qualification of the individual administering the HGN test and the appropriate procedure to be followed." In this case there was no testimony as to these facts, thus denying the defendant the opportunity to challenge the officer's qualifications and administration of the test. Sands, 675 N.E.2d at 373.

III. Purpose and Limits of HGN

The Court did not address this issue.

Michigan

I. Evidentiary Admissibility

Court found that HGN test is scientific evidence and is admissible under the *Frye* standard of admissibility.

State v. Berger, 551 N.W.2d 421, 424 (Mich. Ct. App. 1996).

II. Police Officer Testimony Needed to Admit HGN Test Result

Only foundation necessary for the introduction of HGN test results is evidence that the police officer properly performed the test and that the officer administering the test was qualified to perform it.

Berger, 551 N.W.2d at 424.

III. Purpose and Limits of HGN

HGN test results are admissible to indicate the presence of alcohol. *Berger*, 551 N.W.2d at 424 n.1.

Minnesota

I. Evidentiary Admissibility

Court found that HGN meets the *Frye* standard of admissibility. *State v. Klawitter*, 518 N.W.2d 577, 585 (Minn. 1994).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officers must testify about their training in and experience with the HGN test.

See generally Klawitter, 518 N.W.2d at 585-86.

III. Purpose and Limits of HGN

HGN admissible as evidence of impairment as part of a Drug Evaluation Examination in the prosecution of a person charged with driving while under the influence of drugs.

See generally Klawitter, 518 N.W.2d at 585.

Mississippi

I. Evidentiary Admissibility

HGN is a scientific test. However, it is not generally accepted within the relevant scientific community and is inadmissible at trial in the State of Mississippi. Young v. City of Brookhaven, 693 So.2d 1355, 1360-61 (Miss. 1997).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officers cannot testify about the correlation between the HGN test and precise blood alcohol content. *Young*, 693 So.2d at 1361.

III. Purpose and Limits of HGN

HGN test results are admissible only to prove probable cause to arrest. *Young*, 693 So.2d at 1361.

HGN test results cannot be used as scientific evidence to prove intoxication or as a mere showing of impairment. *Young*, 693 So.2d at 1361.

Missouri

I. Evidentiary Admissibility

Court found that HGN test meets the *Frye* standard of admissibility. State v. Hill, 865 S.W.2d 702, 704 (Mo. Ct. App. 1993), rev'd on other grounds, State v. Carson, 941 S.W.2d 518, 520 (Mo. 1997).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officer must be adequately trained and able to properly administer the test. *Hill*, 865 S.W.2d at 704.

III. Purpose and Limits of HGN

HGN can be admitted as evidence of intoxication. *Hill*, 865 S.W.2d at 704.

Montana

I. Evidentiary Admissibility

Court found that HGN is neither new nor novel; thus, Daubert does not apply. Court still finds that

HGN must meet the state's rules of evidence that are identical to the Federal Rules of Evidence.

Hulse v. DOJ, Motor Vehicle Div., 961 P.2d 75, 88 (Mont. 1998).

III. Police Officer Testimony Needed to Admit HGN Test Result

The court held that before an arresting officer may testify as to HGN results, a proper foundation must show that the officer was properly trained to administer the HGN test and that he administered the test in accordance with this training. Before the officer can testify as to the correlation between alcohol and nystagmus, a foundation must be established that the officer has special training in the underlying scientific basis of the HGN test.

Hulse, 961 P.2d 75 (Mont. 1998).

III. Purpose and Limits of HGN

HGN test results admissible as evidence of impairment. *State v. Clark*, 762 P.2d 853, 856 (Mont. 1988).

Nebraska

I. Evidentiary Admissibility

HGN meets the *Frye* standard for acceptance in the relevant scientific communities, and when the test is given in conjunction with other field sobriety tests, the results are admissible for the limited purpose of establishing impairment that may be caused by alcohol.

State v. Baue, 607 N.W.2d 191 (Neb. 2000)

A police officer may testify to the results of HGN testing if it is shown that the officer has been adequately trained in the administration and assessment of the HGN test and has conducted the testing and assessment in accordance with that training. State v. Baue, 607 N.W.2d 191 (Neb. 2000)

III. Purpose and Limits of HGN

"Testimony concerning HGN is admissible on the issue of impairment, provided that the prosecution claims no greater reliability or weight for the HGN evidence than it does for evidence of the defendant's performance on any of the other standard field sobriety tests, and provided further that the prosecution makes no attempt to correlate the HGN test result with any particular blood-alcohol level, range of blood-alcohol levels, or level of impairment."

State v. Baue, 607 N.W.2d 191 (Neb. 2000) (quoting Ballard v. State, 955 P.2d 931, 940 (Alaska App. 1998))

New Jersey

I. Evidentiary Admissibility

In New Jersey, the party offering the results of a scientific procedure into evidence must comply with <u>Frye</u> and show that the procedure is generally accepted in the relevant scientific communities. A party may prove this general acceptance via "(1) testimony of knowledgeable experts[,] (2) authoritative scientific literature[, or] (3) [p]ersuasive judicial decision." Based on the testimony of Dr. Marcelline Burns and Dr. Jack Richman, the Court found the HGN test to be generally accepted and the results thus admissible. The Court also noted the "significant number" of jurisdictions that have accepted the HGN test as admissible scientific evidence. State v. Maida, 2000 N.J. Super. LEXIS 276 (N.J. Super. Ct. Law Div. 2000).

*But See, State v. Doriguzzi, 760 A.2d 336 (N.J. Super. 2000), which held that HGN is scientific evidence that must meet Frye Standard. However, in each trial, sufficient foundation evidence must be laid by expert testimony to assure defendants that a conviction for DUI, when based in part on HGN testing, is grounded in reliable scientific data. In this case, the appellate court reversed defendant's conviction because at trial no such foundation was presented. The court found that because HGN testing has not achieved general acceptance in the community, it is not a matter of which a court can take judicial notice.

The Court did not address this issue.

III. Purpose and Limits of HGN

The Court found the HGN test admissible "as a reliable scientific indicator of likely intoxication."

New Mexico

I. Evidentiary Admissibility

HGN is a scientific test. New Mexico follows the *Daubert* standard, which requires a showing of reliability before scientific evidence can be admitted. The court held that a scientific expert must testify to the underlying scientific reliability of HGN and that a police officer cannot qualify as a scientific expert. Because the State failed to present sufficient evidence regarding the HGN test's reliability, the court remanded the case stating it would be appropriate for the trial court, on remand, to make the initial determination of whether HGN testing satisfies *Daubert*. In addition, the court found HGN to be "beyond common and general knowledge" and declined to take judicial notice of HGN reliability.

State v. Torres, 976 P.2d 20 (N.M. 1999).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officers can qualify as non-scientific experts based on their training and experience. Non-scientific experts may testify about the administration of the test and specific results of the test provided another scientific expert first establishes the reliability of the scientific principles underlying the test. In order to establish the "technical or specialized knowledge" required to qualify as an expert in the administration of the HGN test, "there must be a showing: (1) that the expert has the ability and training to administer the HGN test properly, and (2) that the expert did, in fact, administer the HGN test properly at the time and upon the person in question."

State v. Torres, 976 P.2d 20 (N.M. 1999).

III. Purpose and Limits of HGN

The Court did not address this issue.

New York

I. Evidentiary Admissibility

Quinn held that HGN test results are admissible under Frye standard of "general acceptance." However, the case no longer has precedential value as it was later reversed on other grounds.

People v. Quinn, 580 N.Y.S.2d 818, 826 (Dist. Ct. 1991), rev'd on other grounds, 607 N.Y.S.2d 534 (N.Y. App. Div. 1993).

II. Police Officer Testimony Needed to Admit HGN Test Result

The Court did not address this issue.

III. Purpose and Limits of HGN

The Court did not address this issue.

North Carolina

I. Evidentiary Admissibility

HGN is a scientific test. It "does not measure behavior a lay person would commonly associate with intoxication but rather represents specialized knowledge that must be presented to the jury by a qualified expert." As a result, "until there is sufficient scientifically reliable evidence as to the correlation between intoxication and nystagmus, it is improper to permit a lay person to testify as to the meaning of HGN test results."

State v. Helms, 504 S.E.2d 293 (N.C. 1998).

II. Police Officer Testimony Needed to Admit HGN Test Result Testimony of one police officer, whose training consisted of a "forty hour training class dealing with the HGN test", was inadequate foundation for admission of HGN test results.

Helms, 504 S.E.2d 293 (N.C. 1998).

IV. Purpose and Limits of HGN HGN test results are evidence of impairment. Helms, 504 S.E.2d 293 (N.C. 1998).

North Dakota

I. Evidentiary Admissibility

Court found that HGN test is admissible as a standard field sobriety test. City of Fargo v. McLaughin, 512 N.W.2d 700, 706 (N.D. 1994).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officer must testify as to training and experience and that the test was properly administered.

City of Fargo, 512 N.W.2d at 708.

III. Purpose and Limits of HGN

"... HGN test results admissible only as circumstantial evidence of intoxication, and the officer may not attempt to quantify a specific BAC based upon the HGN test."

City of Fargo, 512 N.W.2d at 708.

Ohio

I. Evidentiary Admissibility

HGN test is objective in nature and does not require an expert interpretation. *State v. Nagel*, 506 N.E.2d 285, 286 (Ohio Ct. App. 1986).

Court determined that HGN was a reliable indicator of intoxication without specifically ruling on whether HGN meets *Frye* or some other standard of admissibility.

State v. Bresson, 554 N.E.2d 1330, 1334 (Ohio 1990).

Court held that SFSTs, including HGN, must be administered in *strict compliance* with NHTSA's directives in order for the test results to be admissible. *State v. Homan*, 732 N.E.2d 952 (Ohio 2000).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officer need only testify to training in HGN procedure, knowledge of the test and ability to interpret results.

Bresson, 554 N.E.2d at 1336.

III. Purpose and Limits of HGN

HGN can be used to establish probable cause to arrest and as substantive evidence of a defendant's

guilt or innocence in a trial for DUI, but not to determine defendant's BAC. *Bresson*, 554 N.E.2d at 1336.

Oklahoma

I. Evidentiary Admissibility

HGN test results excluded because state failed to lay adequate foundation regarding HGN's scientific admissibility under the *Frye* standard of admissibility. Police officer's testimony alone was insufficient.

Yell v. State, 856 P.2d 996, 996-97 (Okla. Crim. App. 1993).

The *Daubert* rationale replaces the *Frye* standard as the admissibility standard for scientific evidence.

Taylor v. State, 889 P.2d 319, 328-29 (Okla. Crim. App. 1995).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officer testified to training on how to administer HGN test and how the test was administered in this case. Officer also testified as to his training in analyzing HGN test results.

Yell, 856 P.2d at 997.

III. Purpose and Limits of HGN

If HGN testing was found to satisfy the *Frye* standard of admissibility, HGN test results would be considered in the same manner as other field sobriety test results. HGN test results are inadmissible as scientific evidence creating a presumption of intoxication.

Yell, 856 P.2d at 997.

Oregon

I. Evidentiary Admissibility

HGN test results are admissible under the Oregon Rules of Evidence. HGN test results are scientific in nature, are relevant in a DUI trial, and are not unfairly prejudicial to the defendant.

State v. O'Key, 899 P.2d 663, 687 (Or. 1995).

"Admissibility is subject to a foundational showing that the officer who administered the test was properly qualified, that the test was administered properly, and that the test results were recorded accurately."

O'Key, 899 P.2d at 670.

III. Purpose and Limits of HGN

"... HGN test results are admissible to establish that a person was under the influence of intoxicating liquor, but is not admissible...to establish a person's BAC...."

O'Key, 899 P.2d at 689-90.

Officer may not testify that, based on HGN test results, the defendant's BAC was over .10.

State v. Fisken, 909 P.2d 206, 207 (Or. Ct. App. 1996).

Pennsylvania

I. Evidentiary Admissibility

The state laid an inadequate foundation for the admissibility of HGN under the *FryelTopa* standard.

Commonwealth v. Moore, 635 A.2d 625, 629 (Pa. Super. Ct. 1993).

Commonwealth v. Apollo, 603 A.2d 1023, 1028 (Pa. Super. Ct. 1992).

Commonwealth v. Miller, 532 A.2d 1186, 1189-90 (Pa. Super. Ct. 1987).

Testimony of police officer is insufficient to establish scientific reliability of HGN test.

Moore, 635 A.2d at 692.

Miller, 532 A.2d at 1189-90.

Testimony of behavioral optometrist did not establish general acceptance of HGN test.

Apollo, 603 A.2d at 1027-28.

II. Police Officer Testimony Needed to Admit HGN Test Result

County detective certified as HGN instructor. Court did not comment on whether this would be enough foundation to allow the detective to testify about HGN test results.

Moore, 635 A.2d 629.

Police officer had one-day course on HGN. Court did not comment on whether this would be enough foundation to allow the officer to testify about HGN test results. *Miller*, 603 A.2d at 1189.

III. Purpose and Limits of HGN

Not addressed by court.

South Carolina

I. Evidentiary Admissibility

HGN admissible in conjunction with other field sobriety tests. By implication, HGN is not regarded as a scientific test.

State v. Sullivan, 426 S.E.2d 766, 769 (S.C. 1993).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officer given twenty hours of HGN training. *Sullivan*, 426 S.E.2d at 769.

III. Purpose and Limits of HGN

HGN test results admissible "to elicit objective manifestations of soberness or insobriety . . . [E]vidence from HGN tests is not conclusive proof of DUI. A positive HGN test result is to be regarded as merely circumstantial evidence of DUI. Furthermore, HGN test shall not constitute evidence to establish a specific degree of blood alcohol content." Sullivan, 426 S.E.2d at 769.

Tennessee

I. Evidentiary Admissibility

HGN is a scientific test. To be admissible at trial, such evidence must satisfy the requirements of Tenn. Rules of Evidence 702 and 703. State provided an inadequate amount of evidence to allow the court to conclude that HGN evidence meets this standard.

State v. Murphy, 953 S.W.2d 200 (Tenn. 1997).

HGN must be offered through an expert witness. To qualify as an expert, a police officer must establish that he is qualified by his "knowledge, skill, experience, training or education" to provide expert testimony to "substantially assist the trier of fact to understand the evidence or determine a fact in issue." Although the court did not rule out the possibility that the officer can be considered an expert, the court set a high level of proof. In this case, the court felt that although the officer had attended law enforcement training in DUI offender apprehension and the HGN test, this training was not enough to establish him as an expert.

State v. Grindstaff, 1998 Tenn. Crim. App. Lexis 339 (March 23, 1998).

III. Purpose and Limits of HGN

The Court did not address this issue.

Texas

I. Evidentiary Admissibility

HGN admissible under the Texas Rules of Evidence. Emerson v. State, 880 S.W.2d 759, 769 (Tex. Crim. App. 1994).

II. Police Officer Testimony Needed to Admit HGN Test Result

A police officer must qualify as an expert on the HGN test, specifically concerning its administration and technique, before testifying about a defendant's performance on the test. Proof that the police officer is certified in the administration of the HGN test by the Texas Commission on Law Enforcement Officer Standards and Education satisfies this requirement.

Emerson, 880 S.W.2d at 769.

III. Purpose and Limits of HGN

HGN admissible to prove intoxication. *Emerson*, 880 S.W.2d at 769.

Utah

I. Evidentiary Admissibility

HGN test admissible as other field sobriety test. Court reserved judgment as to the scientific reliability of HGN. Salt Lake City v. Garcia, 912 P.2d 997, 1001 (Utah Ct. App. 1996).

Police officer need only testify as to training, experience and observations when HGN admitted as a field test. *Garcia*, 912 P.2d at 1001.

III. Purpose and Limits of HGN

Admissible as any other field sobriety test. *Garcia*, 912 P.2d at 1000-01.

Washington

I. Evidentiary Admissibility

It is "undisputed" in the relevant scientific communities that "an intoxicated person will exhibit nystagmus". HGN testing is not novel and has been used as a field sobriety test for "decades" and is administered the same whether investigating alcohol impairment or drug impairment. Thus, the use of HGN in drug and alcohol impaired driving cases is acceptable.

State v. Baity, 140 Wn.2d 1, 991 P.2d 1151 (Wash. 2000).

"[T]he *Frye* standard applies to the admission of evidence based on HGN testing, unless . . . the State is able to prove that it rests on scientific principles and uses techniques which are not 'novel' and are readily understandable by ordinary persons." The state failed to present any evidence to this fact and the court declined to take judicial notice of HGN.

State v. Cissne, 865 P.2d 564, 569 (Wash. Ct. App. 1994).

II. Police Officer Testimony Needed to Admit HGN Test Result

The Court did not address this issue.

III. Purpose and Limits of HGN

The Court did not address this issue.

West Virginia

I. Evidentiary Admissibility

The state did not present evidence for the court to reach "the question of whether the HGN test is sufficiently reliable to be admissible." However, the court did conclude "that even if the reliability of the HGN test is demonstrated, an expert's testimony as to a driver's performance on the test is admissibile only as evidence that the driver was under the influence. Estimates of blood alcohol content based on the HGN test are inadmissible."

State v. Barker, 366 S.E.2d 642, 646 (W. Va. 1988).

The West Virginia Supreme Court modified *State v. Barker* to the extent that the <u>Daubert</u> analysis of FRE 702 is applicable to the question of admissibility of expert testimony under the West Virginia Rules of Evidence Rule 702. *Wilt v. Buracker*, 443 S.E. 2d 196 (W.Va. 1993).

II. Police Officer Testimony Needed to Admit HGN Test Result

Police officer's training consisted of a one-day, eight-hour training session conducted by the state police. Officer testified to giving the HGN test about 100 times. Court did not reach question of whether this would be enough to allow the officer to testify about the HGN test results.

Barker, 366 S.E.2d at 644.

III. Purpose and Limits of HGN

HGN test results admissible to show probable cause in a civil hearing. Muscatell v. Cline, 474 S.E.2d 518, 525 (W. Va. 1996). Boley v. Cline, 456 S.E.2d 38, 41 (W. Va. 1995).

"[I]f the reliability of the HGN test is demonstrated, an expert's testimony as to a driver's performance on the test is admissible only as evidence that the driver was under the influence," the same as other field sobriety tests. *Barker*, 366 S.E.2d at 646.

Wisconsin

I. Evidentiary Admissibility

The court held that the HGN test results are admissible in this case because the test results were not the only evidence. The results were accompanied by the expert testimony of the officer.

State v. Zivcic, 598 N.W.2d 565 (Wisc. Ct. App. 1999).

A police officer who is properly trained to administer and evaluate the HGN test can testify to the test results. A second expert witness is not needed. State v. Zivcic, 598 N.W.2d 565 (Wisc. Ct. App. 1999).

III. Purpose and Limits of HGN

The Court did not address this issue.

Wyoming

I. Evidentiary Admissibility

SFSTs, including HGN, are admissible to establish probable cause when administered in *substantial compliance* with NHTSA guidelines. Strict compliance is not necessary. The court took judicial notice of the number of states that allow HGN evidence on the basis of the "officer's training, experience and ability to administer the test".

Smith v. Wyoming, 2000 Wyo. LEXIS 202 (Wyo. October 4, 2000).

II. Police Officer Testimony Needed to Admit HGN Test Result

A police officer that is properly trained to administer and evaluate the HGN test can testify to HGN results.

Smith v. Wyoming, 2000 Wyo. LEXIS 202 (Wyo. October 4, 2000).

III. Purpose and Limits of HGN

HGN test results are admissible to show probable cause. Smith v. Wyoming, 2000 Wyo. LEXIS 202 (Wyo. October 4, 2000).

United States

I. Evidentiary Admissibility

HGN test was admitted as part of series of field tests. Its admission was not challenged on appeal.

U.S. v. Van Griffin, 874 F.2d 634 (9th Cir. 1989).

II. Police Officer Testimony Needed to Admit HGN Test Result

The Court did not address this issue.

III. Purpose and Limits of HGN

The Court did not address this issue.

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SCIENTIFIC PUBLICATIONS AND RESEARCH REPORTS ADDRESSING NYSTAGMUS

- 1. Anderson, Schweitz & Snyder, <u>Field Evaluation of Behavioral Test Battery for DWI</u>, U.S. Dept. of Transportation Rep. No. DOT-HS-806-475 (1983) (field evaluation of the Standardized Field Sobriety Test Battery (HGN, one-leg stand, and walk and turn) conducted by police officers from four jurisdictions indicated that the battery was approximately 80% effective in determining BAC above and below .10 percent).
- 2. Aschan, <u>Different Types of Alcohol Nystagmus</u>, 140 ACTA OTOLARYNGOL SUPP. 69 (Sweden 1958) ("From a medico-legal viewpoint, <u>simultaneous</u> recording of AGN (Alcohol Gaze Nystagmus) and PAN (positional alcoholic nystagmus) should be of value, since it will show in which phase the patient's blood alcohol curve is...").
- 3. Aschan & Bergstedt, <u>Positional Alcoholic Nystagmus in Man Following Repeated Alcohol Doses</u>, 80 ACTA OTOLARYNGOL SUPP. 330 (Sweden 1975) (abstract available on DIALOG, file 173: Embase 1975-79) (degree of intoxication influences both PAN I and PAN II).
- 4. Aschan, Bergstedt, Goldberg & Laurell, <u>Positional Nystagmus in Man During and After Alcohol Intoxication</u>, 17 Q.J. OF STUD. ON ALCOHOL, Sept. 1956, at 381. Study distinguishing two types of alcohol-induced nystagmus, PAN (positional alcoholic nystagmus) I and PAN II, found intensity of PAN I, with onset about one-half hour after alcohol ingestion, was proportional to amount of alcohol taken.
- 5. Baloh, Sharma, Moskowitz & Griffith, <u>Effect of Alcohol and Marijuana on Eye Movements</u>, 50 AVIAT. SPACE ENVIRON. MED., Jan 1979, at 18 (abstract available on DIALOG, file 153: Medline 1979-79) (smooth pursuit eye movement effects of alcohol overshadowed those of marijuana).
- 6. Barnes, The Effects of Ethyl Alcohol on Visual Pursuit and Suppression of the Vestibulo-Ocular Reflex, 406 ACTA OTOLARYNGOL SUPP. 161 (Sweden 1984) (ethyl alcohol disrupted visual pursuit eye movement by increasing number of nystagmic "catch-up saccades").
- 7. Burns & Moskowitz, <u>Psychophysical Tests for DWI Arrest</u>, U.S. Dept. of Transportation Rep. No. DOT-HS-802-424 (1977) (recommended the three-test battery developed by SCRI (one-leg stand, walk and turn, and HGN) to aid officers in discriminating BAC level).

- 8. Church & Williams, <u>Dose- and Time-Dependent Effects of Ethanol</u>, 54 ELECTROENCEPHALOGRAPHY & CLIN. NEUROPHYSIOL., Aug. 1982, at 161 (abstract available on DIALOG, file 11: Psychinfo 1967-85 or file 72: Embase 1982-85) (positional alcohol nystagmus increased with dose levels of ethanol).
- 9. Compton, <u>Use of the Gaze Nystagmus Test to Screen Drivers at DWI Sobriety Checkpoints</u>, U.S. Dept. of Transportation (1984) (field evaluation of HGN test administered to drivers through car window in approximately 40 seconds: "the nystagmus test scored identified 95% of the impaired drivers" at 2; 15% false positive for sober drivers, <u>id.</u>).
- 10. Fregly, Bergstedt & Graybiel, <u>Relationships Between Blood Alcohol, Positional Alcohol Nystagmus and Postural Equilibrium</u>, 28 Q.J. OF STUD. ON ALCOHOL, March 1967, at 11, 17 (declines from baseline performance levels correlated with peak PAN I responses and peak blood alcohol levels).
- 11. Goldberg, <u>Effects and After-Effects of Alcohol, Tranquilizers and Fatigue on Ocular Phenomena</u>, ALCOHOL AND ROAD TRAFFIC 123 (1963) (of different types of nystagmus, alcohol gaze nystagmus is the most easily observed).
- 12. Helzer, <u>Detection DUIs Through the Use of Nystagmus</u>, LAW AND ORDER, Oct. 1984, at 93 (nystagmus is "a powerful tool for officers to use at roadside to determine BAC of stopped drivers...(O)fficers can learn to estimate BACs to within an average of 0.02 percent of chemical test readings." Id. at 94).
- 13. L.R. Erwin, DEFENSE OF DRUNK DRIVING CASES (3d ed. 1985) ("A strong correlation exists between the BAC and the angle of onset of (gaze) nystagmus." <u>Id</u>. at 8.15A(3).
- 14. Lehti, The Effect of Blood Alcohol Concentration on the Onset of Gaze

 Nystagmus, 136 BLUTALKOHOL 414 (West Germany 1976) (abstract
 available on DIALOG, file 173: Embase 1975-79) (noted a statistically highly
 significant correlation between BAC and the angle of onset of nystagmus with
 respect to the midpoint of the field of vision).
- 15. Misoi, Hishida & Maeba, <u>Diagnosis of Alcohol Intoxication by the Optokinetic Test</u>, 30 Q.J. OF STUD. ON ALCOHOL 1 (March-June 1969) (optokinetic nystagmus, ocular adaptation to movement of object before eyes, can also be used to detect central nervous system impairment caused by alcohol. Optokinetic nystagmus is inhibited at BAC of only .051 percent and can be detected by optokinetic nystagmus test. Before dosage subjects could follow a speed of 90 degrees per second; after, less than 70 degrees per second).

- 16. Murphree, Price & Greenberg, <u>Effect of Congeners in Alcohol Beverages on the Incidence of Nystagmus</u>, 27 Q.J. OF STUD. ON ALCOHOL, June 1966, at 201 (positional nystagmus is a consistent, sensitive indicator of alcohol intoxication).
- 17. Nathan, Zare, Ferneau & Lowenstein, <u>Effects of Congener Differences in Alcohol Beverages on the Behavior of Alcoholics</u>, 5 Q.J. OF STUD. ON ALCOHOL SUPP., may 1970, at 87 (abstract available on DIALOG, file 11: Psychinfo 1967-85) (incidence of nystagmus and other nystagmoid movements increased with duration of drinking).
- 18. Norris, The Correlation of Angle of Onset of Nystagmus With Blood Alcohol Level: Report of a Field Trial, CALIF. ASS'N CRIMINALISTICS NEWSLETTER, June 1985, at 21 (The relationship between the ingestion of alcohol and the inset of various kinds of nystagmus "appears to be well documented." Id. "While nystagmus appears to be useful as a roadside sobriety test, at this time, its use to predict a person's blood alcohol level does not appear to be warranted." Id. at 22).
- 19. Nuotto, Palva & Seppala, <u>Naloxone Ethanol Interaction in Experimental and Clinical Situations</u>, 54 ACTA PHARMACOL. TOXICOL. 278 (1984) (abstract available on DIALOG, file 5: Biosis Previews 1981-86) (ethanol alone dose-dependently induced nystagmus).
- 20. Oosterveld, Meineri & Paolucci, <u>Quantitative Effect of Linear Acceleration on Positional Alcohol Nystagmus</u>, 45 AEROSPACE MEDICINE, July 1974, at 695 (G-loading brings about PAN even when subject has not ingested alcohol; however when subjects ingested alcohol, no PAN was found when subjects were in supine position, even with G-force at 3).
- 21. Penttila, Lehti & Lonnqvist, <u>Nystagmus and Disturbances in Psychomotor Functions Induced by Psychotropic Drug Therapy</u>, 1974 PSYCHIAT. FENN. 315 (abstract available on DIALOG, file 173: Embase 1975-79) (psychotropic drugs cause nystagmus).
- 22. Rashbass, <u>The Relationship Between Saccadic and Smooth Tracking Eye</u>
 <u>Movements</u>, 159 J. PHYSIOL. 326 (1961) (barbiturate drugs interfere with smooth tracking eye movement).
- 23. Savolainen, Riihimaki, Vaheri & Linnoila, <u>Effects of Xylene and Alcohol on Vestibular and Visual Functions in Man</u>, SCAND. J. WORK ENVIRON. HEALTH 94 (Sweden 1980) (abstract available on DIALOG, file 172: Embase 1980-81 on file 5: Biosis Previews 1981-86) (the effects of alcohol on vestibular functions (e.g., positional nystagmus) were dose-dependent).

- 24. Seelmeyer, Nystagmus, A Valid DUI Test, LAW AND ORDER, July 1985, at 29 (Horizontal Gaze Nystagmus test is used in "at least one law enforcement agency in each of the 50 states" and is "a legitimate method of establishing probable cause." Id.).
- 25. Tharp, Burns & Moskowitz, <u>Circadian Effects on Alcohol Gaze Nystagmus</u> (paper presented at 20th annual meeting of Society for Psychophysiological Research), abstract in 18 PSYCHOPHYSIOLOGY, March 1981 (highly significant correlation between angle of onset of AGN and BAC).
- 26. Tharp, Burns & Moskowitz, <u>Development and Field Test of Psychophysical</u>

 <u>Tests for DWI Arrests</u>, U.S. Dept. of Transportation Rep. No. DOT-HS-805-864
 (1981) (standardized procedures for administering and scoring the SCRI
 three-test battery; participating officers able to classify 81% of volunteers above or below .10).
- 27. Umeda & Sakata, <u>Alcohol and the Oculomotor System</u>, 87 ANNALS OF OTOLOGY, RHINOLOGY & LARYNGOLOGY, May-June 1978, at 392 (in volunteers whose "caloric eye tracking pattern" (CETP) was normal before alcohol intake, influence of alcohol on oculomotor system appeared consistently in the following order: (1) abnormality of CETP, (2) positional alcohol nystagmus, (3) abnormality of eye tracking pattern, (4) alcohol gaze nystagmus).
- 28. Wilkinson, Kime & Purnell, <u>Alcohol and Human Eye Movement</u>, 97 BRAIN 785 (1974) (oral dose of ethyl alcohol impaired smooth pursuit eye movement of all human subjects).
- 29. Zyo, Medico-legal and Psychiatric Studies on the Alcohol Intoxicated Offender, 30 JAPANESE J. OF LEGAL MED., No. 3, 1976, at 169 (abstract available on DIALOG, file 21: National Criminal Justice Reference Service 1972-85) (recommends use of nystagmus test to determine somatic and mental symptoms of alcohol intoxication as well as BAC).

Two Hours and Thirty Minutes

SESSION IV

OVERVIEW OF DRUG RECOGNITION EXPERT PROCEDURES

SESSION IV OVERVIEW OF DRUG RECOGNITION EXPERT PROCEDURES

Upon successfully completing this session, the participant will be able to:

- o Name the components of the drug evaluation and classification process.
- o State the purposes of each component.
- o Describe the activities performed during each component.
- o Correctly answer the "Topics for Study" questions at the end of this section.

Content Segments Learning Activities A. Components of the Process Instructor Led Presentations В. Interview of the Arresting Officer Instructor Led Demonstrations C. The Preliminary Examination Video Presentations 0 . D. Examinations of the Eyes Reading Assignments E. Divided Attention Psychophysical Tests F. **Examinations of Vital Signs** G. Dark Room Checks of Pupil Size H. Examination of Muscle Tone T. Examination for Injection Sites J. Toxicological Examination K. Video Tape Demonstration

Aides	Lesson Plan	Instructor Notes
	OVERVIEW OF DRUG EVALUATION AND CLASSIFICATION PROCEDURES	Total Lesson Time: Approximately 150 Minutes Session title on wall chart.
IV-O Objectives		
	A. Components of the Process	
35 Minutes	The DRE procedure is a standardized and systematic method of examining a subject to determine:	
	a. Whether subject is impaired.	
	b. Whether the impairment is caused by drugs or a medical condition.	
	c. And if drugs, the category (or categories) of drugs that is (or are) the likely cause of the subject's impairment.	
	2. The process is <u>systematic</u> in that it is based on a careful assessment of a variety of observable signs and symptoms that are known to be reliable indicators of drug impairment.	Write on chalkboard or flip-chart: "A SYSTEMATIC PROCESS"
, 22 ¢	a. Some of these observable signs and symptoms relate to the subject's <u>appearance</u> .	Write "appearance" on chalkboard or flip chart.
	b. Some of the signs and symptoms relate to the subject's <u>behavior</u> .	Write "behavior" on chalkboard or flip chart.

Aides	Lesson Plan	Instructor Notes
	c. Some relate to the subject's performance of carefully administered psychophysical tests.	Write "psychophysical testing" on chalkboard or flip chart. Ask students: "What does 'psychophysical' mean?" Point out that "psycho-physical" relates to the subject's mind (psyche) and body (physique).
	 o Drugs impair the subject's ability to control his or her mind and body. o Psychophysical tests can disclose that the subject's ability to control mind and body is impaired. o The specific manner in which the subject performs the psychophysical tests may help indicate the category or 	
	categories of drugs causing the impairment. d. Some of the observable signs and symptoms relate to <u>automatic responses</u> of the subject's body to the specific drugs that are present. e. <u>All</u> of these reliable indicators are examined and carefully considered before a judgment is made concerning what categories of drugs are affecting the subject.	Write "automatic responses of the body" on the chalkboard or flip chart.



IV-1 ("Standardized and Systematic")

- 3. The process is <u>standardized</u> in that it is administered exactly the same way, to every subject, by every Drug Recognition Expert.
 - a. Standardization helps to ensure that no mistakes are made.
 - o No examinations are left out.
 - o No extraneous or unreliable "indicators" are included.

Ask students: "Why is it so important to perform the drug evaluation and classification examination in exactly the same way, every time?"

Probe to draw out all major reasons for standardization.

- b. Standardization helps to promote professionalism among drug recognition experts.
- c. Standardization helps to secure acceptance in court.
- 4. The Drug Evaluation and Classification Process has <u>twelve</u> components.
 - a. The <u>Breath Alcohol Test</u> is needed to determine Blood Alcohol Concentration (BAC).



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Instructor Notes

- o The purpose of the breath test is to determine whether the specific drug, alcohol, may be contributing to the impairment observable in the subject.
- o Obtaining an accurate measurement of BAC enables the drug recognition expert to assess whether alcohol may be the sole cause of the observable impairment, or whether it is likely that some other drug or drugs, or other complicating factors are contributing to the impairment.

b. The <u>Interview of the</u> <u>Arresting Officer.</u>

- o In most cases, the suspects you will examine will <u>not</u> be people that <u>you</u> arrested.
- o The arresting officer may have seen or heard things that would be valuable indicators of the kinds of drugs the suspect has ingested.
- o The arresting officer, in searching the suspect, may have uncovered drug related paraphernalia, or even drugs themselves.

Remind students that many suspects who are under the influence of drugs other than alcohol <u>also</u> have alcohol in their bodies.



IV-3 (Interview of ... Officer)

Aides	Lesson Plan	Instructor Notes	
IV-4A&B (Preliminary Examination)	o The arresting officer also may be able to alert you to important information about the suspect's behavior that could be very valuable for your own safety. c. The Preliminary Examination. o The preliminary examination is your first opportunity to observe the suspect closely and directly.		
	o A major purpose of the preliminary examination is to determine if the suspect may be suffering from an injury or some other medical condition not necessarily related to drugs.	Analogy: The preliminary examination is a "fork in the road." It can help you decide whether to continue with the drug examination, to pursue possible medical complication or to proceed with a DWI (alcohol) case.	
	o Another major purpose of the preliminary examination is to begin systematically assessing the suspect's appear- ance, behavior and auto-	Emphasize that the term "preliminary" does not imply "unimportant". Very valuable evidence often comes to light during the preliminary examination.	

matic bodily responses for signs of drug induced

impairment.

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Instructor Notes

o The preliminary examination consists of a series of questions dealing with possible injuries or medical problems; observations of the suspect's face, speech and breath; pupil size and tracking ability; initial checks of the suspect's eyes; and, an initial examination of the suspect's pulse.

While you are assessing the suspect's tracking ability, you can also perform a preliminary assessment of whether Horizontal Gaze Nystagmus is present in the suspect's eyes. In particular, if the nystagmus or "jerking" is observed, an initial estimation of the angle of onset can be made. The approximate angle of onset may help to determine whether the suspect has consumed some drug other than alcohol.

Emphasize that courts generally accept these questions as not being in conflict with the suspect's Constitutional rights. However, the students must comply with their own departments' policies as to whether they should advise suspects of their Constitutional rights before asking these questions.



d. Examinations of the Eyes.

o Certain Drugs produce very easily observable effects on the eyes.

<u>Ask</u> students: "What do we look for, in a suspect's eyes, to determine if he or she may be under the influence of alcohol?"

Probe, as necessary, to draw out the response "nystagmus".

o One of the most dramatic of these effects is <u>nystagmus</u>, which means an involuntary jerking of the eyes.

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Instructor Notes

- Persons under the influence of alcohol usually will exhibit Horizontal Gaze Nystagmus, which is an involuntary jerking of the eyes occurring as the eyes gaze to the side.
- Alcohol is not the only drug that causes Nystagmus.
- Horizontal Gaze Nystagmus is not the only observable effect on the eyes that will be caused by various drugs.
- Divided Attention Psychophysical tests.
 - All drugs that impair driving ability will also impair the suspect's ability to perform certain carefully designed divided attention tests.
 - These tests are familiar to you in the context of examining alcohol impaired suspects.
 - The same tests are very valuable for disclosing evidence of impairment due to drugs other than alcohol.
- f. Examinations of Vital Signs.

Point out that the examinations of the eyes will be covered in much greater depth subsequently.

Ask students: "What does 'divided attention' mean?"

Probe, as necessary, to draw out responses indicating the concept of "concentrating on more than one thing at a time".

Point out that students will have opportunities to practice administering these tests subsequently in the course.



tention Tests)



Examinations)

		-	
Α	Ī	ď	es

Instructor Notes

- o Many categories of drugs affect the operation of the heart, lungs and other major organs of the body.
- o These effects show up during examination of the suspect's <u>vital signs</u>.
- o The vital signs that are reliable indicators of drug influence include blood pressure, pulse, and temperature.

Point out that examinations of vital signs will be covered in depth subsequently, and that students will have ample opportunity to practice measuring vital signs.



IV-8A&B (Dark Room Examinations)

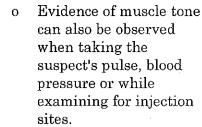
g. Dark Room Examinations

- o Many categories of drugs affect how the pupils will appear, and how they respond to light.
- o Certain kinds of drugs will cause the pupils to widen dramatically, or dilate.
- o Some other drugs cause the pupils to narrow, or constrict...
- o By systematically changing the amount of light entering the suspect's eyes, we can observe the pupils' appearance and reaction under controlled conditions.

Aides	Lesson Plan	Instructor Notes	
	o We carry out these examinations in a dark room, using a penlight to control the amount of illumination entering the suspect's eyes.	Exhibit a penlight.	
	o We use a device called a pupillometer to estimate the size of the suspect's pupils.	Exhibit a pupillometer. Point out that the pupillometer has a series of black circles of various sizes.	
	o Other examinations are also conducted in the darkroom, using the penlight: i.e., examination of the nasal area and mouth for signs of drug use and for concealed contraband.	By lining the circles up along side the suspect's pupil, the pupil's size can be determined Point out that students will have several opportunities to practice conducting dark room examinations subsequently in the course.	
	h. Examination for Muscle Tone.	. ,	
IV-9A&B (Muscle Tone)	o Certain categories of drugs can cause the user's muscles to become markedly tense, and rigid. Others may cause flaccidity, or "rubbery-like" muscle tone.		
	o Evidence of this muscle tone may come to light when the suspect attempts to perform the divided attention test.	Point out that examination for muscle tone will be covered in greater depth subsequently in the course.	



IV-10A&B (Examination for Injection Sites)



- i. Examination for <u>Injection</u> Sites.
 - o Certain drugs are commonly injected by their users, via hypodermic needles.
 - o Heroin is probably most commonly associated with injection, but several other types of drugs also are injected by many users.
 - o Uncovering injection sites on a suspect provides evidence of possible drug use.
- j. Suspect's statements and other observations.
 - o At this point in the examination, the trained DRE should have reasonable grounds to believe that the suspect is under the influence of a drug or drugs.
 - o The DRE should also have at least an articulable suspicion as to the category or categories of drugs causing the impairment.

Ask students: "What drug is most often associated with injection via hypodermic needle?"



Observations)

		-		
А	i	d	Δ	

Instructor Notes

- to interview the suspect to confirm their opinion concerning the drug category or categories involved.
- o The DRE must carefully record the suspect's statements, and any other observations that may constitute relevant evidence of drug induced impairment.

k. Opinions of Evaluator

- o Based on all of the evidence and observations gleaned from the preceding ten steps, the DRE must reach an informed conclusion as to:
 - whether the suspect is under the influence of a drug or drugs
 - if so, the probable category or categories of drugs causing the impairment
- o The DRE must record a narrative summary of the facts forming the basis for their conclusion.

1. Toxicological Examination

Emphasize that any such interview can proceed only in conformance with formal admonition and strict observance of the suspect's Constitutional rights.

Point out that the appropriate procedures for interviewing suspects vary with the probable category or categories of drugs involved.



IV-12 (Opinions of the Evaluator)



IV-13 (Toxicological Examination)

Aides	Lesson Plan	Instructor Notes
·	o The toxicological examination is a chemical test or tests designed to obtain scientific, admissible evidence to substantiate the DRE's conclusion.	
	o Departmental policy and procedures must be carefully and completely followed in requesting, obtaining and handling the toxicological sample.	questions concerning this preview of the Drug Evaluation
	B. Interview of the Arresting Officer	
10 Minutes	1. The purpose of the interview of the arresting officer is to obtain a summary of the suspect's actions, behaviors, etc. that led to the arrest and the suspicion that drugs other than alcohol may be involved.	Emphasize that DREs should form the habit of posing explicit questions to arresting officers. A cursory or open ended interview (e.g., "What do we have here?") may fail to elicit some relevant information, because arresting officers won't always know what is relevant to a drug examination.
	2. Issues concerning the suspect's behavior.	
IV-14A ("Interview: Behavior")	a. Was the suspect operating a vehicle?	·
	b. What actions, maneuvers, etc. were observed?	
	c. Was there a collision? If yes, was the suspect injured?	
	d. Was the suspect observed smoking, drinking or eating?	

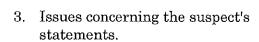
Aides
IV-14B ("Interview: Statements")

Instructor Notes

- e. Was the suspect apparently inhaling any substance?
- f. How did the suspect respond to the arresting officer's command to stop?
- g. Did the suspect attempt to conceal or throw away any items or materials?
- h. What has been the suspect's attitude and demeanor during contact with the arresting officer and have there been any changes?

Ask students to suggest any other questions that might be relevant concerning the arresting officer's observations of the suspect's behavior.

Note: Remind the students that they are acting as investigators and advisors to the arresting officers.



- a. Has the suspect complained of an illness or injury?
- b. Has the suspect used any "street terms" or slang associated with drugs or drug paraphernalia?
- c. How has the suspect responded to the arresting officer's questions?
- d. Does the suspect's speech appear to be slurred, slow, rapid, thick, mumbled, etc.?
- e. What, specifically, has the suspect said to the arresting officer?

Ask students to suggest any other questions that might be relevant concerning statements the suspect made in the arresting officer's presence.

Aides

Lesson Plan

Instructor Notes



IV-14C ("Interview: Physical Evidence")

- 4. Issues concerning physical evidence.
 - a. What items or materials were uncovered during the search of the suspect or vehicle?
 - b. Were any smoking paraphernalia uncovered?
 - c. Were any injection materials, i.e., needles, syringes, leather straps, rubber tubes, spoons, bottle caps, etc. found?
 - d. Were there any balloons, plastic bags, small metal foil wrappings, etc. found?
 - e. What was the suspect's blood alcohol concentration?

NOTE: Emphasize that the suspect should be requested to submit to a breath test, if that has not already been done.

<u>Ask</u> students to suggest any other relevant questions concerning physical evidence.

Solicit students' comments and questions concerning the interview of the arresting officer.



20 Minutes



IV-15 (Overview of Preliminary Examination)

- C. The Preliminary Examination
 - 1. The preliminary examination consists of:
 - a. Questions
 - b. Observations of face, breath and speech.

		•	
Aides		Lesson Plan	Instructor Notes
	c.	Initial checks of the eyes.	
	ď.	The initial check of the suspect's pulse.	Point out that the pulse check actually is part of the examination of the suspect's vital signs. Pulse is checked three times during the Drug Evaluation and Classification Examination.
IV-16	or	ne questions deal with injuries medical problems the suspect ay have.	
("Preliminary Examination Questions")	a.	Are you sick or injured?	Point out that these questions are incorporated into the
questions)	b.	Do you have any physical defects?	Standardized Drug Influence Evaluation Form, which the students will use during all of
	c.	Are you diabetic or epileptic?	their practice sessions.
	d.	Do you take insulin?	Briefly discuss the relevance of each question.
	e.	Are you under a doctor or dentist's care?	
	f.	Are you taking medication?	Show video segment, "Preliminary Examination Questions"

The initial checks of the

pupil.

suspect's eyes include several

particularly important items.

a. Checks of the size of each

A pupillometer is

utilized for this check



("Initial Checks of Eyes")

Point out that, if the two pupils are of unequal size, this may indicate that the suspect is suffering from a head injury, brain tumor, or other condition that may require prompt medical attention.

Aides	Lesson Plan	Instructor Notes	
		Also point out that the influence of certain categories of drugs may be indicated if the pupils are dilated or constricted.	
	b. Assessment of the ability of the eyes to track a moving object.	Demonstrate how to use a stimulus to assess the ability of eyes to track a moving object.	
	o The presence of Nystag- mus indicates the pos- sible presence of certain categories of drugs.	Point out that, if the two eyes do not exhibit the same tracking ability, this too may indicate a head injury or other medical problem.	
	c. Initial estimation of the angle of onset of Horizontal Gaze Nystagmus.	Point out that certain categories of drugs enhance Horizontal Gaze Nystagmus. For example, this will be true of CNS Depressants; PCP; and certain inhalants.	
	o The approximate angle of onset <u>may</u> indicate the presence of some drug other than alcohol.	Remind students that there is a general correspondence, or correlation, between blood alcohol concentration and the onset angle of nystagmus. Generally speaking, the higher the BAC, the earlier will be the angle of onset.	
		But, if the suspect has also ingested some other drug that also enhances Nystagmus, the onset angle may occur even earlier than the Blood Alcohol Concentration would indicate.	
	·	Example: Suppose you are examining a suspect who is known to have a BAC of 0.05.	

Aides	Lesson Plan Instructor N	
	•	Based on that alcohol level alone, you would expect that the angle of onset of nystagmus would be somewhere in the neighborhood of 45 degrees. The formula to estimate the angle of onset, 50 minus the blood alcohol (BA). But if that suspect has also ingested PCP, the onset could occur much earlier, perhaps as soon as the eyes start to move to the side.
		Emphasize if the Nystagmus onset occurs much earlier than would be expected from the alcohol level alone, the DRE should be alert to the possible presence of some drug other than alcohol.
		But also emphasize the Nystagmus onset angle could correspond very closely to what would be expected from the alcohol level alone even though the suspect has ingested large quantities of other drugs.
	·	For example, Cannabis, Narcotic Analgesics, CNS Stimulants and Hallucinogens do <u>not</u> cause nystagmus, and will <u>not</u> affect the onset angle.
	D. Examinations of the Eyes	
IV-18 ("Eye Examinations")	1. The Examinations of the Eyes consist of three tests:	Selectively reveal the items on the slide.

Aides	Lesson Plan	Instructor Notes
	a. Horizontal Gaze Nystagmus.	Emphasize that this test is a full scale, formal and precise examination, unlike the initial estimation of angle of onset conducted during the preliminary examination.
	b. Vertical Gaze Nystagmus.	Point out that Vertical Gaze Nystagmus is an involuntary jerking of the eyes that occurs when the eyes gaze upwards. Select a student, and demonstrate how to perform a test of Vertical Gaze Nystagmus on that student. The instructor should hold the stimulus horizontally in front of the subject's face and about 12-15 inches in front of their nose. Instruct the person to focus on the center of the stimulus, and to keep the head steady. Raise the stimulus until the suspect's eyes are elevated as far as possible. Hold the eyes at that position for four seconds. If the eyes are observed to jerk noticeably, Vertical Gaze Nystagmus is present. Point out that certain types of drugs tend to cause Vertical Gaze Nystagmus, while others do not. Also point out that Vertical Gaze Nystagmus tends to develop with relatively high doses of certain drugs for that individual.
IV-19	c. Lack of Convergence.	Point out that Lack of Convergence is the inability of both eyes to draw in toward the center while fixating on a stimulus being pushed in to the bridge of the nose.

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Aides				

Instructor Notes

- 2. Lack of Convergence is checked by first getting the subject to focus on and track the object while it is slowly moving in a circle in front of the subject's face.
- 3. Then, the object is slowly pushed in and touched to the bridge of the subject's nose and held for approximately 1 second.
- 4. Under the influence of certain types of drugs, the eyes may not be able to converge.

<u>Point out</u> that the circular motion (either left or right) serves to demonstrate that the subject is tracking the object.

<u>Demonstrate</u> this circular motion, using the student volunteer.

<u>Demonstrate</u>, using the student volunteer.

Illustrate on flip chart different examples of Lack of Convergence.

Point out that many people may not be able to converge their eyes.

Excuse the student volunteer and thank him or her for participating.

Solicit students' comments and questions concerning the Examinations of the Eyes.



10 Minutes



IV-20 ("Divided Attention Tests")

- E. Divided Attention Psychophysical Tests
 - 1. The Divided Attention tests used for drug examinations are the same familiar tests used for examining alcohol impaired subjects.

Aides	Lesson Plan	Instructor Notes	
	a. Romberg Balance b. Walk and Turn c. One Leg Stand d. Finger to Nose	Point out that the Romberg Test is administered by asking the suspect to tilt their head back slightly and close the eyes, and estimate 30 seconds, when they believe 30 seconds have passed they are to tilt their head forward, open their eyes and say stop.	
		Point out that the One Leg Stand is administered twice during the drug evaluation and classification examination (once on each leg).	
	2. Walk and Turn demonstration.	Point out that complete demonstrations of all four tests will be given later. For the present, we will demonstrate only the Walk and Turn.	
	a. Instructions stage.	Select a student known to be proficient in administering the Walk and Turn test.	
	b. Walking stage.	Select another student to serve as the test subject.	
		Instruct the student administrator to administer the Walk and Turn test to the student subject.	
		Excuse the students, following the demonstration, and thank them for participating.	
		Point out that students will have numerous opportunities to observe and practice the divided attention tests during the remainder of the course.	

Lesson Plan **Instructor Notes** Aides F. Examinations of Vital Signs 5 Minutes The Vital Signs consist of three things routinely measured in IV-21 ("Vital basic physical examinations. Sign Mea-Blood pressure surements b. Pulse and Instru-Temperature mentation") These measurements require some familiar instruments. Display these items. Stethoscope Blood pressure cuff and gauge (sphygmomanometer) Thermometer (digital, with disposable mouthpieces) Point out that procedures for d. Timepiece capable of measuring blood pressure, measuring in seconds. pulse and temperature will be explained and practiced subsequently. Solicit students' comments and questions concerning examinations of vital signs. G. Dark Room Checks of Pupil Size 15 Minutes



IV-22 ("Dark Room Checks of Pupil Size")

The principal activity that takes place during the dark room examinations is the estimation of pupil size under three lighting conditions.

Aides	Lesson Plan	Instructor Notes
	a. Room light b. Near total darkness c. Direct light	
	2. Another officer should always accompany you and the suspect into the dark room.	Point out that this is essential for officer safety. Remind students that no one should be carrying a weapon when in the presence of a suspect during a drug evaluation and classification examination.
, ·		Point out that some departments require that the suspect be handcuffed before going into the darkroom.
	3. Before turning off the lights, you will estimate the size of the suspect's pupils under room light.	
	a. You must always first estimate the <u>left</u> pupil, then the right .	Point out that the subject should be instructed not to try to focus on you or on the penlight, but to look "slightly up and at a specific focal point" (several feet away) during the estimation of pupil size.
	b. You must position the pupillometer alongside the eye to ensure an accurate estimation.	
	c. After you have completed the room light estimations, turn off the lights and wait 90 seconds to allow your eyes and the suspect's eyes to adapt to the dark.	,

4. The next check will be of pupil size under near total darkness.

Aides	Lesson Plan		Instructor Notes	
	a.	You will need the bare minimum amount of light necessary to see the suspect's pupils and the pupillometer.		
	b,	You can create the necessary light by covering the tip of the penlight with your finger.	Demonstrate this. Point out the reddish glow that emanates through the skin.	
			If possible, darken the room and exhibit the reddish glow.	
	be	ne third and final check will of the pupil size under direct tht.		
	a.	You will shine the full strength of the penlight directly into the subject's eye for 15 seconds.	Point out that it is necessary to maintain reasonably fresh batteries in the penlight.	
	b.	Do this by bringing the light in from the side of the student's face.	<u>Demonstrate</u> this, using the student volunteer.	
	c.	The penlight should be held close enough to the subject's eye so that its beam fills the eye socket.	Demonstrate this. Point out that this will illuminate the area that usually would be discolored if the subject had a "black eye".	
	d.	When the light is initially shown into the eye, you will check for the pupils reaction to light. Then immediately estimate the pupil size under direct light.	If possible, darken the room and exhibit the illumination of the student volunteer's eye socket.	
i				

Aides	Lesson Plan	Instructor Notes
	6. Two other activities are conducted while in the darkroom. a. Examination of the nasal area.	Emphasize that it is very important not to position the penlight too closely or too far away, since this will affect the constriction or dilation of the pupil.
·	b. Examination of the oral cavity.	Excuse the student and thank him or her for participating.
		Solicit students' comments and questions concerning these checks of pupil size.
	H. Examination of Muscle Tone.	
10 Minutes	Starting with the left arm, examine the arm muscles.	
	2. Firmly grasp the upper arm and slowly move down to determine muscle tone.	
	3. The muscles will appear flaccid, normal or rigid to the touch.	
	4. Examine the right arm in the same fashion.	Demonstrate.
	I. Examination for Injection Sites.	
10 Minutes	Some injection sites may be relatively easy to notice.	
	a. Persons who frequently inject certain drugs develop lengthy scars, called "tracks", from repeated injections in the same veins.	

b. Injection of certain drugs

may result in severe caustic action against the skin and flesh, producing easily observable sores.

Aides	Lesson Plan	Instructor Notes	
	2. Often, a <u>fresh</u> injection site may not be readily observable.		
	3. Frequently, a DRE will locate the injection site initially by touch, running the fingers along such commonly used locations	Emphasize that gloves should be worn when touching the suspect.	
,	as the neck, forearms, wrists, back of hand, etc.	Select a student and demonstrate a tactile search for injection sites.	
	4. When the DRE locates a possible injection site, a light	"Ski": short for schematic.	
	magnifying lens, commonly known as <u>ski light</u> is used to	<u>Display</u> this instrument. <u>Demonstrate</u> its use.	
	provide a magnified visual examination.	Solicit students' comments and questions concerning examination for injection sites.	
	5. During this step, the third pulse is taken.	•	
	J. Suspect Statements		
10 Minutes	All spontaneous statements and suspect's response to questions should be documented. Ask additional probing questions as	Note: Give specific examples of probing questions, admissions and denials.	
	appropriate.	Ask students for additional examples and list all on chalkboard or flip chart.	
	K. Opinion of Evaluator		
20 Minutes	1. By this point in the evaluation, the DRE should have formed an opinion of the category or categories of drugs responsible for any observed impairment.		
	2. This opinion is based on the totality of the investigation.		

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Instructor Notes



20 Minutes

L. Toxicological Examination.

1. Toxicology Samples

Your State's implied consent statues will dictate the type of sample you can obtain; urine, blood, breath or saliva.

2. Specimen Containers

- a. The type of container for collecting the sample will be dictated by the type of sample taken and the laboratory requirements where it will be tested.
- b. Containers should be sterile and have a lid that will seal tightly. Make sure the seal is tight to prevent leakage.
- c. Containers will differ depending on the type of specimen collected.
 Containers are uniquely designed to accommodate specific samples such as blood, urine, saliva, breath, etc.

Review the students' department's policy and procedures for requesting, obtaining and handling toxicological samples.

Ask the students to relate the laws of their state. The implied consent laws may vary significantly from state to state.

Have the students discuss their individual laws and possibly write their requirements on the flip chart for comparison.

3. Obtaining a Sample

- a. Urine An officer must witness the drawing of the sample.
- b. Blood Should be drawn by a qualified technician and witnessed by the officer.

The sample must include a preservative. This is often pre-packaged in the container intended for this use.

Samples should be refrigerated or frozen as soon as possible to minimize degeneration during storage.

4. Chain of Custody

- a. Establish a policy dictating the chain of custody, if one does not already exist.
- b. Establish a policy for your Department on:

The sealing of evidence to include officer identification markings; (i.e. initials, labels, tags and packaging)

Paperwork for the chain of custody and laboratory analysis of your sample.

Note: If possible, discourage the use of the mail for delivering the sample to the lab.

Aides	Lesson Plan	Instructor Notes
	Transportation of the sample to the laboratory. Return reporting of the laboratory analysis.	Note: These are issues that must be addressed with the individual agencies to insure proper and standardized procedures. Students should follow-up with the appropriate representatives from their agencies to coordinate this activity.
		Solicit students' comments and questions concerning toxicological examinations.
15 Minutes	M. Video Demonstration	Instruct students to refer to their checklists as they watch the video.
		Show the Video Tape "Overview of DRE Procedures". (This is the same video that is shown during Session II of the PRE-School and subsequently in Session VIII of this school.)
		Solicit students' comments and questions.

Session IV

Overview of Drug Recognition Expert Procedures



Overview of Drug Recognition Expert Procedures

Upon successfully completing this session, the participant will be able to:

- Name the components of the drug evaluation and classification process
- · State the purpose of each component
- Describe the activities performed during each component
- Correctly answer the "Topics for Study" questions at the end of this section

Drug Evaluation & Classification Training

N/O

The Drug Evaluation:

A standardized and systematic process

Drug Evakuation & Classification Training

IV-1

Drug Evaluation Steps

1. The breath alcohol test

Drug Evaluation & Classification Training

IV-2

Drug Evaluation Steps (continued)

2. Interview of the arresting officer



Drug Evaluation & Classification Training

IV-3

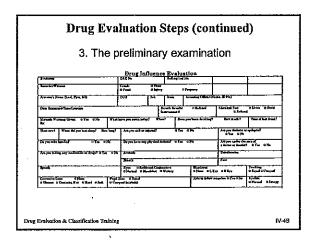
Drug Evaluation Steps (continued)

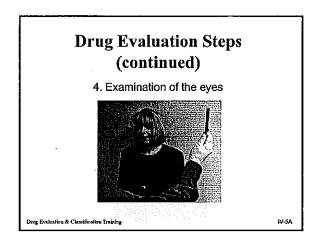
3. The preliminary examination

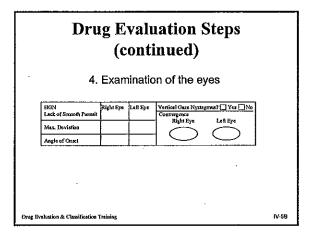


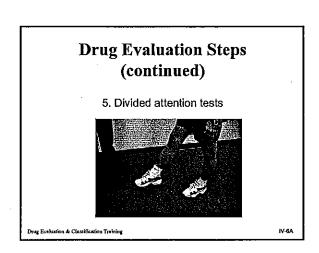
Drug Evaluation & Classification Training

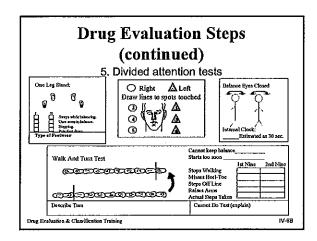
IV-4A

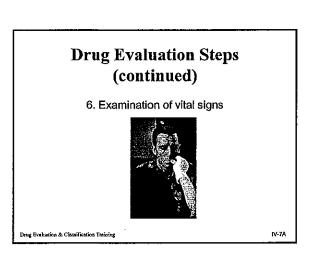


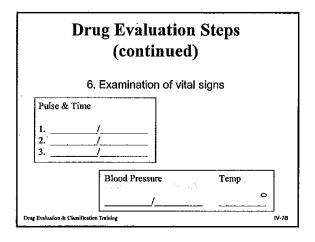




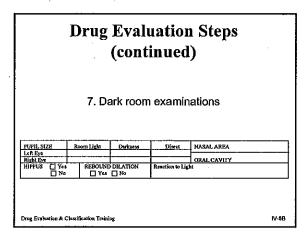


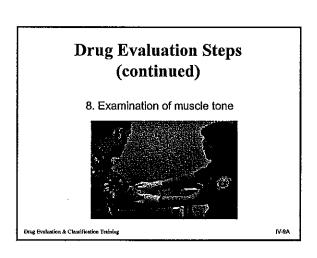




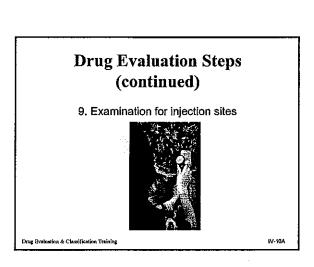


Drug Evaluation Steps (continued) 7. Dark room examinations Drug Evaluation & Classification Training 1148A



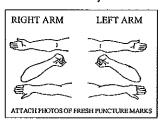


Drug Evaluation Steps (continued)			
8. Examinatio	on of muscle	tone	
MUSCLE TONE:			
Near Normal	Flaccid	Rigid	
Comments:			
Drug Evaluation & Classification Training			IV-9B



Drug Evaluation Steps (continued)

9. Examination for injection sites



Drug Evaluation & Classification Training

IV-10B

Drug Evaluation Steps (continued)

10. Suspect's statements and other observations



Drug Evaluation & Classification Training

IV-11A

Drug Evaluation Steps (continued)

10. Suspect's statements and other observations

io. Suspects s	laternerits (and our	ei onzeiva	uons
What medicine or drug have you been using? How much?		Time of use?	Where were the drugs used? (Location)	
Date/Time of Arrest	Time DRE Notified		Eval Start Time	Time Completed
Member Signature (Include Rank)	ID No.	Reviewe	d By	
Opinion of Evaluator; Rule Out	Akohol Depressant	Stimulant	PCP Narcolic Anal	□lnhalant Igesic □ Cannabis
1.8			1.04	

Drug Evaluation Steps (continued)

11. The opinion of the evaluator



Drug Evaluation & Classification Training

IV-12

Drug Evaluation Steps (continued)

12. The toxicological examination



Drug Evaluation & Classification Training

IV-13

Interview of Arresting Officer: Issues Concerning Suspect's Behavior

- · Was suspect operating a vehicle?
- What actions, maneuvers, etc. were observed?
- · Was there a collision?
- · Was suspect observed smoking, drinking or eating?
- · Was suspect inhaling any substance?
- · How did suspect respond to stop command?
- · Did suspect try to conceal or throw away any items?
- · What has been suspect's attitude and demeanor?

Drug Evaluation & Classification Training

IV-14A

Interview of Arresting Officer: Suspect's Statements

- · Has suspect complained of illness or injury?
- Has suspect used drug-related "street terms" or slang?
- · How has suspect responded to questions?
- Is suspect's speech slurred, slow, thick, rapid, mumbled, etc.?
- · What, specifically, has suspect said?

Drug Evaluation & Classification Training

IV-148

Interview of Arresting Officer: Physical Evidence

- What items or materials were uncovered during search of suspect and vehicle?
- · Was any smoking paraphernalia uncovered?
- Were there any injection materials (e.g., needles, syringes, leather straps, rubber tubes, spoons, bottle caps, etc.)?
- Were there any balloons, plastic bags, small metal foil wrappings, etc.?
- · What was the suspect's BAC?

rug Evaluation & Classification Training

IV-140

Overview of the **Preliminary Examination**



- · Questions
- · Observations of face, breath and speech
- · Initial checks of the eyes
- · The first check of the pulse

Drug Evaluation & Classification Training

IV-19

Preliminary Examination Questions

- · Are you sick or injured?
- · Do you have any physical defects?
- · Are you diabetic or epileptic?
- · Do you take insulin?
- · Are you under a doctor's or dentist's care?
- · Are you taking medication?

Drug Evaluation & Classification Training

IV.s

Initial Checks of the Eyes

- · Check pupil size
- · Assessment of tracking ability
- Initial estimate of nystagmus angle of onset



Drug Evaluation & Classification Training

IV-17

Eye Examinations



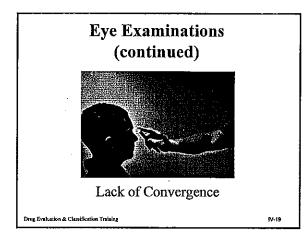
Horizontal Gaze Nystagmus

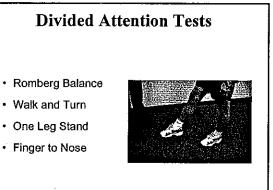


Vertical Gaze Nystagmus

Drug Evaluation & Classification Training

IV-1





Vital Signs Measurements

- · Blood pressure
- Pulse
- Temperature



Drug Evaluation & Classification Training

FV-21

Dark Room Checks of Pupil Size

- Room light
 - Near-total darkness
 - · Direct light

Drug Evaluation & Classification Training

Drug Evaluation & Classification Training

IV-22



Drug Influence Evaluation 000306 Rolling Log No. Evaluator □ None Recorder/Witness Crash: □ Fatal □ Injury □ Property stee's Name (Last, First, MI) DOB Arresting Officer (Name, ID No.) Sex Race Date Examined/Time/Location Breath Results: □ Refused Chemical Test □ Urine □ Blood □ Refused Instrument # Miranda Warning Given: □ Yes □ No What have you eaten today? When? Have you been drinking? How much? Time of last drink? When did you last sleep? Are you sick or injured? □ Yes Are you diabetic or epileptic? Time now? How long? □ No □ Yes □ No Are you under the care of □ Yes □ No Do you have any physical defects?

Yes Do you take insulin? □ No a doctor or dentist? □ Yes □ No Coordination Are you taking any medication or drugs? □ Yes □ No Attitude Breath Face Speech ☐ Reddened Conjunctiva Blindness: Tracking: □ Equal □ Unequal □ Normal □ Bloodshot □ Watery □ None □ L Eye □ R Eye Able to follow stimulus: ☐ Yes ☐ No Corrective Lens: □ None Pupil Size: □ Equal Eyelids: □ Hard □ Soft □ Unequal (explain) □ Normal □ Droopy □ Glasses □ Contacts, if so One Leg Stand HGN Left Eye Right Eye Vertical Nystagmus?

Yes □ No Pulse & Time Lack of Smooth Pursuit Convergence Right Eye Left Eye Max. Deviation Angle of Onset Romberg Balance Walk and Turn Test Cannot Keep Balance_ Starts Too Soon Approx. Арргох. 1st Nine 2nd Nine Sways While Balancing Stops Walking Misses Heel-Toe ☐ Uses Arms to Balance Steps Off Line □ Hopping Raises Arms Puts Foot Down Actual Steps Taken Internal Clock Describe Turn Cannot Do Test (Explain) Type of Footwear _Estimated At 30 Sec. Pupil Size Room Light Darkness Direct Nasal Area ○ Right △ Left Left Eye Draw lines from spot touched Oral Cavity Right Eye Rebound Dilation Hippus □ Yes Reaction To Light □ No □ No □ Yes Blood Pressure Temp Muscle Tone: □ Normal □ Flaccid □ Rigid Attach Photos Of Fresh Puncture Marks Comments: How Much? Where Were The Drugs Used? (Location) t Medicine or Drug Have You Been Using? Time of Use? Time DRE Notified Date/Time of Arrest Eval Start Time Time Completed Reviewed By: Member Signature (Include Rank) ID No. п РСР □ Rule Out □ Inhalant Opinion of Evaluator: □ Alcohol □ Stimulant □ Medical □ Cannabis □ Depressant □ Hallucinogen □ Narcotic Analgesic

000307	•	
	DRUG INFLUENCE EVALUATION	Page <u> 2</u> of <u>2</u>
LOG NO.	DRE:	ARRESTEE:
5. INITIAL OBSERVAT	IESS 3. BREATH TEST 4. NOTIFICATION / IONS 6. MEDICAL PROBLEMS 7. PSYCHO N 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS
1. LOCATION:		
2. WITNESS:	·	
3. BREATH TEST:		
4. NOTIFICATION/	INTERVIEW of ARRESTING OFFICER:	
5. INITIAL OBSERV	ATIONS:	•
6. MEDICAL PROB	LEMS:	
7. PSYCHOPHYSIC	AL TESTS:	
8. CLINICAL INDIC	ATORS:	
9. SIGNS of INGEST	TION:	
10. STATEMENTS:		·
11. OPINION of EVA	LUATOR:	
12. TOXICOLOGICA	AL SAMPLE:	
13. MISCELLANEO	US:	

One Hour and Forty-Five Minutes

SESSION V

EYE EXAMINATIONS: NYSTAGMUS, CONVERGENCE, PUPIL SIZE AND REACTION TO LIGHT

<u>SESSION V</u> EYE EXAMINATIONS: NYSTAGMUS, CONVERGENCE, PUPIL SIZE AND REACTION TO LIGHT

Upon successfully completing this session, the participant will be able to:

- o State the purposes of various eye examinations in the Drug Evaluation and Classification Process.
- o Describe the administrative procedures for the eye examinations.
- o Describe the clues of interest in each eye examination.
- o Conduct the eye examinations and note the clues that come to light.
- o Prepare complete, clear and accurate records of the eye examinations.

Content Segments		<u>Lear</u>	Learning Activities	
A.	Purpose of the Examinations	O	Instructor Led Presentations	
В.	Procedures and Clues	0	Instructor Led Demonstrations	
C.	Demonstrations	0	Student Led Demonstrations	
D.	Documentation Procedures	0	Students' Hands On Practice	
E.	Practice	0	Reading Assignments	

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Instructor Notes





V-0A&B (Session Objectives)



15 Minutes



V-1 ("The Eye Examinaions")

EYE EXAMINATIONS

A. Purposes of the Eye Examinations

- 1. The principal purpose of all of the eye examinations is to obtain articulable facts indicating the presence or absence of specific categories of drugs.
 - a. Certain drug categories usually cause the eyes to react in specific ways.
 - b. Other drug categories usually do not cause those reactions.
- 2. The tests of <u>Horizontal and Vertical Gaze Nystagmus</u> provide important indicators of the drug categories that may or may not be present.
 - a. If Horizontal Gaze
 Nystagmus is observed, it is
 likely that the suspect may
 have taken alcohol or
 another CNS depressant,
 PCP, an inhalant, or a
 combination of those.

Total Lesson Time: Approximately 105 Minutes

Session title on wall chart.

Briefly review the content, objectives and activities of this session.

Ask students "What causes Horizontal Gaze Nystagmus?" Alcohol and certain other drugs will cause Horizontal Gaze Nystagmus.

- b. If Vertical Gaze Nystagmus is observed, the implication may be that the suspect took PCP, or fairly large doses of alcohol or other Depressants or Inhalants.
- c. By comparing the suspect's blood alcohol concentration with the angle of onset of Horizontal Gaze Nystagmus, it may be possible to determine that alcohol is or is not the sole cause of the observed Nystagmus.
- d. The consistency of onset angle and BAC can be compared using the following formula:

BAC = 50 - A

unlikely that a suspect would exhibit Vertical Gaze
Nystagmus without also exhibiting HGN.

Point out that it is very

Clarification: If the onset angle is significantly inconsistent with the BAC, the implication may be that the suspect has also taken PCP, an inhalant, or some CNS Depressant other than alcohol.

Write the formula on the chalkboard or flip-chart.

Note: Emphasize that this is not an absolute mathematical formula.

Explanation:

BAC = $100 \times \text{blood alcohol}$ (i.e., if blood alcohol is 0.10, BAC = 10)

A = onset angle (in degrees)

Example: If onset angle is 35 degrees, then BAC = 50 - 35 = 15.

The corresponding blood alcohol concentration would be approximately 0.15.

Emphasize this point: The formula can easily be "off" by 0.05 or more, even though the subject has consumed no drug other than alcohol.



e. Keep in mind that this formula is only a statistical approximation. It is not an exact relationship for all subjects at all times.

Instructor Notes

- f. The purpose of comparing BAC and onset angle is to obtain a gross indication of the possible presence of another CNS Depressant, PCP, or an Inhalant.
- 3. The check for <u>Lack of</u>
 <u>Convergence</u> can provide
 another clue as to the possible
 presence of Depressants, PCP,
 or Inhalants.
- 4. Lack of Convergence is also an indicator of the possible presence of Cannabis.
- 5. The checks of <u>pupil size</u> and <u>reaction to light</u> provide useful indicators of the possible presence of many drug categories.
 - a. CNS Depressants, CNS
 Stimulants and Narcotic
 Analgesics will usually
 cause the pupils to react
 very slowly or not visibly at
 all to light.
 - b. CNS Stimulants and Hallucinogens usually will cause the pupils to dilate.
 - c. Cannabis usually causes dilation of the pupils, although this isn't always observed.
 - d. Some specific Inhalants may cause pupil dilation.
 - e. Narcotic Analgesics will usually cause observable constriction of the pupils.

Emphasize that many other facts will also be considered that will help to determine whether PCP, inhalants or CNS Depressants may be present.

Point out that a Drug Recognition Expert might begin to suspect the presence of cannabis if Lack of Convergence was observed but <u>no</u> nystagmus was observed.

Point out that pupil dilation due to cannabis isn't always observed in laboratory studies, but this may be due to the fact that laboratory dose levels are less than "street" doses.

Aides

Lesson Plan

Instructor Notes





V-2 "Hippus And Rebound"

- 6. You will also check for <u>hippus</u> and <u>rebound dilation</u>.
 - a. "Hippus" means a rhythmic pulsating of the pupils as they dilate and constrict within fixed limits.
 - b. "Rebound dilation" is a period of constriction followed by dilation with a change equal to or greater than 2 mm the final size determination being estimated at the end of a 15-second time period in which the light from the penlight is directed into the eye.
 - c. Hippus occurs under various conditions, including -- at times -- withdrawal from Narcotic Analgesics.
 - d. Rebound dilation has been reported with persons under the influence of Cannabis.
- B. Procedures and Cues
 - 1. Horizontal Gaze Nystagmus test consists of <u>three separate</u> <u>checks</u>, administered independently to each eye.
 - a. The first check is for "lack of smooth pursuit".
 - o If the subject is wearing eyeglasses, have him or her remove them.

Print on chalkboard:
"HIPPUS"
"REBOUND DILATION".

Note: Instructors are encouraged to use additional visual aides to demonstrate if necessary (i.e., balloon, videos, etc.).

Point out that these terms are defined in the glossary at the front of the Student's Manual.

Solicit students' comments and questions concerning the purposes of the eye examinations.

Select a student, and demonstrate the first check of HGN on that student.



50 Minutes



V-3 ("HGN Procedures and Cues")



Pursuit")

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Instructor Notes

- o If the subject is wearing contact lenses, note that fact on the report, but don't have the subject remove them.
- o Position the stimulus about 12 -15 inches in front of subject's nose.
- o Hold the tip of the stimulus slightly above the level of the subject's eye.
- o Instruct the subject to hold the head still and follow the stimulus with the eyes.
- o Move the stimulus smoothly, all the way to the subject's left side and back all the way to the right side.
- o Make at least two complete passes of the stimulus: to the left side, to the right side, back to the left side, and finally back to the right side.
- o When doing this, don't pause at the center of the subject's face; move all the way to the left, then all the way to the right, then again all the way to the left and back all the way to the right, in a smooth, continuous fashion.

<u>Point Out</u> that this procedure ensures that the subject's eyes will be wide open and easy to observe.

Point out that the stimulus should be moved at a speed that requires 2 seconds to bring it from the center out all the way to the side. It should then be moved from side to side at the same speed. This means it should take 4 seconds to move from the extreme left to the extreme right.

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Instructor Notes

b. While the eyeball is moving, examine it for evidence of a lack of smooth pursuit.

Use these or similar analogies:

- (1)A smoothly pursuing eyeball will move without friction, much the way that a windshield wiper glides across the windshield when it is raining steadily. An eyeball showing lack of smooth pursuit will move in a fashion similar to a wiper across a dry windshield.
- (2)A smoothly pursuing eveball will roll in the socket the way that a marble or ball bearing would glide smoothly across a polished pane of glass. An eyeball exhibiting lack of smooth pursuit would move more like that marble rolling over a sheet of heavy gauge sandpaper.

Also, check to be sure that both eyes are tracking in the same way: if one eye is moving smoothly but the other moves hesitantly or not at all, an illness or injury may be present.

Excuse the student volunteer and thank him or her for participating.

d. Students' initial practice of the check for lack of smooth pursuit.

Instruct students to work in pairs, taking turns checking each other's eyes for lack of smooth pursuit.

Monitor, coach and critique the students' practice.

Allow this practice to continue for only about 2 minutes.

Aides

Lesson Plan

Instructor Notes



V-4B ("Distinct...At Maximum")

- e. The second check is for "distinct nystagmus at maximum deviation".
 - o Again position the stimulus as before.
 - o Move the stimulus all the way to the subject's left side and hold it there so that the subject's eye is turned as far to the side as possible.
 - o Hold the eyeball at that position for a minimum of 4 seconds, to check carefully for any jerking that may be present.
 - o When you have completed this check for the left eye, repeat the process for the right eye. Then, do it once again for the left eye, and again for the right, to verify that distinct nystagmus is present.
- f. With this cue, the examiner looks for a <u>very distinct</u>, unmistakable jerking.
 - o A slight or barely visible tremor is not sufficient to consider this cue present.
 - o A definite, sustained jerking must be seen.

Select a student and demonstrate the second check of HGN on that student.

Point out that people exhibit slight jerking of the eye at maximum deviation, even when unimpaired, but this will not be evident or sustained for more than a few seconds. When impaired by alcohol and "D.I.P." drugs, the jerking will be larger, more pronounced, sustained for more than 4 seconds, and easily observable.

Excuse the student volunteer and thank him or her for participating.

Instructor Notes



g. Students' initial practice of the check for distinct nystagmus at maximum deviation. Instruct students to work in pairs, taking turns checking each other's eyes for distinct nystagmus at maximum deviation.

Monitor, coach and critique the students' practice.

Allow this practice to continue for only about 2 minutes.

- h. The final check is for the "angle of onset".
 - o Position the stimulus as before.
 - o Slowly move the stimulus to the subject's left side, carefully watching the eye for the first sign of jerking.
 - o When you think that you see the eyeball jerk, stop moving the stimulus and hold it perfectly still.
 - o Verify that the eyeball is, in fact, jerking.
 - o Once you have established that you have located the point of onset, estimate the angle.
 - o Then, repeat the process for the right eye.

Select a student and demonstrate the third check of HGN on that student.

Note: Stimulus should be moved at a speed that requires four seconds to travel from center all the way out to the side.

Point out that, if the eye is not jerking, it will be necessary to resume moving the stimulus slowly to the side, again observing for the first sign of jerking.

<u>Point out</u> that angle estimation simply requires practice.

Aides	Lesson Plan	Instructor Notes
	o Then, again check onset for the left eye, and again for the right.	
		Exhibit a template.
		Point out that the template will be used during practice.
		Excuse the student volunteer and thank them for participating.
	·	Emphasize that if the clues of Horizontal Gaze Nystagmus are markedly different for the two eyes, a neurological or other medical problem (such as a head injury) may be present.
	i. Students' initial practice of angle estimation.	Instruct students to work in pairs, taking turns estimating angles of each other's eyes.
		Instruct students that they are to try to draw their partners' eyes to three different angles: 30 degrees 35 degrees 40 degrees
		Students will check their accuracy using the template.
	·	Monitor, coach and critique the students' practice.
		Allow this practice to continue for only about 3 minutes.
V-5 ("Vertical Gaze Nystagmus")	2. The <u>Vertical Gaze Nystagmus</u> test is very simple, and consists of a single check.	Select a student and demonstrate the Vertical Gaze Nystagmus test on the student.

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Instructor Notes

- a. Position the stimulus <u>horizontally</u>, about 12 -15 inches in front of the subject's nose.
- b. Instruct the subject to hold the head still and follow the object with the eyes only.
- c. Raise the object until the subject's eyes are elevated as far as possible.
- d. Watch closely for evidence of jerking.

<u>Point out</u> that the examiner should keep the subject's eyes elevated for about 4 seconds to verify that the jerking really is present.

Excuse the student volunteer and thank them for participating.

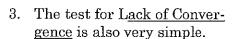
Instruct students to work in pairs, taking turns administering the Vertical Gaze Nystagmus test to each other.

Monitor, coach and critique the students' practice.

Allow this practice to continue for only about 2 minutes.

<u>Select</u> a student and demonstrate the test for Lack of Convergence on that student.

e. Students' initial practice of the Vertical Gaze Nystagmus test.



a. Position the stimulus about 12-15 inches in front of the student's face, with the stimulus pointing toward the nose.



V-6 (Lack of Convergence)

Aides	*	Lesson Plan	Instructor Notes
	b.	Instruct the subject to hold the head still and follow the object with the eyes only.	
	c.	Keep the object 12-15 inches away from the subject's nose, and start to move the object slowly in a circle, approximately the same size as the suspect's face.	
			Point out that this initial circular motion helps to verify that the subject has focused on the stimulus and is able to track it. Emphasize that it doesn't matter whether the circular motion is clockwise or counter-clockwise.
	d.	Once you have verified that the subject is tracking the object, move it slowly and steadily and touch the bridge of the nose.	Note: Hold stimulus on bridge of nose for one (1) second.
	e.	Carefully observe the subject's eyes to determine whether both eyes converge on the bridge of the nose.	Excuse the student volunteer and thank them for participating.
	f.	Students' initial practice of the test for Lack of Convergence.	Instruct students to work in pairs, taking turns testing each other's eyes for Lack of Convergence.
		·	Monitor, coach and critique the students' practice.

Allow this practice to continue for only about 2 minutes.

Aides

Lesson Plan

Instructor Notes



V-7 ("Pupil Size")

- 4. Estimation of <u>pupil size</u> requires use of the pupillometer.
 - a. For the check in room light. Hold the pupillometer alongside the subject's eye. Instruct the subject to focus on a specific point behind the DRE and slightly above the subject's eye level. Utilize the same point for the dark room examinations.

- b. Make sure that the pupillometer is even with the eyeball (neither closer to you nor farther from you than is the subject's eyeball).
- c. Move the pupillometer up or down until you find the darkened circle that appears to be approximately the same size as the subject's pupil. Check the left eye and then the right eye.
- d. Students' initial practice of pupil size estimation.

Exhibit a pupillometer.

Write on flipchart or chalkboard "The Three Lighting Conditions".

<u>Select</u> a student and demonstrate pupil size estimation using the student.

Explain to the students that the pupils will automatically constrict as objects move closer to them. This is called accommodation reflex.

This should not be confused with pupillary light reflex which is the pupil's normal reaction to changes in light.

Demonstrate the accommodation reflex by having the students focus on an object very close and one at a distance.

Point out that the "normal" size of a pupil is about 3.0 - 6.5 mm.

Excuse the student-volunteer and thank him or her for participating.

<u>Instruct</u> students to work in pairs, taking turns checking each other's pupils.

Aides	Lesson Plan	Instructor Notes
		Monitor, coach and critique the students' practice. Allow this practice to continue for only about 2 minutes.
	e. After you have completed the pupil size estimations in room light, you must darken the room, wait 90 seconds, and then proceed with the darkroom exam.	Select a student to participate in demonstrations of darkroom pupil estimations.
	f. For the check under <u>near</u> total darkness, hold your finger over the tip of the penlight, so that only a reddish glow emerges.	<u>Demonstrate</u> this.
	g. For the check under <u>direct</u> <u>light</u> , bring the light from the side of the subject's face, directly into the eye.	<u>Demonstrate</u> this. <u>Emphasize</u> that the penlight
		should be positioned so that the beam just "fits" the eye socket.
	5. Assessment of the pupil's reaction to light takes place immediately before the check of pupil size under direct light.	
	a. Once again, start by bringing the uncovered light from the side of the subject's face directly into his or her left eye.	<u>Demonstrate</u> this.
	b. As you bring the beam of light directly into the subject's eye, note how the pupil reacts.	<u>Demonstrate</u> this.

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Aides		Lesson Plan	Instructor Notes
·	c.	Under ordinary conditions, the pupil should react very quickly, and constrict noticeably when the light beam strikes the eye.	
	d.	Under the influence of certain categories of drugs, the pupil's reaction may be very sluggish, or there may be no visible reaction at all.	Emphasize: We consider the pupil's reaction to be slow if it takes more than one second to reach full constriction.
	e.	Hold the direct light on the subject's eye for <u>15 seconds</u> to assess pupil reaction.	
	f.	Also check for <u>hippus</u> or <u>rebound dilation</u> during this 15 seconds period.	Caution should be used by the officer so as not to move the light beam or allow the bulb to change in light intensity.
	g.	When you have completed this process for the left eye, repeat it for the right eye.	Excuse the student volunteer and thank him or her for participating.
	h.	Students' initial practice in assessing the pupil's reaction to light.	Instruct the students to work in pairs, taking turns shining the light into each other's eye and observing the pupil's reaction.
			Remind students to position the penlight so that the beam exactly "fits" the eye socket when the beam is brought directly into the eye.
			Monitor, coach and critique the students' practice.

Allow the practice to continue for only about 2 minutes.

Solicit students' comments and questions concerning the eye

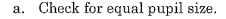
examinations.

Aides	Lesson Plan	Instructor Notes	
	C. Demonstrations		
15 Minutes	1. Demonstration of Horizontal Gaze Nystagmus.	Select two students to come before the class.	
	a. Check for lack of smooth pursuit.	Instruct one student to demontrate the administration of Horizontal Gaze Nystagmus to the other student.	
	b. Check for distinct nystagmus at maximum deviation.	Coach and critique the student administrator's performance.	
	do viation.	Make sure that the student administrator checks both eyes.	
	c. Estimation of onset angle.	When the student administrator has completed the HGN test, instruct the student administrator to draw the student subject's eye to an angle of 35 degrees. Check the accuracy of this estimate, using the template.	
		Excuse the two students and thank them for participating.	
	2. Demonstration of Vertical Gaze Nystagmus and Lack of Convergence.	Select two other students to come before the class.	
	Convergence.	Instruct one student to check the other for Vertical Gaze Nystagmus.	
		Coach and critique the student administrator's performance.	
		Instruct the second student to check the eyes of the first student for Lack of Convergence.	
		Coach and critique the student administrator's performance.	

Aides	Lesson Plan	Instructor Notes
		Excuse the two students and thank them for participating.
	3. Demonstration of pupil size checks and test for reaction to light.	Select two other students to come before the class.
	a. Pupil size estimation under room light.	Instruct one student to check the other's pupils under room light.
		Coach and critique the student administrator's performance.
		Instruct the second student to demonstrate how to perform the dark room checks of pupil size.
	b. Darkroom checks of pupil size.	Coach and critique the student administrator's performance.
	o near total darkness o direct light	Point out that assessment of the pupil's reaction to light takes place in conjunction with the direct light check.
		Excuse the two students and thank them for participating.
	·	Solicit students' comments and questions concerning these demonstrations of the eye examinations.
5 Minutes	D. Documentation Procedures	Instruct students to turn to the Standardized Drug Influence Evaluation Form in their manuals.
	1. A brief examination of the eyes is made during the <u>Preliminary Examination</u> .	

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V-8 (Sample
Eye Data)
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Instructor Notes



- b. Assessment of tracking ability.
- c. Initial assessment of Nystagmus.
- 2. The next section of the Form is devoted to the Eye Examinations.
 - a. Horizontal Gaze Nystagmus
 - b. Vertical Gaze Nystagmus
 - c. Lack of Convergence
- 3. The darkroom eye examinations are documented in a subsequent section of the Form.

<u>Point out</u> that section of the Form.

Emphasize that all three checks of the HGN test must be documented for each eye.

<u>Point out</u> that "yes" implies that Vertical Gaze Nystagmus <u>was</u> observed, "no" implies that it was <u>not</u> observed.

<u>Point out</u> that it will be necessary to diagram the movement of the eyes.

<u>Point out</u> the location of that section.

Emphasize that all darkroom checks of the eyes must be performed and documented independently for each eye.

Solicit students' comments and questions concerning procedures for documenting the eye examinations.

<u>Instruct</u> students to practice in pairs.

Each student will conduct a complete set of eye examinations on his or her partner.



20 Minutes

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E. Practice

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Instructor Notes

- 1. Preliminary eye exams
 - a. Check for equal pupil size.
 - b. Assessment of tracking ability.
 - Initial estimation of Nystagmus onset angle.
- Eye exams.
 - a. Horizontal Gaze Nystagmus
 - b. Vertical Gaze Nystagmus
 - c. Lack of Convergence
- 3. Pupil Size Estimations
 - a. Room light
 - b. Near total darkness
 - c. Direct light
- 4. Reporting out of Pupil Size Estimations.

Students then will "reverse roles".

Tell the students to record their estimations of their partners' pupil sizes on the standard Drug Influence Evaluation Form.

<u>Monitor</u>, coach and critique students' practice.

<u>Make sure</u> each student administers a complete series of eye examinations at least once.

<u>NOTE</u>: If possible, the training room should be at least somewhat darkened for this final stage of practice.

Instructor: While the student's practice is still going on, print the matrix at the end of this session on the chalkboard or flip-chart.

Tell students that they should refer to the Drug Influence Evaluation forms on which they recorded their partners' pupil sizes.

Tell the students that we will tabulate the pupil sizes of everyone in the class, for each of the three lighting conditions.



Aides	Lesson Plan	Instructor Notes
		For simplicity, tell the students that we will tabulate the <u>left</u> <u>eye</u> pupil sizes only.
	a. Room light tabulation.	Direct the students' attention to the first column of the matrix.
		Say: "Let's concentrate now only on the room light estimations."
		Ask: "How many of you found that your partners had pupils of 2.0 mm or less in room light?" (Get a show of hands; count them; print the number in the first box of the first column.
		Then ask: "How many had partners with a 2.5 mm pupil in room light?" (Count the hands and print the number in the 2nd box.)
		Continue this until you get to the last box in the 1st column: "How many had partners with pupils of 7.5 mm or larger?" (Count the hands; print the number.)
	b. Near total darkness tabulation.	Repeat this process for each of the other two lighting conditions.
	c. Direct light tabulation.	Make appropriate comments about the number of students whose pupils are outside the normal range of size under the various lighting levels.

pupil size	room light	near total darkness	direct light
≤ 2.0			
2.5			
3.0			
3.5			
4.0			
4.5			
5.0			,
5.5			
6.0			
6.5			
7.0			
≥ 7.5			

Session V

Eye Examinations





Eye Examinations: Nystagmus, Convergence, Pupil Size, and Reaction to Light

Upon successfully completing this session, the participant will be able to:

- State the purposes of various eye examinations in the Drug Evaluation and Classification process
- Describe the administrative procedures for the eye examinations

Drug Evaluation & Classification Training

V.OA

Eye Examinations: Nystagmus, Convergence, Pupil Size, and Reaction to Light (continued)

- Describe the clues of interest in each eye examination
- Conduct the eye examinations and note the clues that come to light
- Prepare complete, clear and accurate records of the eye examinations

Drug Evaluation & Classification Training

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The Eye Examinations





Drug Evaluation & Classification Training

V-1

"Hippus" and "Rebound"

- Hippus a rhythmic pulsating of the pupils as they dilate and constrict within fixed limits.
- Rebound is a period of constriction followed by dilation with a change equal to or greater than 2 mm.

Drug Evaluation & Classification Training

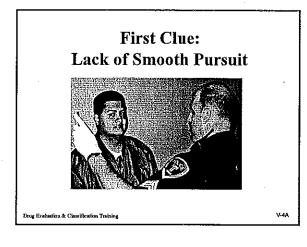
V-2

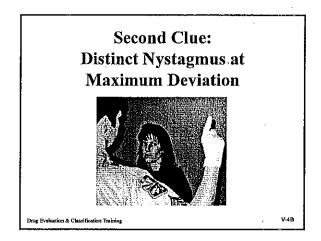
Three Clues of Horizontal Gaze Nystagmus

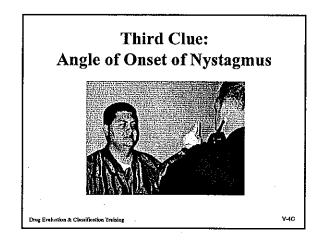
- 1. Lack of smooth pursuit
- 2. Distinct nystagmus at maximum deviation
- 3. Angle of onset of nystagmus

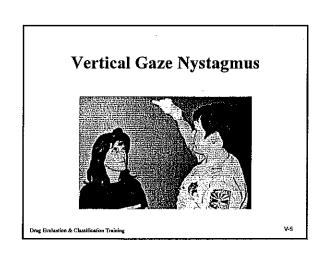
Drug Evaluation & Classification Training

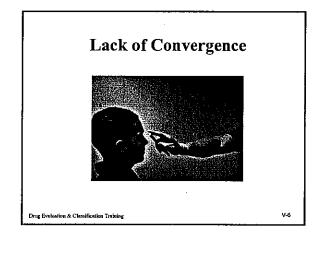
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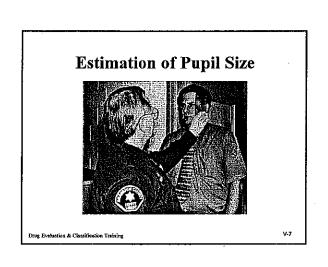




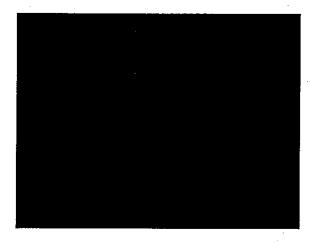








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Two Hours

SESSION VI

 $\begin{array}{c} {\bf PHYSIOLOGY\ AND\ DRUGS:} \\ {\bf AN\ OVERVIEW} \end{array}$

SESSION VI PHYSIOLOGY AND DRUGS: AN OVERVIEW

Upon successfully completing this session, the participant will be able to:

- o Explain in layman's terms the general concept of human physiology.
- o Explain in layman's terms the purpose and functions of major systems in the body (nervous system, circulatory system, respiratory system, etc.).
- o Explain in layman's terms how drugs work in the body.
- o Explain in general terms how the drug evaluation is used to detect signs or symptoms indicative of drug impairment.
- o Correctly answer the "topics for study" questions at the end of this Section.

Content Segments

A. Body Systems

- B. Body Systems and Body Functions Relevant to Drug Evaluations
- C. How Drugs Work
- D. Physiologic Signs and Symptoms of Drugs or Medical Impairment
- E. Medical Conditions
- F. Summary

Learning Activities

- o Instructor Led Presentations
- o Reading Assignments

Aides

Lesson Plan

Instructor Notes



VI-0A&B (Session Objectives)

PHYSIOLOGY AND DRUGS: AN OVERVIEW

Total Lesson Time: Approximately 120 Minutes

Briefly review the content, objectives and activities of this session.



5 Minutes

A. Introduction

1. Before we can understand how drugs work we must have a basic understanding of how the body works.

Point out that it is not necessary to have detailed knowledge of specific functions or medical terminology. Students will not become medical specialists as a result of this limited overview, however, they should be encouraged to learn as much as possible about human physiology through additional instruction and independent reading.

2. We will review general concepts of how the body functions in a "normal" or "standard" human.

Point out that all human beings are different and a "normal" or "standard" human does not exist. However, experience has produced a range of normal values that can be used for comparison purposes.



VI-1

15 Minutes

- 3. We will briefly review the chief functions of the body systems.
- 4. Primary focus will be on the systems or component parts of those systems that are examined during the drug evaluation.
- B. Human Physiology

- o Central Nervous System
- o The Eyes
- o Blood Pressure and Pulse
- o Body Temperature
- o Balance and Coordination

Instructor Notes



VI-2 Definition



VI-3A ("Murders, Inc.")



VI-3B ("The Ten Systems")

- 1. Physiology is the study of the functions of living organisms and their parts.
- 2. A convenient way of discussing human physiology is to list the ten major systems of the body.
 - a. The phrase "MURDERS, INC." helps us remember the names of the ten systems.
 - b. Each letter stands for the name of one system.
- 3. M stands for the MUSCULAR SYSTEM.
 - a. The body has three different kinds of muscles.
 - (1) the heart, or cardiac muscle.
 - (2) smooth muscles, which control the body's involuntary operations.
 - (3) striated muscles, which carry out our voluntary movements.
 - All three types of muscles are examined at various stages of the drug influence evaluation.

Selectively reveal the systems as you discuss each of them.

<u>Point out</u> that we assess the muscular system in the drug influence evaluation when we test coordination and balance by administering divided attention tests, and when we check for muscle rigidity.

Examples: Smooth muscles control breathing, the operation of the pyloric valve (a muscle located at the base of the stomach), dilation and constriction of the pupils, and all other things that we do not consciously control.

- 4. U is for the URINARY SYSTEM.
 - a. The system consists of two kidneys, the bladder, ureters connecting the kidneys to the bladder, and the urethra, which transports the urine out of the body.
 - b. Kidneys filter waste or harmful products, such as drugs and their metabolites, from the blood, and dump these waste products into the bladder.
- 5. The first R in "MURDERS, INC." stands for the RESPIRATORY SYSTEM.
 - a. The major parts of the Respiratory System are the lungs and the diaphragm.
 - b. The diaphragm is a smooth muscle that draws the air into the lungs and forces it out.
 - c. Lungs take in oxygen and transfer it to the blood, and remove carbon dioxide and some other waste products from the blood, and expel them into the outside air.
- 6. D stands for the DIGESTIVE SYSTEM.
 - a. Major components of this system are the tongue, teeth, esophagus, stomach, intestines, liver and pancreas.

Point out that drugs can usually be detected in the urine, and that collection of a urine specimen or other suitable bodily substance is an important part of the drug influence evaluation.

<u>Point out</u> that some drugs cause the user to breathe slowly and shallowly, while others cause rapid breathing.

<u>Point out</u> that important clues of drug use, i.e., odors of alcohol beverages, marijuana, chemicals, etc. may be present on a suspect's breath.

Aides

Lesson Plan

Instructor Notes

- b. The Digestive System breaks down large particles of food, until they are of a size and chemical composition that can be absorbed in the blood.
- Remind students that, when drugs are taken orally, they might be retained in the stomach for a while, until any food that is there has been broken down sufficiently to allow passage into the small intestine.
- E is for the ENDOCRINE SYSTEM.
 - The Endocrine system is made up of a number of different glands, that secrete hormones.
- INSTRUCTOR, FOR YOUR INFORMATION: The glands that make up the Endocrine System include the Thyroid, Parathyroid, Pituitary and Adrenal glands, as well as portions of the pancreas, testes and ovaries.
- b. Hormones are complex chemicals that travel through the blood stream and that control or regulate certain body processes.
- Print HORMONES on the chalkboard or flip-chart.
- Some drugs can mimic the effects of certain hormones. or can react with the hormones in ways that alter the hormones' effects.
- INC." stands for the REPRODUCTIVE SYSTEM.

The second R in "MURDERS.

The functions of the reproductive system fall into two categories: 1) self-producing (cytogenic), and 2) hormoneproducing (endocrinic). We are primarily concerned with hormone production since the hormones produced by the reproductive system aid the nervous system in its regulatory role.



- 9. S is for the SKELETAL SYSTEM.
 - a. Consists of bones, cartilage and ligaments.
 - b. The Skeletal System provides support to the body, permits movement, and forms blood cells.
- 10. The I in "INC" stands for the INTEGUMENTARY SYSTEM.
 - a. Consists of the skin, hair, finger and toe nails, and accessory structures.
 - b. The chief functions of the Integumentary System include protection of the body, control of body temperature, excretion of wastes (i.e., through the sweat) and sensory perception.
- 11. N is for the NERVOUS SYSTEM.
 - a. This system consists of the brain, the brain stem, the spinal cord and the nerves.
 - b. Nerves keep the brain informed of changes in the body's external and internal environments.

<u>Point out</u> that the Reproductive and Skeletal Systems are the only major components of physiology and that are not directly involved in the drug influence evaluation.

<u>Point out</u> that DREs examine the skin for hypodermic injection sites, and for sweating, clamminess, and temperature.

EMPHASIZE that the Nervous System is one of the most important components of physiology, as far as the drug influence evaluation is concerned.

CLARIFICATION: Nerves carry messages to the brain from the sense organs (eyes, ears, nose, etc., and also from pain sensors).

Instructor Notes

- Nerves also carry messages from the brain to the body's muscles, tissues and organs.
- d. The nervous system controls, coordinates and integrates all physiological processes, so that normal body functions can be maintained.
- 12. C is for the CIRCULATORY SYSTEM.
 - a. For our purposes, the most important parts of the Circulatory System are the heart, the blood vessels (e.g., arteries, veins, capillaries, etc.) and the blood.
 - b. Blood is the body's primary transport mechanism: it carries food, water, oxygen, hormones, antibodies, etc. to the body's tissues and organs.
 - c. Blood is also primarily responsible for carrying heat throughout the body.
 - d. And, blood is the main transport mechanism for bringing drugs to the brain.
 - e. The heart, of course, pumps the blood, and causes it to circulate through the body.

CLARIFICATION: The brain uses nerves to send messages commanding the heart to beat, the fingers to move, the pupils to dilate, etc.

<u>Point out</u> that this is another very important component of physiology, as far as the drug influence evaluation is concerned.

Solicit students' comments and questions about "MURDERS, INC", the ten major systems of human physiology. Point out that much more will be said about the last two systems (Nervous and Circulatory) later in this session.



VI-3C (Interrelated Body Systems)

Instructor Notes



VI-3D (Homeostasis).

13. Homeostasis

- a. Human body is exposed to constantly changing external environment.
- b. Changes are neutralized by the <u>internal</u> environment the blood.
- c. Oxygen, foods, water and other substances are constantly leaving body fluids to enter cells, while carbon dioxide and other wastes are leaving the cells to enter these fluids...
- d. Yet, the chemical composition of these fluids remains within very narrow limits.
- e. This phenomenon is called homeostasis.

Homeostasis is the dynamic balance, or steady state, involving levels of salts, water, sugars and other materials in the body's fluids.

Point out that the rhythm of the heart, breathing, constancy of body temperature, and the steady level of blood pressure under specific circumstances or conditions are all manifestations of homeostatic mechanisms at work within the body.

Drugs interfere with the homeostatic mechanisms and produce signs and symptoms that can be recognized by a trained DRE.



C. Major Systems and Body Functions of Concern in Drug Evaluations

1. Heart and circulatory system.

Aides

Lesson Plan

Instructor Notes



VI-4 Basic plan of the circulatory system.

- Circulation is a closed system, round which blood is propelled by contractions of the heart.
- b. Blood is driven into arteries, arteries divide into smaller and smaller branches and finally into meshwork of fine capillaries which pervade body tissues.
- c. Meshwork joins up again to form small veins which become larger trunks as they travel centrally towards the heart.
- d. There are two separate circulation systems:
 - (1) A systemic circulation concerned with the body as a whole and driven by the left side of the heart.
 - (2) A pulmonary circulation concerned with passage of blood through the lungs and driven by the right side of the heart.
- The heart acts as two pumps:
 - (1) Left side pumps blood through the aorta and the arteries to the tissues.
 - (2) Blood, after passing through the tissues, returns via the veins to the right side.

Point out that arteries constrict to aid distribution of blood.

Point out that blood does not come into direct contact with the cells, but rather stays in the blood vessels.



VI-5

System

Consists of the left atrium and ventricle. The upper chamber (atrium) receives blood from the great veins, the lower chamber discharges blood into the great arteries.

Aides	Lesson Plan	Instructor Notes
	(3) Right side pumps blood through the pulmonary artery to the lungs and returns it to the left side of the heart again via the four pulmonary veins.	Consists of the right atrium and ventricle. Note: The Pulmonary Artery is the only artery that carries de-oxygenated blood; all other arteries carry blood that has received fresh oxygen from the lungs. Likewise, the Pulmonary Vein is the only vein that carries blood rich in oxygen; all other veins carry blood depleted of oxygen back to the heart.
	f. The normal heart continues to beat regularly and continuously, with a rest interval never longer than a fraction of a second.	
	(1) Heart rate is the number of beats per minute.	Point out that heart rate is regulated by the autonomic nervous system: sympathetic nerve fibers insure that heart beats fast enough to maintain circulation during any activity. Parasympathetic nerve fibers tend to slow the heart. This coordinated nerve supply assures that the heart does not beat too fast or too slowly.
	(2) Pulse rate is the number of pulsations per minute.	For the DRE Program, the normal range is 60-90 pulsations per minute.
		Point out that some people may exhibit <u>irregular</u> (or arrhythmic) heart beats, i.e., where the interval between pulses varies.
VI-6	(3) Blood pressure (BP) is the force of the blood circulating in the arteries.	

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Instructor Notes

- (4) BP is categorized as systolic or diastolic BP.
- (5) Systolic pressure is the maximum force that occurs during contraction.
- (6) Diastolic pressure represents the minimum force that occurs when the heart relaxes.
- (7) Both systolic and diastolic pressures are measured and is recorded as follows:

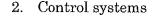
 120 systolic
 80 diastolic

Ask students to define "systolic" and "diastolic".

Point out that physical conditioning can also affect blood pressure and pulse rate.

Demonstrate proper method of recording on flip chart or chalkboard.

Point out that the normal range of BP varies widely based on a number of factors, including age. The normal range of systolic pressure is 120 to 140. The normal range of diastolic is 70 to 90.



- a. The functions of the organs of the body are controlled in two ways:
 - (1) One, by sending
 "chemical messengers"
 known as hormones via
 the blood stream from
 an endocrine gland
 where they are
 produced.
 - (2) Second system of control is by means of the nervous system.

This is a function of the endocrine system.

Remind students that the hormones modify the activity of specific organs.



Aides

Lesson Plan

Instructor Notes

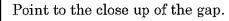


- VI-7 (Nerve Concept)
- b. A Simplified Concept of a <u>nerve</u>.
 - (1) The nerves that carry messages to and from the brain often are pictured as "wires" that carry electrical signals.
 - (2) A more accurate, but still simplified concept would envision a nerve as a series of broken wire segments, with the segments separated by short spaces, or gaps.
 - (3) We can imagine messages running along the "wire segments" in much the same manner that electrical impulses run along telephone wires.
 - (4) When the message reaches the end of the "wire segment", it triggers the release of chemicals that flow across the gap, and contact the next "wire segment".
 - (5) When the chemical contacts the next wire segment, it generates an electrical impulse which runs along the wire until it reaches the next gap.

CLARIFICATION: Nerves are often pictured as telephone or telegraph wires.

Point to a "wire segment".





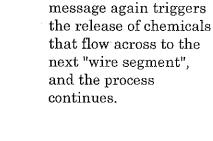


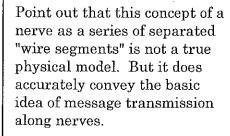


V1-7

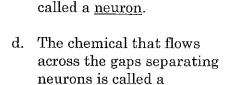
(6) At that gap, the







Solicit students' questions about this concept.



neurotransmitter.

nerves, each "wire segment"

corresponds to a nerve cell,

c. In our simple model of

e. The body has a number of different neurotransmitter; each carries a different chemical message.

f. Each neuron, or "wire segment" has three main parts:

- (1) the cell body.
- (2) the axon.
- (3) the dendrite.
- g. The <u>axon</u> is the part of the neuron that sends out the neurotransmitter, or chemical messenger.
- h. The dendrite is the part that receives the neurotransmitter.
- The gap between two neurons is called a <u>synapse</u>, or <u>synaptic</u> gap.

CLARIFICATION: neurotransmitter are the body's chemical messengers.



Point out the cell body, the axon and the dendrite on VI-8A.

Solicit students' questions about nerve cells (neurons).



VI-9 (Classification of Nerves)

- 3. Classifications of nerves.
 - a. Some nerves carry messages away from the brain, to the body's muscles and organs.
 - (1) These are called <u>Motor</u>, or <u>efferent</u> nerves.
 - (2) The brain uses motor nerves to send commands to the heart to beat, the lungs to breathe, the muscles to contract or expand, and so forth.
 - b. Other nerves carry
 messages to the brain, i.e.,
 from the eyes, ears and
 other senses, from the
 muscles, etc.
 - (1) These are called <u>Sensory</u>, or <u>afferent</u> nerves.
 - (2) The brain decodes the messages that come along the sensory nerves to monitor the condition of the body and of the outside world.
 - c. A Fundamental Notion: If something interferes with the messages the brain sends along the motor nerves, the brain's control over the heart, the lungs, the muscles and other organs will be distorted.

Instructor Notes





- d. Another Fundamental Notion: If something interferes with the messages the brain receives from the sensory nerves, the brain's perception of the outside world and of the body's status will be distorted.
- e. Focus on the <u>Motor</u> nerves. There are two sub-systems of motor nerves.
 - (1) The <u>voluntary</u> nerves send messages to the striated muscles that we consciously control.
 - (2) The <u>autonomic</u> nerves send messages to the muscles and organs that we do not consciously control, i.e., smooth muscle and cardiac muscle.
- f. The <u>Autonomic Sub-system</u> divides into two groups.
 - (1) The <u>Sympathetic</u> nerves command the body to react in response to fear, stress, excitement, etc.

Point out that, basically, this is how drugs work: they interfere with transmission or reception of the messages that travel along nerves.

On the chalkboard, print the word "autonomic", and draw two lines from the word one line angling down toward the left, the other angling down toward the right.

Write "Sympathetic" at the end of one line, "Parasympathetic" at the end of the other.

CLARIFICATION:

<u>Sympathetic</u> nerves control the body's "fight or flight" responses.

EXAMPLES: Sympathetic nerves carry the messages that cause:

- o blood pressure to elevate
- o pupils to dilate
- o sweat glands to activate
- o hair to stand on end
- o heartbeat to increase & strengthen



Aides		Lesson Plan	Instructor Notes
			o blood vessels of the skin to constrict o the walls of the hollow viscera to relax (inhibiting digestion)
		(2) Parasympathetic nerves carry messages that produce relaxed and tranquil activities.	EXAMPLES: Parasympathetic nerves carry messages that cause: o pupils to constrict o heartbeat to slow o peripheral blood vessels to dilate o blood pressure to decrease o digestion to be facilitated
	g.	Certain <u>neurotransmitter</u> (i.e., chemical messengers) aid in the transmission of messages along sympathetic and parasympathetic nerves.	
	h.	Some drugs mimic the action of these neurotransmitters: When taken into the body, these drugs artificially cause the transmission of messages along sympathetic or parasympathetic nerves.	
S S S S S S S S S S S S S S S S S S S	i.	Drugs that mimic the neurotransmitter associated with sympathetic nerves are called sympathomimetic drugs.	Write "Sympathomimetic" on the chalkboard or flip chart.
		(1) Sympathomimetic drugs artificially cause the transmission of messages that produce elevated blood pressure, dilated pupils, etc.	Ask students to name a category of drugs that would be considered sympathomimetic.

Aides	Lesson Plan	Instructor Notes
	(2) Examples: CNS Stimulants, Halluci- nogens, and to some extent PCP and Cannabis.	
	j. Drugs that mimic neuro- transmitters associated with parasympathetic nerves are called <u>para-</u> <u>sympathomimetic</u> drugs.	Write "Parasympathomimetic" on the chalkboard or flip chart.
	(1) Parasympathomimetic drugs artificially cause the transmission of messages that produce lowered blood pressure, drowsiness, etc.	Ask students to name a drug category that would be considered parasympathomimetic.
	(2) Examples: Narcotic Analgesics and CNS Depressants.	
	4. Although there are more than 100 chemicals in the brain, only about two dozen probably are true neurotransmitters.	
	a. Among the primary neurotransmitters that have been identified are:	Write these neurotransmitter on the chalkboard or flip chart.
ansatt Etham	o Norepinephrine (also called Noradrenaline)	Point out that Norepinephrine is a <u>neurotransmitter</u> that produces effects on the body that are similar to the effects produced by Adrenaline (a <u>hormone</u>). Many neurotransmitter correspond to hormones that produce similar effects.
	o Acetylcholine	Acetylcholine plays a role in muscle control, and affects neuromuscular or myoneural junctions.

Aides	Lesson Plan	Instructor Notes
	o Dopamine	Dopamine plays a role in mood control and is used in treating Parkinson Disease.
	o Serotonin	Serotonin is a vasoconstrictor, thought to be involved in sleep, wakefulness and sensory perception. Tryptophan is a precursor to serotonin, and has been used to treat insomnia.
	o Gama Amino Butric Acid (Abbreviated GABA)	GABA inhibits various neuro- transmitter and also causes a release of growth hormones.
	o Endorphins and Enkephalins	These are the body's natural pain relievers.
	b. There are many drugs that artificially induce the effects of neurotransmitter and hormones.	Solicit students' questions and comments about nerves and neurotransmitter.
	D. How Drugs Work	
30 Minutes	1. In very simple terms, drugs usually work by artificially creating natural body reactions generally associated with the work of neurotransmitters and hormones.	
	a. Therapeutic doses of legitimate prescriptive and over the counter drugs are designed to produce mild and carefully controlled simulations of the natural action of neurotransmitters and hormones.	Ask students: What drug do many people take to overcome artificially the drowsiness they feel in the morning?

- b. Large, abusive doses of drugs may produce greatly exaggerated simulations of the natural action of hormones and neurotransmitters, sometimes with disastrous results.
- 2. When a person takes a drug and artificially simulates the natural action of hormones and neurotransmitters, the body's dynamic balance is disrupted.
 - a. The body automatically responds to the presence of the drug by producing other hormones and chemicals that can oppose the drug's effects, and bring the body back into balance.
 - (1) Example #1: If a person takes a stimulant drug that mimics neurotransmitters associated with the sympathetic nerves, the body may react by excreting hormones that depress the bodily functions that the drug is exciting.
 - (2) Example #2: If a person takes a drug that depresses some bodily function, the body may pour out one of its natural chemicals that stimulate that same function.
 - b. An interesting situation can occur when the drug is no longer psychoactive.

Example: Cocaine (a sympathomimetic drug) may artificially create a message commanding the heart to beat so rapidly that cardiac arrest results.

Remind students that the body struggles to maintain homeostasis, the dynamic balance of salts, sugars and other substances.

If a person ingested Cocaine, for example, the Cocaine would artificially stimulate the body functions. The body would then produce hormones and neurotransmitters to slow down the body functions to try to maintain homeostasis.

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Instructor Notes

- (1) The chemicals produced by the body in an effort to counteract the drug may still be active.
- (2) These natural chemicals have exactly the opposite effect on the body that the drug had: after all, that is precisely why the body produced those chemicals.
- (3) As a result, the person may feel, appear and act in a manner exactly opposite to the way he or she would feel, appear and act when under the influence of the drug.

Example: Ask students if they have ever experienced this situation...After drinking several drinks, they become drowsy, go to bed and fall asleep quickly. But, after a few hours, when it is still the middle of the night, they suddenly awaken and are wide awake, unable to fall asleep again. What has happened is that the alcohol has worn off, but the natural CNS Stimulants the body produced to counteract the alcohol are still around.

Write "Downside of the Drug" on the chalkboard or flip chart.

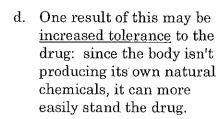


c. We call this situation being on the "downside" of the drug.

Aides	Lesson Plan	Instructor Notes
	(1) It is not uncommon for a DRE to encounter someone on the "downside".	·
- F H 2- 1		Example: with cocaine (a drug that is metabolized, or broken down by the body fairly quickly) the user may be exhibiting drowsiness and general depression by the time the DRE is called to the scene.
	(2) The concept of "Downside" will be	DRAW this diagram on the chalkboard or flip-chart:
encon grand	especially important to us when we discuss the effects of drug combinations.	DREs <u>do not</u> classify a subject as being "under the influence" of the downside of a drug.
<u> </u>	·	Solicit students' questions about <u>Downside</u> .
	3. Another interesting effect that drugs can produce is called Negative Feedback.	Write "Negative Feedback" on the chalkboard or flip chart.
·.	a. By taking the drug, the person artificially simulates the action of certain hormones and/or neurotransmitters.	
	b. If the person continues to take the drug, the body may simply cease producing the natural chemicals that the drug simulates.	•
F SI E	c. In effect, the body comes to rely on the drug to supply itself with those chemicals.	Write "The Body Quits Producing The Natural Chemicals" on the chalkboard or flip chart.

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Instructor Notes



Write "Increased Tolerance" on the chalkboard or flip chart.

Emphasize: Habitual users of drugs may develop tolerance to the drug. As a result, they may exhibit relatively little evidence of impairment on the psychophysical tests. Even tolerant drug users, when impaired, usually exhibit clinical evidence. (i.e. in the vital signs and eye signs - such as HGN)

e. Example of Negative
Feedback: When people
regularly use heroin,
cocaine or marijuana, their
bodies may cease producing
the neurotransmitters and
hormones known to be crucial for proper pain relief,
stress reduction, mental
stability and motivation.

Point out that because of this Negative Feedback, the user becomes dependent on the drug to cope with the stresses and strains of daily life.





- f. Another result may be <u>physical dependence</u>, or addiction.
- 4. Why do people take drugs?
 - a. In simplest terms, people take drugs because they like the feelings the drugs produce.

Write "Physical Dependence" on the chalkboard or flip chart.

Pose the questions to the class. Solicit responses.

Instructor Notes

- b. The artificial simulation of the natural action of hormones and neurotransmitters appears to permit the user to create any feeling or mood he or she desires.
- c. As time goes on, and negative feedback develops, the user finds that he or she can <u>only</u> achieve those feelings and moods if the drug is taken.
- 5. One final concept is important for an understanding of how drugs work.
 - a. A Metabolite is a product of metabolism, the chemical changes that take place when the drug reacts with enzymes and other substances in the body.

Write "Metabolite" on the chalkboard or flip chart.

<u>Instructor</u>, for your <u>information</u>:

Metabolism is defined as the combined chemical and physical processes that take place in the body involving the distribution of nutrients and resulting in growth, energy production, the elimination of wastes, and other body functions. There are two basic phases of metabolism: anabolism, the constructive phase, during which small molecules resulting from the digestive process are built up into complex compounds that form the tissues and organs of the body; and catabolism, the destructive phase, during which larger molecules are broken down into simpler substances with the release of energy.



Aides	Lesson Plan		Instructor Notes	
		b. The body uses chemical reactions to break down the drug, and ultimately to eliminate it.	Example: When we drink alcohol, we initiate a series of chemical reactions that ultimately transform the alcohol into harmless carbon	
		c. Sometimes, metabolites of the original drug are themselves drugs, and cause impairment.	dioxide and water.	
		d. For example, the body quickly metabolizes heroin into morphine, and it is the morphine that actually produces the effects the heroin user experiences.	Solicit students' questions and comments about how drugs work.	
	E.	Medical Conditions		
15 Minutes				
I II I	1.	Certain medical conditions or injuries may cause signs and symptoms similar to those of drug impairment.	List these conditions on the chalkboard or flip chart.	
VI-11 (Medical Conditions)				
	The state of the s	a. Head Trauma	Point out that head trauma may produce disorientation, confusion, unequal pupil size, unequal tracking ability of the eyes, or the drooping of one eyelid while the other remains normal.	
		b. Stroke	Point out that stroke may produce many of the same indicators as will head trauma.	

Aides		Lesson Plan	Instructor Notes
			In addition, stroke victims may have pupils that are markedly different in size, and one pupil may exhibit no visible reaction to light while the other reacts normally.
	c.	Diabetes Insulin shock (taking too much insulin) can produce tremors, increased blood pressure, rapid respiration, lack of coordination, headache, confusion and seizures.	The most common problem with diabetics arises when they take too much insulin, so that their blood sugar levels become extremely low. They may be very confused, sweat profusely, and exhibit increased pulse rate and increased blood pressure.
	d.	Conjunctivitis	Conjunctivitis is a condition caused by infection, allergy or irritation of the mucous membrane lining of the eyes, resulting in a "pink eye" appearance. A casual observer might mistake this for the bloodshot conditions associated with Cannabis or alcohol.
	е.	Shock	A shock victim may be dazed, uncoordinated, non-responsive.
	f.	Multiple Sclerosis, and similar conditions	Multiple Sclerosis is a progressive disease in which the nerve fibers of the brain and spinal cord lose their myelin cover. Some signs and symptoms are abnormal sensations in the face or extremities, weakness, double vision, ataxia, etc.
	g.	Carbon Monoxide poisoning	

h. Seizures

Endocrine disorders

A	i	d	e	S

Instructor Notes

- j. Neurological conditions
- k. Psychiatric conditions
- l. Infections
- 2. Normal conditions can affect vital signs.
 - a. Exercise
 - b. Excitement
 - c. Fear
 - d. Anxiety
 - e. Depression
 - f. Other
- 3. Point out that often times signs and symptoms can be contradictory.
 - a. Drug combinations may have an additive effect.
 - b. Drug combinations may cause unexpected effects.
 - c. Drug combinations may be used to mask symptoms.

Review physiologic changes that may be mistaken for drug induced symptoms. For example, strenuous exercise increases heart rate and rapidity and rate of respiration; surprise, fear and pain dilate the pupils markedly.

Total effect is greater than the sum of the effects taken independently.

For example, a CNS stimulant/ CNS depressant combination may cause the suspect to look and act like a "wide awake drunk".

For example, a person who has been using Marijuana, Cocaine, or some other drug may also consume a moderate amount of alcohol in the hope that, if they are stopped and asked to submit to a breath test, the arresting officer will be fooled by the low to moderate BAC into thinking that the suspect is simply "slightly" impaired by alcohol alone.

Instructor Notes



10 Minutes

d. Misinterpretation of symptoms of disease or injury in combination with consumption of alcohol.

- F. Summary
 - 1. Briefly review main points of the lesson.
 - a. Basic understanding of how the body works is necessary to:
 - o understand why the drug evaluation is conducted in a systematic manner.
 - o understand why the results, when viewed in their totality, provide reliable indicators of impairment within broad categories of drugs.

b. The body maintains
homeostasis (equilibrium)
by constantly adjusting to
changes in the external and
internal environment:

Suspect alcohol, however, impairment is not consistent with BAC.

Emphasize that research in drug intoxication and the interaction with neurotransmitters and neurohormones is in its infancy. There are many unknowns!

This limited overview will not qualify students as medical specialists!

The knowledge gained during this session must be supplemented by additional reading and/or instruction. The body of knowledge is being constantly expanded.

Point out that the best response to questions regarding bodily functions and or specific drug interactions is "I don't know. I conducted a series of evaluations and documented my observations. Based on my training and experience the results of my observations are consistent with those produced by persons impaired by ____."

Point out that body functions as a total unit in an integrated and coordinated manner.

Aides

	(1) When drugs are introduced into the body this process comes into play.	•
	(2) When drugs interact in the body they tend to:	
	o speed things up, or	Point out that this is a very simplistic overview of how
	o slow things down, or	drugs work.
	o confuse signals, or	,
	o block signals, or	
	o some combination of the above.	
	(3) The effects of drugs can be detected and/or observed in the drug evaluation.	
	2. Drug evaluations	
	a. Detailed instructions on procedures and expected results will be covered in following sessions.	Solicit and answer students' questions.
VI-12 (Physiological Pursuit)	3. Physiological Pursuit	For review of the Physiology and drugs session, questions can be asked of the students as if it were a game of Trivial Pursuit. See attachment.
	·	

Lesson Plan

Instructor Notes

Session VI

Physiology and Drugs: An Overview







Physiology and Drugs: An Overview

Upon successfully completing this session, the participant will be able to:

- Explain in layman's terms the general concept of human physiology
- Explain in layman's terms the purpose and functions of major systems in the body (nervous system, circulatory system, respiratory system, etc.)

Drug Evaluation & Classification Training

VI.OA

Physiology and Drugs: An Overview (continued)

- Explain in layman's terms how drugs work in the body
- Explain in general terms how the drug evaluation is used to detect signs or symptoms indicative of drug impairment
- Correctly answer the "topics for study" questions ate the end of this section

Drug Evaluation & Classification Training

VI-OB

Bodily Functions Examined During Drug Evaluation

- · Central nervous system
- · The eyes
- · Blood pressure and pulse
- · Balance and coordination
- · Body temperature

Drug Evaluation & Classification Training

VL1

Physiology:

The study of the functions of living organisms and their parts

Drug Evaluation & Classification Training

VI-2

Murders, Inc.

Drug Evaluation & Classification Training

VI-3A

The Ten Systems of Human Physiology: Murders, Inc.

M is for Muscular System

U is for Urinary System

I is for Integumentary System

R is for Respiratory System

N is for Nervous System*

D is for Digestive System

C is for Circulatory System*

E is for Endocrine System

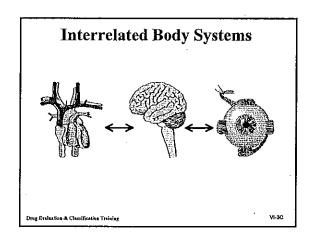
R is for Reproductive System

S is for Skeletal System

*For DRE officers, these are key systems

g Evaluation & Classification Training

VI-3B

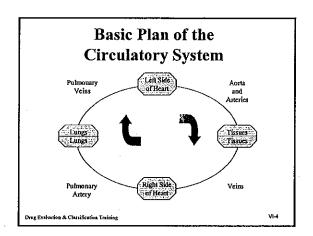


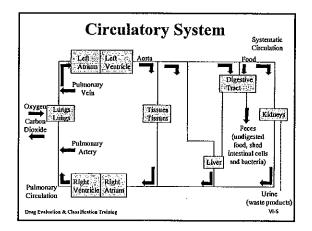
Homeostasis:

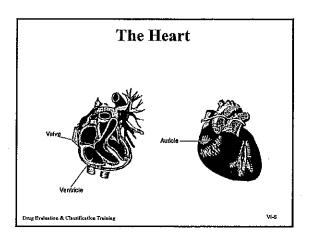
Dynamic balance or steady state involving levels of salts, water, sugars and other material in the body's fluids

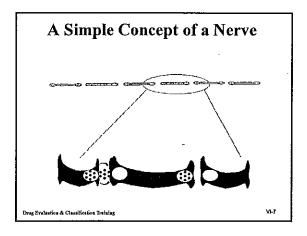
Drug Evaluation & Classification Training

VI-3D









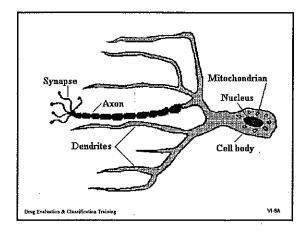
How a Neurotransmitter Works

Steps are numbered sequentially:

- 1. Neuron makes a neurotransmitter
- 2. Vesicles store neurotransmitter
- 3. Neurotransmitter enters gap to transmit electrical impulse to receptor site
- 4. Receptor performs a function

Drug Evaluation & Classification Training

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Classification of Nerves Motor - efferent nerves Sensory - afferent nerves Drug Evaluation & Classification Training

Motor Nerves

- Voluntary
- Autonomic

Drug Evaluation & Classification Training

VI-10

Tolerance

- May exhibit relatively little evidence of impairment on the psychophysical tests.
- Even tolerant drug users, when impaired, usually exhibit clinical evidence (i.e. vital signs, eye signs, etc.).

Drug Evaluation & Classification Training

VI-10A

Medical Conditions

- Head Trauma
- Stroke
- Diabetes
- Conjunctivitis
- Shock
- · Multiple Sclerosis, and similar conditions

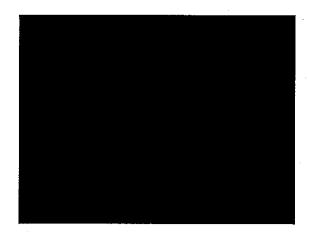
Drug Frahation & Classification Trainin

VI-11

Physiological Pursuit

Drug Evaluation & Classification Training

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INSTRUCTIONS FOR PHYSIOLOGICAL PURSUIT

- 1. Preparation and Rules of the Game
 - a. Ahead of time, secure five like items as prizes (such as lottery scratch off tickets).
 - b. Select two teams of five students each. Appoint a captain for each team. (Usually home team and visitors team. Attempt to balance teams and avoid "sharks".)
 - c. Appoint a time keeper.
 - d. Appoint a score keeper.
 - e. Select a panel of instructor judges.
 - f. On a flipchart or chalkboard, mark as follows:

	Score	e
Questions	<u>Home</u>	$\underline{ ext{Visitor}}$
1.		
2.		
3.		
4.		
5.		
6.		
7 .		•
8.		
9.		
10.		
11.		
12.		
13.		
14.		
15.		

- g. Place the teams on opposite sides of the room in view of the screen.
- h. Tape the overhead with the questions to the overhead projector.

- i. Cover all the questions with two pieces of paper. When a question is selected, reveal the question using the two papers to cover all others and turn the projector on long enough to read the question and repeat it. Then turn the projector off. The team getting the question has 20 seconds to discuss and come up with the "correct" answer. The captain can answer the question or designate a team member to do so.
- j. The judges decide if the answer is correct. If not, the other team may answer. If neither team gets the answer, no points are scored and the game goes on to the next question.

2. Playing the Game

- a. To start the game, flip a coin and have the team captains call the result while the coin is in the air. The winning team captain can elect to receive or pass the first question selection to the opposing team.
- b. The selected team stars with the question selection and the selection alternates until the game ends.
- c. As the questions are selected, the score keeper crosses out those selected. He also awards one point to the team answering the question correctly.
- d. "No coaching from the audience."
- e. The team with the most points after 14 questions wins. If the score is tied, use the last question to the break tie.

QUESTIONS FOR PHYSIOLOGICAL PURSUIT

- Name the major body systems.
 Muscular, Urinary, Respiratory, Digestive, Endocrine, Reproductive, Skeletal, Integumentary, Nervous, and Circulatory.
- 2. What vein carries oxygenated blood?

 Pulmonary vein. The pulmonary vein returns oxygenated blood from the lungs to the left side of the heart. The left side of the heart then pumps the oxygenated blood via arteries throughout the body. The pulmonary artery carries de-oxygenated blood from the right side of the heart to the lungs.
- 3. What is the function of the endocrine system?

 The endocrine system is composed of ductless glands that release chemical messengers, called hormones, into the bloodstream. The function is the regulation of various bodily processes by the production and release of hormones.
- 4. Explain the "downside" effect of a drug.

 The "downside" effect of a drug refers to the post euphoric stage of a drug's effects. As the effects of a drug wear off, the individual may display effects that are essentially the opposite of the "high" state that was brought about by the drug. This effect is in part due to the body's attempt to counteract the effects of a drug by releasing hormones into the bloodstream. DRE's should never write a report stating the suspect was under the influence of the "downside" of a drug.
- 5. Define homeostasis.

 Homeostasis is basically a physiological equilibrium or dynamic balance.

 Homeostasis refers to the body's mechanisms that keep the levels of fluids, salts, chemicals and other internal substances in a safe balance. The regulation of temperature is an example of homeostasis at work.
- 6. Hair and nails are part of what system?

 The Integumentary system. This system also includes the skin.
- 7. Name the two circulatory systems.

 The systemic circulatory system, which is driven by the left side of the heart, and pulmonary circulatory system, driven by the heart's right side.
- 8. The functions of the organs of the body are controlled by what two systems? The endocrine and nervous system.

- 9. Define synapse, axon, and dendrite.
 - These structures are all part of the nerve cell, or neuron. The axon is the part of the neuron that releases neurotransmitter from a terminal into the synapse. An electrical impulse causes the axon to release the neurotransmitter. The synapse is the gap between nerve cells and is also called the synaptic gap. The dendrite refers to a structure that receives the chemical message from the neurotransmitter. There are often many dendrites on each neuron. The neurotransmitter fit into receptor sites on the dendrite and cause an electrical message to be sent to the neuron's body.
- 10. Define neurotransmitter and hormone.

Both are chemical messengers. Neurotransmitter are chemicals that send messages within the nervous system. Hormones are released by glands in the endocrine system into the bloodstream.

- 11. _____ nerves carry messages AWAY from the brain to the body's muscles and organs.
 - Efferent, or Motor nerves. These nerves cause a motor response. Afferent nerves send sensory messages to the brain. The central nervous system interprets these messages and if appropriate, calls for a response through the efferent nerves.
- 12. The _____ nervous system commands the body to react to stress, fear, and excitement.

The Sympathetic nervous system, a division of the Autonomic Nervous System, produces the body's "fight or flight" response to real or perceived danger. Drugs that mimic the activation of the sympathetic nervous system are "sympathomimetics". CNS Stimulants have effects closest to the effects of sympathetic nervous system activation.

- 13. Explain "negative feedback."
 - Refers to the body's response to taking a drug that has effects similar to natural internal chemicals. After repeated exposure to the drug, the body responds by slowing, or even stopping the production of the internal chemical. In time, the body begins to rely on the drug. An example of negative feedback involving legitimate substances is insulin dependant diabetics. Once an individual begins to take insulin, the person's body will eventually stop making its own insulin. The person must obtain insulin by administering it.

- 14. What two types of nerves make up the autonomic nervous subsystem? The Sympathetic and Parasympathetic nerves. The sympathetic nervous system initiates the body's "fight or flight" response to real or perceived danger. The parasympathetic nervous system parallels or balances the sympathetic nervous system. This system initiates calming and digestive processes.
- 15. Define metabolite.

A metabolite is the by-product of the body's chemical breakdown of various substances for elimination. Metabolites may or may not be psychoactive by themselves. Often times a toxicological analysis will disclose various metabolites of a drug, rather than the parent drug.

Two Hours

SESSION VII EXAMINATION OF VITAL SIGNS

SESSION VII EXAMINATION OF VITAL SIGNS

Upon successfully completing this session, the participant will be able to:

- o Explain the purposes of the various vital signs examinations in the drug evaluation and classification process.
- o Explain the administrative procedures for these examinations.
- o Explain the cues obtained from these examinations.
- o Document the examinations of vital signs accurately and completely.
- o Correctly answer the "topics for study" at the end of this Section.

Content Segments		Lear	<u>Learning Activities</u>		
A.	Purpose of the Examinations	0	Instructor Led Presentations		
В.	Procedures and Cues	0	Instructor Led Demonstrations		
C.	Demonstrations	0	Audio Tape Presentation		
D.	Documentation Procedures	0	Student Led Demonstrations		
E.	Practice	0	Students' Hands On Practice		
		0	Reading Assignments		

		-		
А	1	М	Ω	C

Instructor Notes





VII-0 (Session Objectives)



5 Minutes



EXAMINATIONS OF VITAL SIGNS

Total Lesson Time: Approximately 120 Minutes

Session title on wall chart.

Briefly review the content, objectives and activities of this session.

- A. Purposes of the Examinations
 - 1. The Vital Signs that are relevant to the Drug Evaluation and Classification Process include:
 - a. Pulse rate
 - b. Blood pressure
 - c. Temperature
 - 2. Different types of drugs affect these vital signs in different ways.
 - a. Certain drugs tend to"speed up" the body andelevate these vital signs.
 - b. Other drugs tend to "slow down" the body and <u>lower</u> these vital signs.
 - 3. Systematic examination of the vital signs gives us much useful information concerning the possible presence or absence of various categories of drugs.

<u>Point out</u> these vital signs on the wall chart.

Clarification

- o pulse may quicken
- o blood pressure may rise
- o temperature may rise

Clarification

- o pulse may slow
- o blood pressure may drop
- o temperature may fall

Instructor Notes



75 Minutes



Definitions")

B. Procedures and Cues

- 1. Measurement of pulse rate.
 - a. <u>Pulse</u> is the expansion and relaxation of an artery generated by the pumping action of the heart.
 - b. <u>Pulse Rate</u> is the number of pulsations in an artery per minute.
 - c. An <u>artery</u> is a strong, elastic blood vessel that carries blood <u>from the heart</u> to the body tissues.

<u>Point out</u> that pulse rate is equal to the number of contractions of the heart per minute.

Instructor, for your information: Technically speaking, pulse rate is not quite the same thing as heart beat rate. There are rare and very serious conditions that could cause the heart to beat so weakly that it is unable to force blood through some or all arteries. In that case, there might be no discernable pulse even though the heart is beating. But with a normal, healthy heart, pulse rate will equal heart beat rate.

- d. A <u>vein</u> is a blood vessel that carries blood <u>back to the</u> <u>heart</u> from the body tissues.
- e. When the heart contracts, it squeezes blood out of its chambers into the arteries.
- f. The surging blood causes the arteries to expand.

Instructor Notes

- g. By placing your fingers on the skin next to an artery and pressing down, you can feel the artery expand as the blood surges through.
- h. By keeping your fingers on the artery and counting the number of pulses that occur in one minute, you will measure the pulse rate.
- Pulse is easy to measure, once you locate an artery close to the surface of the skin.
- One convenient pulse point involves the radial artery.
 - o The radial artery can be located in or near the natural crease of the wrist, on the side of the wrist next to the thumb.
 - o Hold your left hand out, with the palm down.
 - o Place the tips of your right hand's index finger and middle finger into the crease of your left wrist, and exert a slight pressure.
 - o Allow your left hand to curl downward.
 - o You should be able to feel the pulse in your radial artery.

Emphasize: The "surge" can be felt as the blood is squeezed from the heart through an artery. The pulse cannot be felt in a vein.

<u>Demonstrate this</u>, by holding your fingers on your own radial artery.

<u>Point to</u> the radial artery pulse point on your own wrist.

Demonstrate this.

Demonstrate this.

Demonstrate this.

<u>Ask</u> students whether they can feel their pulses. <u>Coach</u> any students who have difficulty in locating the pulse.



Aides

Lesson Plan

Instructor Notes



VII-3 ("Brachial Artery")

- k. Another pulse point involves the brachial artery.
 - o The brachial artery can be located in the crook of the arm, halfway between the center of the arm and the side of the arm closest to the body.
 - o Hold your left hand out, with the palm up.
 - o Place the tips of your right hand's index and middle fingers into the crook of your left arm, close to the body, and exert a slight pressure.
 - o You should be able to feel the pulse in your brachial artery.
- 1. Another pulse point involves the carotid artery.
 - o The carotid artery can be located in the neck, on either side of the Adam's apple.
 - o Place the tips of your right hand's index and middle fingers alongside the right side of your Adam's apple.
 - o You should be able to feel the pulse in your carotid artery.

<u>Point to</u> the brachial artery pulse point in your own arm.

<u>Instruct</u> students to roll up their sleeves, if necessary, to expose their brachial artery pulse points.

<u>Demonstrate</u> this.

Demonstrate this.

Ask students whether they can feel their pulses. Coach any students who have difficulty locating the pulse.

<u>Point out</u> the carotid artery pulse point on your own neck.

Demonstrate this.

<u>Ask</u> students whether they can feel their pulses. <u>Coach</u> any students who have difficulty locating the pulse.

Instructor Notes

- m. Basic do's and don'ts of measuring pulse.
 - o <u>Don't</u> use your thumb to apply pressure while measuring a subject's pulse.
 - o If you use the carotid artery pulse point, <u>don't</u> apply pressure to both sides of the Adam's apple: this can cut off the supply of blood to the brain.
 - o When measuring the pulse rate, use time intervals of 30 seconds.
- n. Some technical terms associated with pulse rate:
 - (1) <u>Tachycardia</u>: Abnormally rapid heart rate.
 - (2) <u>Bradycardia</u>: Unusually slow heart rate.
 - (3) <u>Arrhythmia</u>: Abnormal heart rhythm.
- o. Students' initial practice at measuring pulse rate.

<u>Note</u>, however, that there is wide variation in "normal" human pulse rate.

Point out that there is an artery located in the thumb close to the surface of the skin. If you apply pressure with the thumb, you may wind up measuring your own pulse when you think you are measuring the suspect's.

<u>Point out</u> that pulse rate is always expressed as "beats per minute". When you count the beats during an interval of 30 seconds, you must double the result to obtain the pulse rate.

<u>Instruct</u> students to work in pairs, taking turns measuring each other's pulse.



Aides	Lesson Plan	Instructor Notes
		Tell students to record on paper their partner's pulse rate.
		Monitor, coach and critique the students' practice.
		Allow the practice to continue for only about 5 minutes.
4 0 4		PRINT the following lists on the chalkboard or flip-chart:
AND		50 or less 76-78 52-54 80-82 56-58 84-86 60-62 88-90 64-66 92-94 68-70 96-98 72-74 100 or more
		TABULATE the numbers of students whose pulse rates were in each of the listed intervals.
		POINT OUT that the "normal range" of pulse rate is 60-90 beats per minute.
	2. Measurement of blood pressure.	
VII-4 ("Blood Pressure Definitions")	a. <u>Blood Pressure</u> is the force that the circulating blood exerts on the walls of the arteries.	
	o Blood pressure is measured in millimeters of mercury.	

Lesson Plan

Instructor Notes





- o Example: a blood pressure of 120 means that the blood is pressing on the walls of the artery with enough force to push liquid mercury 120 millimeters up a glass tube.
- o We commonly abbreviate "millimeters of mercury" as mmHg.

- b. Blood Pressure changes constantly as the heart contracts and relaxes.
- c. Blood Pressure reaches its maximum as the heart contracts and sends the blood surging through the arteries. This is called the systolic pressure.
- d. Blood Pressure reaches its minimum when the heart is fully expanded. This is called the diastolic pressure.
- e. It is always necessary to measure and record both the systolic and diastolic blood pressure.
- f. The device used for measuring blood pressure is called a sphygmomanometer.

<u>Point out</u> that 120 millimeters is approximately four and three-quarter inches.

Print "mmHg" on the chalkboard or flip-chart.

Instructor, for your information: "Hg" is the chemical symbol for the element mercury. It comes from Hydrargyrum, the Latin word for mercury.

Remind students that "systolic" is the higher number, "diastolic" the lower number.

Memory aid:

Systolic: "S" for "Superior" Diastolic: "D" for "Down"

Exhibit a sphygmomanometer.

Aides		Lesson Plan	Instructor Notes
			Write "SPHYGMOMANOMETER" on the chalkboard or flip chart.
	g.	The sphygmomanometer has a special cuff that can be wrapped around the subject's arm and inflated with air pressure.	Select a student to come before the class. Have the student sit in a chair facing the class, and roll up a sleeve (if necessary) to expose a bicep.
			Advise students to check for birth control implants in the upper left arm. If subject has an implant, blood pressure should be taken on the right arm and documented.
			Instruct the student to elevate the arm and squeeze the fist several times; explain that this helps to drain blood from the arm.
	h.	As the pressure in the cuff increases, the cuff squeezes tightly on the arm.	Wrap the cuff around the student volunteer's arm and inflate it.
	i.	When the pressure gets high enough, it will squeeze the artery completely shut.	Ask the student volunteer whether they can feel the pressure of the cuff.
	j.	Blood will cease flowing through the brachial artery. And, since the brachial artery "feeds" the radial artery, blood will also cease flowing through the radial artery.	Ask students: "What artery is located in the crease of the elbow?" (Point to that location on the student volunteer's arm).
	k.	If we <u>slowly</u> release the air in the cuff, the pressure on the arm and on the artery will start to drop.	Release the pressure in the cuff on the student volunteer's arm.

- Eventually, the pressure will drop enough so that blood will once again start to flow through the artery.
 - o Blood will start flowing in the artery once the pressure <u>inside</u> the artery equals the pressure outside the artery.
 - o The two pressures will become equal when the air pressure in the cuff drops down to the systolic pressure.
 - o When that happens, blood will spurt through the artery each time the heart contracts.

Ask students: "How far must the pressure in the cuff drop before the blood can start to squeeze through the artery.

Ask students: "What would happen if we allowed the pressure in the cuff to drop down to the <u>systolic</u> level, and held the air pressure at that level?"

<u>Point out</u> that the blood would spurt through the artery each time the heart <u>contracted</u>, but would cease flowing when the heart expanded.

Ask students: "How far down must the air pressure in the cuff drop before the blood will flow through the artery continuously?"

o Once the air pressure in the cuff drops down to the <u>diastolic</u> level, the blood will flow continuously through the artery.

Aides

Lesson Plan

Instructor Notes



VII-5 ("The Basics of Blood Pressure Measurement")

- m. Overview of procedures for measuring blood pressure.
 - o Apply enough air pressure to the cuff to cut off the flow of blood through the artery.
 - o Slowly release the air pressure until the blood just begins to spurt through the artery: that level will be the systolic pressure.
 - o Continue to release the air pressure until the blood flows continuously through the artery: that level will be the diastolic pressure.

- n. We can <u>listen</u> to the spurting blood, using a <u>stethoscope</u>.
 - o Apply the stethoscope to the skin directly above the artery.
 - o Apply pressure to the cuff, enough to cut off the flow of blood.

<u>Demonstrate</u>, using the student-volunteer (apply pressure to the cuff).

Slowly release the pressure in the cuff.

Ask students:

- (1) "How can we tell when the blood starts to spurt through the artery?"
- (2) "How can we tell when the blood is flowing continuously through the artery?"

Exhibit a stethoscope.

<u>Demonstrate</u>, using the student volunteer.

<u>Inflate</u> the cuff on the student volunteer's arm.

Lesson Plan

Instructor Notes

- o When no blood is flowing through the artery, we hear <u>nothing</u> through the stethoscope.
- o Slowly release the air from the cuff, letting the pressure start to drop.
- o When we drop to the systolic pressure, we start to hear a spurting sound.
- o As we continue to allow the air pressure to drop, the surges of blood become steadily longer.
- o When we drop to the diastolic pressure, the blood flows steadily and all sounds cease.
- o. The sounds that we listen to are called <u>Korotkoff Sounds</u>.

 They are divided into 5 phases.
 - o Phase 1 the first appearance of clear, tapping sounds that gradually increase in intensity.
 - o Phase 2 the sounds change to a murmur and take on a swishing quality.
 - o Phase 3 the sounds develop a loud, knocking quality (not quite as clear as the Phase 1 sounds).

Release the air in the cuff.

<u>NOTE</u>: This begins as a clear, tapping sound.

<u>NOTE</u>: The sounds take on a swishing quality, and become fainter.

Excuse the student volunteer and thank them for participating.

<u>Point out</u> that the beginning of Phase 1 corresponds to the systolic pressure.



Sounds")

Aides	Lesson Plan	Instructor Notes	
	o Phase 4 - the sounds become muffled and again have a faint swishing quality.		
	o Phase 5 - the sounds cease.	Point out that the beginning of Phase 5 corresponds to the diastolic pressure.	
VII-7 ("Sphygmoma nometer")	q. Familiarization with the sphygmomanometer.	Hand out stethoscopes and sphygmomanometers (one per each student is desirable. At a minimum, there should be one for every four students).	
VII-7	o The <u>compression cuff</u> contains an inflatable rubber bladder.	<u>Point out</u> the components of the sphygmomanometer on visual.	
		Point out that blood pressure cuffs come in three sizes, child, adult and extra large, depend- ing on the size of the bladder.	
	o A tube connects the bladder to the <u>manometer</u> , or pressure gauge.	Clarification: The manometer displays the air pressure inside the bladder. In the DRE Program, we use an aneroid (without fluid) pressure gauge.	
	o Another tube connects the bladder to the <u>pressure bulb</u> , which can be squeezed to inflate the bladder.		
	o The <u>pressure control</u> <u>valve</u> permits inflation of the bladder and regulates the rate at which the bladder is deflated.		

- To <u>inflate</u> the bladder, the pressure control valve must be twisted all the way to the right.
- When the valve is twisted all the way to the right, air can be pumped into the bladder, but no air can escape from the bladder.
- To <u>deflate</u> the bladder, twist the valve to the left.
- The more the valve is twisted to the left, the faster the bladder will deflate.
- r. Details of blood pressure measurement.
 - o If it proves difficult to hear the Korotkoff sounds, simply have the subject elevate the arm and squeeze the fist several times, to drain the arm: this will make the Korotkoff sounds louder.
 - o The manometer (pressure gauge) may be clipped on the subject's sleeve, so that it is readily viewable.
 - o Twist the pressure control valve all the way to the right.

Demonstrate this.



<u>Select</u> a student to serve as a blood pressure subject.
Demonstrate the procedures using the student.

Aides	Lesson Plan	Instructor Notes
	o Put the stethoscope earpieces in your ears.	Make sure the earpieces are turned forward, i.e., toward the nose.
	o Place the diaphragm or bell of the stethoscope over the brachial artery.	
	o Rapidly inflate the bladder to a pressure of at least 180.	Point out that, if the subject's blood pressure is very elevated, it may be necessary to inflate the bladder to a higher pressure.
	o Twist the pressure control valve slightly to the left to release the pressure slowly.	EMPHASIZE the need to release the pressure slowly. If the pressure drops too fast, the needle will sweep down the gauge too quickly to be read accurately.
		The pressure should be released at a speed that takes one full second for the needle to move a single gradation (i.e., 2 millimeters of mercury) on the gauge.
	o Keep your eyes on the gauge and listen for the Korotkoff sounds.	Point out that the needle on the pressure gauge generally will "bounce" slightly when blood starts to spurt through the artery.
		Excuse the student and thank him or her for participating. <u>Solicit</u> students' questions concerning these procedures.
	•	Point out that "normal" values of blood pressure are: Systolic 120 - 140 Diastolic 70 - 90

Note, however, that "normal" people can have significantly different blood pressures: there is wide variation in human blood pressure.

- s. Do's and Don'ts of blood pressure measurement.
 - o If you inflate the bladder and then need to repeat the measurement, wait at least three minutes to allow the subject's artery to return to normal.
 - o Hold the bell of the stethoscope with your fingers; don't slide it under the cuff: that will distort the measurement.
- t. Some technical terms associated with blood pressure:
 - (1) <u>Hypertension</u>: Abnormally high blood pressure.
 - (2) <u>Hypotension</u>: Abnormally low blood pressure.
- u. Students initial practice at measuring blood pressure.

If at least one sphygmomanometer and stethoscope are available for every two students, instruct students to practice in pairs. Otherwise, assign students to practice in teams of 3 or 4 members.

Monitor, coach and critique the students' practice.

Allow this practice to continue for only about 10 minutes.



Terms)



Aides	Lesson Plan	Instructor Notes	
	3. Measurement of temperature. a. Temperature is measured orally using an electronic thermometer.	Exhibit this.	
	b. Make sure that a fresh disposable mouthpiece is used each time.	Solicit students' comments and questions concerning this overview of procedures and cues.	
	C. Demonstrations		
15 Minutes	Pulse rate measurement demonstrations.	Select two students to come before the class.	
	a. Radial artery pulse point. Instruct the first student to measure the second student's pulse using the radial artery pulse point. (Simultaneously, the instructor should measure the subject's pulse using a carotid artery pulse point).		
	b. Carotid artery pulse point.	Instruct the second student to measure the first student's pulse using the carotid artery pulse point. (Simultaneously, the instructor should measure the subject's pulse using a radial artery pulse point.)	
		Excuse the two students and thank them for participating.	
	2. Blood pressure measurement demonstrations.	Select two other students to come before the class.	
		Instruct the first student to measure the second student's blood pressure.	
		Have the students reverse roles.	

Aides	Lesson Plan	Instructor Notes
		Excuse the two students and thank them for participating.
5 Minutes	D. Documentation Procedures	Review the sections of the Standardized Form used to record vital signs measurements.
20 Minutes	E. Practice	Instruct students to practice in teams of 2-4 members, taking turns measuring each other's vital signs.
		Monitor, coach and critique the students' practice.

Session VII

Examinations of Vital Signs



Examination of Vital Signs

Upon successfully completing this session, the participant will be able to:

- Explain the purposes of the various vital signs examinations in the drug evaluation and classification process
- Explain the administrative procedures for these examinations
- Explain the cues obtained from these examinations
- Document the examinations of vital signs accurately and completely
- Correctly answer the "topics for study" at the end of this section

Drug Evuluation & Classification Training

Vil-Q

Definitions Concerning "Pulse"

- Pulse
 - The expansion and relaxation of an artery due to the pumping action of the heart
- Pulse Rate
 - The number of pulsations in an artery per minute
- Artery
 - A strong, elastic blood vessel that carries blood from the heart to the body's tissues
- Vain
 - A blood vessel that carries blood back to the heart from the body's tissues

Drug Systemation & Classification Training

VII-1

Radial Artery Pulse Point



Dete Evaluation & Classification Training

VII-2

Brachial Artery Pulse Point



Drug Evaluation & Classification Training

VI)-3

Technical Terms Associated With Pulse Rate

- Tachycardia
 - Abnormally rapid heart rate
- · Bradycardia
 - Abnormally slow heart rate
- Arrhythmia

Abnormal heart rate rhythm

Drug Evaluation & Classification Training

VII-3A

Definitions Concerning Blood Pressure

· Blood Pressure

The force that the circulating blood exerts on the walls of the arteries

Systolic Pressure

The maximum blood pressure, reached as the heart contracts

Diastolic Pressure

The minimum pressure, reached when the heart is fully expanded

Drug Evaluation & Classification Training

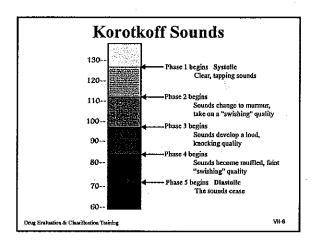
VII.4

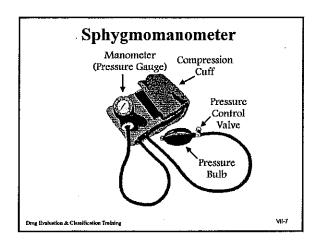
The Basics of Blood Pressure Measurement

- Apply enough air pressure to cut off the flow of blood through the artery
- Slowly release the air, 2 mmHg per second, until the blood just begins to spurt through the artery: that will be the systolic pressure
- Continue to release the air until the blood flows continuously: that will be the diastolic pressure

Drug Evaluation & Classification Training

VII-5





Details of Blood Pressure Measurement

- 1. Position cuff on bicep so that tubes extend down middle of arm
- 2. Wrap cuff snugly around bicep
- 3. Clip manometer to subject's sleeve
- 4. Twist pressure control valve all the way to the right
- 5. Put stethoscope earpieces in your ears
- 6. Place stethoscope over brachial artery
- 7. Rapidly inflate bladder to 180 mmHg
- 8. Twist the valve slightly to the left
- 9. Keep your eyes on the gauge and listen for the Korotkoff sounds

Drug Evaluation & Classification Training

VII-8

Technical Terms Associated With Blood Pressure

- Hypertension
 - Abnormally high blood pressure
- Hypotension

Abnormally low blood pressure

Drug Evaluation & Classification Training

VII-9

One Hour and Forty-Five Minutes

SESSION VIII

DEMONSTRATIONS OF THE EVALUATION SEQUENCE

SESSION VIII DEMONSTRATIONS OF THE EVALUATION SEQUENCE

Upon successfully completing this session, the participant will be able to describe the sequence in which examinations and other activities are performed in the Drug Evaluation and Classification process.

Content Segments Learning Activities A. Live Demonstrations o Instructor Led Presentations

- B. Video Demonstrations
- o Instructor Led Demonstrations
- o Video Presentations
- o Reading Assignments

Aides

Lesson Plan

Instructor Notes





VIII-0 (Session Objective)



70 Minutes

DEMONSTRATIONS OF THE EVALUATION SEQUENCE

Total Lesson Time: Approximately 105 Minutes

Session title on wall chart.

<u>Briefly</u> review the objective, content and activities of this session.

A. Live Demonstrations

For these Live Demonstrations, students must be grouped into teams of not more than 12 members. Each team must be taken to a separate classroom. At least two instructors must work with each team. This is to ensure that all students have the opportunity for a close and detailed observation of the demonstrations.

NOTE: Instructors should conduct at least two complete demonstrations of the evaluation sequence, articulating each step in the process.

<u>Instruct</u> students to follow along with copies of the report form.

Select a student or one of the volunteer drinkers for Session XII (prior to drinking) to serve as the "subject" for the preliminary examination.

Ask each question, exactly as it should be asked during an actual preliminary examination.

1. Preliminary Examination.

a. Preliminary eye checks

- o equal tracking
- o equal pupil size
- o blindness
- o eyelids
- o initial check for Nystagmus

Aides	Lesson Plan	Instructor Notes
		Explain the kinds of clues and evidence that may be gleaned during the preliminary examination.
	b. First measurement of pulse rate.	<u>Check</u> the student subject's eyes for tracking, equal pupil size, eyelids.
		Conduct a check of the student subject's pulse.
		Solicit students' comments or questions about the preliminary examination.
		Excuse the student subject and thank them participating in the demonstration.
	2. Eye Examinations (Room Light).a. Horizontal Gaze Nystagmus	Select another student <u>or a</u> <u>volunteer drinker</u> to serve as the "subject" for the eye examinations.
	b. Vertical Gaze Nystagmus c. Lack of Convergence	Conduct a complete demonstration of an eye examination.
		Explain the kinds of clues and other evidence that may be gleaned during the eye examinations.
		Solicit students' comments or questions about the eye examinations.
		Excuse the student subject and thank him or her for participating in the demonstration.

Aides	Lesson Plan	Instructor Notes
	 3. Psychophysical Tests. a. Romberg Balance b. Walk and Turn c. One Leg Stand d. Finger to Nose 	Select another student or a volunteer drinker to serve as the "subject" for the psychophysical tests.
		Conduct a complete set of psychophysical tests on the student subject.
		Explain the kinds of clues and other evidence that may be gleaned during the psychophysical tests.
·		Solicit students' comments or questions about the psychophysical tests.
		Excuse the student subject and thank them for participating in the demonstration.
	4. Vital Signs Examinations. a. Blood Pressure b. Temperature	Select another student to serve as the "subject" for the vital signs examination.
	c. Second Check of Pulse	Conduct a complete set of vital signs examinations on the student subject.
		Explain the kinds of clues and other evidence that may be gleaned during the vital signs examinations.
		Solicit students' comments or questions about the vital signs examination.
		Excuse the student subject, and thank them participating in the demonstration.

Aides	Lesson Plan	Instructor Notes
	5. Dark Room Examinations.	Point out that this portion of the Drug Evaluation and Classification Process is to be carried out in a darkened room.
		However, this demonstration will be conducted in normal room light, so that all students can observe the proper procedures for using the pen light.
		Select another student to serve as the "subject" for the dark room examination.
	a. Pupil Size Examinations o room light o darkness o direct light	Conduct a complete set of "dark room" examinations on the student subject.
	b. Reaction to Light c. Check of Nasal Area	Explain the kinds of clues and other evidence that may be gleaned during the dark room examinations.
	d. Check of Oral Cavity	Point out that the checks of the oral and nasal cavities actually are part of the examination for signs of ingestion.
		Solicit students' comments or questions about the dark room examinations.
		Excuse the student subject and thank them for participating in the demonstration.
	6. Examination for Muscle Tone and Injection Sites; Third Check of Pulse.	Select another student to serve as the "subject" for this portion of the examination.

Aides	Lesson Plan	Instructor Notes
		Point out that heroin is not the only drug that abusers inject: "Puncture marks" in the skin may also be found on the arms (and elsewhere) of abusers of several other drugs.
		Explain how to check for injection sites and muscle rigidity.
		Conduct a complete examination for injection sites and muscle rigidity on the student subject.
		Solicit students' comments or questions about this portion of the examination.
		Excuse the student subject, and thank them participating in the demonstration.
	7. Final Interview. a. Statements made by subject	Explain the kinds of clues and other evidence that may be gleaned during the final interview.
	b. Behavior during entire evaluation	Give examples of typical statements or behaviors of drug impaired subjects.
		Solicit students' comments or questions about the final interview.
	8. Opinions of Evaluator.	Point out that students subsequently will learn the clues and indicators of the various categories of drugs.
		Solicit students' comments and questions concerning the entire Drug Evaluation and Classification Process.

Aides	Lesson Plan	Instructor Notes
		NOTE: Be sure to conduct at least two complete, live demonstrations of the Drug Evaluation and Classification Process.
10 Minutes	B. Review of the 12-Step Process (Video)	

Session VIII

Demonstrations of the Evaluation Sequence



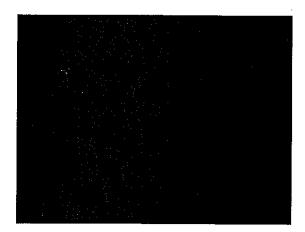
Demonstrations of the Evaluation Sequence

Upon successfully completing this session, the participant will be able to:

 Describe the sequence in which examinations and other activities are performed in the Drug Evaluation and Classification process

Drug Evaluation & Classification Training

VIII-O



One Hour and Forty-Five Minutes

SESSION IX

CENTRAL NERVOUS SYSTEM DEPRESSANTS

SESSION IX CENTRAL NERVOUS SYSTEM DEPRESSANTS

Upon successfully completing this session, the participant will be able to:

- o Explain a brief history of the CNS Depressant category of drugs.
- o Identify common drug names and terms associated with this category.
- o Identify common methods of administration for this category.
- o Explain the symptoms, observable signs and other effects associated with this category.
- o Explain the typical time parameters, i.e., onset and duration of effects, associated with this category.
- o State the clues that are likely to emerge when the drug evaluation and classification process is conducted for a person under the influence of this category of drugs.
- o Correctly answer the "topics for study" questions at the end of this Section.

Content Segments		<u>Lear</u>	Learning Activities	
A.	Overview of the Category	0	Instructor Led Presentations	
В.	Possible Effects	o	Review of Drug Evaluation and Classification Exemplars	
C.	Onset and Duration of Effects	O	Reading Assignments	
D.	Overdose Signs and Symptoms	O	Video Presentations	
E.	Expected Results of the Evaluation	0	Slide Presentations	

Aides	Lesson Plan	Instructor Notes
, 5,	CENTRAL NERVOUS SYSTEM DEPRESSANTS	Total Lesson Time: Approximately 105 Minutes
		Session title on wall chart.
IX-0A&B		Briefly review the objectives, content and activities of this session.
(Session Objectives)		
	A. Overview of the Category.	
20 Minutes	Central Nervous System Depressants slow down the operations of the brain.	Point out that other common names for CNS Depressants are "downers" and "sedative-hypnotics".
	a. Depressants first affect those areas of the brain that control a person's conscious, voluntary actions.	Judgment, inhibitions and reaction time are some of the things that CNS Depressants affect first.
	b. As the dose is increased, depressants begin to affect the parts of the brain that control the body's automatic processes.	
	o heartbeat o respiration o etc.	
	2. The CNS depressant category includes the single most commonly abused drug in America.	Ask this question: "What is the single most commonly abused drug?"
N 1 (NA)	a. Alcohol has been used and abused since prehistoric	

IX-1 ("Alcohol
- The Most
Familiar CNS
Depressant")

familiar to most Americans.

b. Alcohol and its effects are

times.

Aides

Lesson Plan

Instructor Notes



IX-2 ("Chloral Hydrate -- The First CNS Depressant Other Than Alcohol")

- c. Alcohol is a model for the CNS depressant category: with some exceptions, all depressants produce effects that are quite similar to the effects of alcohol.
- 3. Non-Alcohol CNS depressants have been around for more than 150 years.
 - a. The first non-alcohol CNS depressant was <u>chloral</u> <u>hydrate</u>.
 - b. It was developed in 1832.
 - c. It is commonly referred to as "Mickey Finn" or "Knockout drops" because of its fast acting effects.
 - d. Chloral hydrate is still produced and prescribed today.
- 4. There are six major subcategories of CNS depressants other than alcohol.
 - a. Barbiturates
 - o derivatives of barbiturate acid
 - o first produced in 1864
 - o in very common use and abuse today
 - b. Non-barbiturates
 - o synthetic compounds with a variety of chemical structures

<u>Point out</u> that the remainder of this session will focus on the non-alcohol CNS depressants.

Chloral Hydrate was derived from alcohol.

<u>Clarification</u>: "Mickey Finn" was a well known British prizefighter of the 19th Century.

"Felsule" and "Noctec" are two registered brand names of chloral hydrate.

More than 250 different barbiturates have been produced. Of these, about 50 have been accepted for medical use.

<u>Note</u>: Chloral Hydrate belongs to the non-barbiturate subcategory.



IX-3 (Major Types of Non-Alcohol CNS Depressants).

Aides		Lesson Plan	Instructor Notes
		o avoid some of the undesirable side effects of barbiturates	i.e. sleepiness or drowsiness
		o still produce physical and psychological dependence.	·
	c.	Anti-Anxiety Tranquilizers o first produced in 1950	The Anti-Anxiety Tranquilizers are also know as the "Minor Tranquilizers"; They include
		o in very wide spread use	the group of drugs known as the "Benzodiazepines", examples of which are Valium,
•		o frequently abused	Xanax and Librium.
	d.	Anti-depressants o sometimes called the "mood elevators" Anti-psychotic tranquilizers	Point out that it is not a contradiction to call one subcategory of CNS Depressants the Anti-depressants. It is psychological depression that they are "anti". Prozac is an anti-depressant but generally doesn't have psycho-active properties or side effects. Point out that the anti-
		Time physicolic stampanizors	psychotic tranquilizers are generally more powerful than the anti-anxiety tranquilizers.
		o sometimes called the "major tranquilizers"	The most familiar Anti- Psychotic Tranquilizer is "Thorazine".
		o Anti-Psychotic Tranquilizers were first introduced in the early 1950's. They provide a way to manage schizophrenia and other mental disorders, and allow psychiatric patients to be released from hospitals and to lead fairly normal lives.	I MOI dZinic .

Aides	Lesson Plan	Instructor Notes
	f. Combinations of the other five subcategories.	
•	5. Examples of specific common CNS Depressants.	Note: Briefly review these examples.
		Emphasize that students are not expected to memorize the names of these various CNS depressants. But, if they see these names, they should be able to recognize them as depressants.
	a. The Barbiturates	
IX-4A	o <u>Secobarbital</u> (Trade name "Seconal") (Street names "reds";	Mainly manufactured by Eli Lilly Pharmaceutical Co.
	"red devils"; "RDs"; "fender benders"; "F-40s")	The code "F40" is used by Eli Lilly to designate one capsule version of Seconal.
	o <u>Pentobarbital</u> (Trade name "Nembutal") (Street names "yellows"; "yellow jackets")	Mainly manufactured by Abbot Pharmaceutical Co.
	o <u>Amobarbital</u> (Trade name "Amytal") (Street names "blues"; "blue heavens")	Mainly manufactured by Eli Lilly.
	o <u>Amosecobarbital</u> (Trade name "Tuinal") (Street names "rainbows"; "Christmas trees")	Manufactured by Eli Lilly Note: this is a combination of Amobarbital and Secobarbital.
	o <u>Phenobarbital</u>	Manufactured by many

companies, one of which is

Smith, Kline and French.

(Many trade names)

(Street name "pink

ladies")

Aides	Lesson Plan	Instructor Notes
		Phenobarbital has been called the "Model T" of sedatives: it was first introduced in 1912. According to the "Physician's Guide to Psychoactive Drugs", 1 ounce of 80-proof alcohol is equivalent to about 15 milligrams of Phenobarbital.
F7-		<u>If available</u> : display 35mm slides of these various drugs.
	b. The Non-Barbiturates	
IX-4B	o <u>Chloral hydrate</u> (Trade names "Felsule"; "Noctec") (Street names "Knock out drops"; "Mickey Finn")	Point out that primary medical use for the Non-Barbiturates is the treatment of insomnia.
	o <u>Glutethimide</u> (Trade name "Doriden")	Note: the absence of street names implies only that illicitly manufactured versions
	o <u>Methyprylon</u> (Trade Name "Noludar")	of these drugs are not common. The <u>legally</u> manufactured versions are abused, however.
	o <u>Methaqualone</u> (Trade names "Parest"; "Quaalude"; "Sopor"' "Optimil"; "Mandrax") (Street name "ludes")	Note: methaqualone continues to be pharmaceutically manufactured in Mexico, trade name "Mandrax".
	o <u>Ethchlorvynol</u> (Trade name "Placidyl")	
	o <u>Diphenhydramine</u> <u>Hydrochloride</u> (Trade names "Benadryl"; "Sominex")	
	o <u>Ethinamate</u> (Trade name "Valmid")	

Aides	Lesson Plan	Instructor Notes
	o <u>Paraldehyde</u> (Trade name "Paral")	If available: display 35mm slides of these various drugs.
	o <u>Carisoprodol</u> (Trade name "Soma")	
	o <u>Diphenhylhydantoin</u> <u>Sodium</u> (Trade name "Dilantin")	
	o <u>Gamma-</u> <u>Hydroxybutyrate</u> (Street name "GHB", "Liquid X")	
	c. The Anti-Anxiety Tranquilizers	
IX-4C	o <u>Chlordiazepoxide</u> (Trade name "Librium")	Point out that <u>tens of millions</u> of prescriptions for these anti-anxiety tranquilizers are
	o <u>Diazepam</u> (Trade name "Valium")	written in America each year.
	o <u>Alprazolam</u> (Trade name "Xanax")	
	o <u>Lorazepam</u> (Trade name "Ativan")	
••	o <u>Triazolam</u> (Trade name "Halcion")	
,	o <u>Flurazepam</u> (Trade name "Dalmane")	<u>If available</u> : display 35mm slides of these various drugs.
	o <u>Estazolam</u> (Trade name "ProSom")	
	o <u>Temazepam</u> (Trade name "Restoril")	
	o <u>Oxazepam</u> (Trade name "Serax")	

Aides	Lesson Plan	Instructor Notes
	o <u>Clonazepam</u> (Trade name "Klonopin")	
	o <u>Flunitrazepam</u> (Trade name "Rohypnol") (Street Name "Roofies", "Roches")	
	d. The Anti-Depressants	
IX-4D	o <u>Phenelzine Sulfate</u> (Trade name "Nardil")	
	o <u>Amitriptyline</u> <u>Hydrochloride</u> (Trade names "Elavil"; "Endep")	
	o <u>Desipramine</u> <u>Hydrochloride</u> (Trade names "Nor- pramin"; "Pertofrane")	
	o <u>Doxepin Hydrochloride</u> (Trade names "Adapin"; "Sinequan")	
	o <u>Fluoxetine</u> (Trade name "Prozac")	Prozac generally does not have psychoactive properties in
IX-4E	o <u>Imipramine</u> (Trade name "Tofranil")	therapeutic doses.
	e. The Anti-Psychotic Tranquilizers	,
	o <u>Lithium Carbonate</u> (Trade name "Lithane")	
	o <u>Lithium Citrate</u>	
	o <u>Droperidol</u> (Trade names "Inapsine"; "Innovar")	

Aides	Lesson Plan	Instructor Notes
	o <u>Haloperidol</u> (Trade name "Haldol")	
	o <u>Chlorpromazine</u> (Trade name "Thorazine")	
	f. The Combinations	
IX-4F	o <u>Chlordiazepoxide</u> in combination with <u>Amitriptyline</u> (Trade name "Limbitrol")	Point out that "Limbitrol" is a combination of an Anti-Anxiety Tranquilizer and an Anti-Depressant.
	o <u>Perphenazine</u> in combi- nation with <u>Amitri-</u> <u>ptyline Hydrochloride</u> (Trade name "Triavil")	<u>Point out</u> that "Triavil" is a combination of an Anti- Psychotic Tranquilizer and an Anti-Depressant.
	o <u>Chlordiazepoxide</u> <u>Hydrochloride</u> in combination with <u>Clidinium Bromide</u> (Trade name "Librax")	
IV 5	6. Methods of ingestion of CNS Depressants.	
1X-5	a. Most common and easiest method is <u>orally</u> .	
	b. Some abusers prefer to use intravenous injection for Barbiturates.	
	c. Some abusers experience a "flash" or "rush" from intravenous injection of Barbiturates, that they do not experience from oral	

ingestion.

P	۱i	d	e	S

Lesson Plan

Instructor Notes

d. The injection paraphernalia used for Barbiturates are very similar to those used for Heroin.

- e. However, the Barbiturate abuser will use a larger hypodermic needle, because the barbiturate solution is thicker than the heroin solution.
- f. The injection sites on the skin of a Barbiturate abuser appear quite different from those of an Heroin addict.
- g. A large swelling, about the size of a quarter or fifty cent piece frequently will appear at the Barbiturate injection site.
- h. Necrosis may occur: i.e., a decaying of the body's tissue at the injection site.
- I. The dead tissue may begin to separate from the living tissue, producing ulcerations.
- j. The Barbiturate user who injects the drug usually will not display the same type of track marks as the heroin addict who uses repeated injections along the same vein.

Examples:

- o spoon, for heating and dissolving the barbiturate.
- o cotton, for filtering the solution when drawing it into the needle.
- o hypodermic syringe.
- o tourniquet.

Note: The "gauge" of a hypodermic needle indicates the width of the needle's inside diameter. The smaller the number, the larger the needle. For example, a 16 gauge needle is larger in diameter than a 20 gauge needle.

Point out that these effects result from the skin's reaction to the high alkaline content of the barbiturate solution.

<u>If available</u>, display a 35 mm slide showing ulcerated injection sites.

<u>Point out</u> that these ulcerations resemble burns placed on the skin by the tip of a cigarette.

Aides		Lesson Plan	Instructor Notes
	k B. P	Barbiturate abusers often will inject in parts of the body other than the forearm, and will commonly exhibit the characteristic swellings at random locations on the extremities.	Solicit students' questions and comments about the overview of CNS depressants.
5 Minutes IX-6	ir m	NS Depressants produce apairments of the human ind and body that essentially	<u>Point out</u> that these effects will not necessarily appear in a predictable sequence as dose
		irror alcohol impairment. reduce social inhibitions	increases.
	b.	impair the ability to divide attention	<u>Clarification</u> : impede the person's ability to concentrate on more than one thing at a time.
	c.	slow reflexes	U
	d.	impede judgment and concentration	
	e.	impair vision	Elaboration: ability to focus
	f.	impair coordination	eyes may be impaired; "double vision" may develop.
	g,	cause speech to become slurred and incoherent	Emphasize: The extent to which a CNS depressant user will exhibit these effects will depend, in part, on the user's tolerance to these drugs. Persons habituated to a drug often won't exhibit its effects as clearly as will a novice user.
	h.	produce a variety of emo- tional effects, such as eu- phoria, depression, suicidal tendencies, laughing or	

crying without provocation,

etc.

Aides	Lesson Plan	Instructor Notes
	2. Analogy: a person under the influence of CNS Depressants is like a 45 rpm record played at 33 1/3.	
	3. Generally speaking, a person under the influence of CNS Depressants will look and act drunk.	Solicit students' questions and comments concerning possible effects of CNS depressants.
	C. Onset and Duration of Effects	
IX-7 ("Onset and Duration Classes")	Depressant drugs can be grouped loosely into four classes, based on how quickly they take effect and how long their effects last.	Selectively reveal.
	a. <u>Ultrashort</u> : very fast acting, very brief effects.	
	o take effect in a matter of seconds.	
	o effects last only a few minutes.	•
	o very rarely are the "drugs of choice" for drug abusers.	Ask students: "Why is there little or no street abuse of the ultrashort CNS depressants"?
·		Solicit responses.
		Guide respondents to bring out the point that abusers seek drugs that will produce reason- ably long lasting effects. Effects that last for only a few minutes aren't attractive or satisfying to most drug abusers.
		·

Aides	Lesson Plan	Instructor Notes
	o ultrashort depressants are sometimes used at the beginning of a surgical operation, in conjunction with an inhaled anesthetic.	Clarification: to provide a momentary sedation to ease the patient's anxiety and allow for the proper administration of the anesthetic.
	o psychiatrists sometimes use ultrashort depressants at the beginning of a session, to reduce the client's inhibitions and foster a free and open communication.	
	o common example of an ultrashort depressant is thiopental sodium, brand name "Pentothal".	<u>Point out</u> that this is sometimes called "truth serum".
9900	b. <u>Short</u> : fairly fast acting, effects last for several hours.	·
	o generally take effect in 10-15 minutes.	
	o effects last for approximately 4 hours.	
	o this is the most commonly abused class of CNS Depressants.	<u>Point out</u> that short acting depressants are attractive to many drug abusers because:
		o they produce effects reasonably quickly.
		o the effects last long enough to "enjoy".
		o the effects don't last so long

that the user is in a prolonged state of impairment.

Aides	Lesson Plan	Instructor Notes
	o short acting Depressants frequently are prescribed as a treatment for insomnia.	
	o they also may be used as a pre-anesthetic medication to calm a patient prior to surgery.	
	o common example of a short acting Depressant: Secobarbital, brand name "Seconal".	
	c. <u>Intermediate</u> : relatively slow acting, but prolonged effects.	
•	o generally take effect in about 30 minutes.	
	o effects typically last about 6-8 hours.	
	o fairly often abused, especially by users who desire a longer lasting state of intoxication.	
	o medical use of this class of drugs is similar to that of short acting Depressants. (i.e., treat insomnia, etc.)	
	o common example of an intermediate Depressant: amobarbital, brand name "Amytal".	

Aides	Lesson Plan	Instructor Notes
	o a popularly abused drug is Amobarbital in combination with Secobarbital.	Point out that this amosecobarbital ("Tuinal") combination offers a fast acting drug (10-20 minutes onset, thanks to the Seconal) with prolonged effects (up to 8 hours, thanks to the Amytal).
	d. <u>Long</u> : delayed but long lasting effects.	
	o generally take effect about one hour after ingestion.	
	o effects typically last 8-14 hours.	
	o generally not the "drugs of choice" for abusers.	Ask students: "Why don't drug abusers usually prefer the long acting depressants?"
	o however, some people will abuse the long act- ing Depressants if the more popular short and intermediate types are not readily available.	Solicit students' questions and comments about the overview of CNS depressants.
	o long acting depressants are used medically in the control of epilepsy and of other conditions that can cause convulsions.	
	o they can also be used to provide continuing sedation to patients suffering from extreme anxiety.	
	o example of a long acting Depressant: Barbital, brand name "Veronal".	

Aides	Lesson Plan	Instructor Notes
	2. Alcohol as a specific example.	Ask students: "How would you classify alcohol in terms of the onset and duration of its effects?"
		Probe question: Suppose an average person drank two shots of whiskey. How long would it be before he or she started to feel the effects?
		(solicit responses).
		Probe question: How long would the average person continue to feel the effects of those two shots?
		(solicit responses).
		Guide students toward the conclusion that alcohol would be classified as a short or short to intermediate depressant.
IX-8 (examples of short to intermediate depressants)	3. Other examples of short to intermediate Depressants.	
	a. Barbiturates o Seconal ("reds") o Nembutal ("yellows") o Tuinal ("rainbows")	Point out that these are frequently abused CNS depressants, but they are not the only depressants that are abused.
	o Amytal ("blues")	
	b. Non-barbiturates	
	o Noctec or Felsule ("Mickey Finn") o Doriden	
,	o Noludar	•

Aides	Lesson Plan	Instructor Notes
	o Quaalude ("ludes") o Placidyl o Valmid o Equanil or Miltown o Soma	
	c. Anti-anxiety tranquilizers	
	o Valium o Librium o Xanax o Serax	
	D. Overdose Signs and Symptoms	
5 Minutes	1. Overdoses of Central Nervous System Depressants produce symptoms essentially identical to those of alcohol overdoses.	
·	a. Subject will become extremely drowsy and may pass out.	·
	b. The heartbeat (pulse) will slow.	
	c. Respiration will become shallow.	
	d. Skin may feel cold and clammy.	
	2. One major danger with CNS Depressant overdoses is death from respiratory failure.	
	a. A sufficiently high dose of CNS Depressant will suppress the portions of the brain that control respiration.	

Aides		Lesson Plan	Instructor Notes	
		b. This situation only rarely occurs from alcohol intoxication: usually, a drinker will pass out before he or she consumes enough alcohol to suppress respiration completely.		
	·	c. With other Depressants, it is relatively easy to take a fatal overdose.	Point out that CNS depressants are often used as a means of suicide.	
	3.	Another major danger with CNS Depressants occurs when they are combined with alcohol.	•	
		a. There is <u>at least</u> an additive effect when alcohol and another Depressant are taken together.		
		b. With many CNS Depressants, there may be a more than additive effect.	Clarification: the combination of alcohol and certain other CNS Depressants may produce an effect greater than the sum	
		c. Coroners have reported a number of cases in which neither the Alcohol level nor the Depressant level independently, would have been close to a fatal dose.	of the effects of the two drugs independently.	
		d. It is not possible to predict how great an effect will occur when Alcohol is mixed with another Depressant.		
•		e. However, it is clear that the combination is always risky.	Solicit students' questions and comments concerning overdoses of CNS depressants.	
60 Minutes	E.	Expected Results of the Evaluation	·	

Aides

Lesson Plan

Instructor Notes



IX-9A ("SFST evidence")

- 1. Observable evidence of impairment.
 - a. Standardized field sobriety tests.
 - o Horizontal Gaze
 Nystagmus will be
 present with suspects
 under the influence of
 CNS Depressants.
 - o Vertical Gaze

 Nystagmus <u>may</u> be

 present, with high

 doses, of Depressants

 for that individual.
 - o Performance on Walk and Turn and One Leg Stand will be similar to that of suspects impaired by alcohol.
 - o Performance on Romberg and Finger to Nose tests will be similar to that of suspects impaired by alcohol.
 - b. General indicators
 - o drowsiness
 - o droopy eyes (ptosis)
 - o thick, slurred speech
 - o uncoordinated
 - o fumbling
 - o slow reactions, sluggish
 - o muscle tone flaccid

2. Evidence associated with the Physiologic Examinations.

Point out that, if a person is under the influence of a combination of alcohol and some other CNS Depressant, the onset angle of HGN will not be consistent with the person's BAC: in other words, the eyes will start to jerk earlier than would be expected due to the alcohol alone.

Point out that suspect's perception of time (on Romberg) may be slowed, i.e., may estimate "30 seconds" after more than 30 seconds have elapsed.

Note: speech may also be incoherent.

Analogy: drunken behavior without the odor of alcoholic beverages.

But remind students: suspects may have consumed alcohol and some other CNS depressant. Hence, odor of alcoholic beverage may also be present.



("General

Indicators")

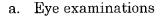
Aides

Lesson Plan

Instructor Notes



IX-9C ("Eye Examinations")

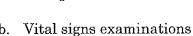


o Lack of Convergence generally will be present

o pupil size generally will be normal

o pupillary reaction to light will be slowed

Exception: Methaqualone or Soma usually will cause pupils to dilate.

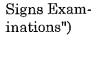


o blood pressure will be down

o pulse will be down

o body temperature generally will be normal Possible exceptions:

Methaqualone and alcohol may cause the pulse to be <u>increased</u>.



IX-9D ("Vital



IX-10 ("CNS Depressant Symptomatology Chart")

3. Summary

4. Demonstrations

a. Video tape demonstrations

 b. Drug Evaluation and Classification Exemplar Demonstrations Show video tape of subject(s) under the influence of CNS Depressants. Relate behaviors and observations to the CNS Depressant Symptomatology Chart.

Refer students to the exemplars found at the end of section IX of their student manuals.

Relate the items on the exemplars to the CNS Depressant Symptomatology Chart.

Solicit students' questions or suggestions concerning Expected Results of the Evaluation of subjects under the influence of Depressants.

Session IX

Central Nervous System
Depressants





Central Nervous System Depressants

Upon successfully completing this session, the participant will be able to:

- Explain a brief history of the CNS depressant category of druos
- Identify common drug names and terms associated with this category
- Identify common methods of administration for this category

Drug Evaluation & Classification Training

IX-OA

Central Nervous System Depressants (continued)

- Explain the symptoms, observable signs and other effects associated with this category
- Explain the typical time parameters, i.e., on-set and duration of effects, associated with this category
- State the clues that are likely to emerge when the drug evaluation and classification process is conducted for a person under the influence of this category of drugs
- Correctly answer the "topics for study" questions at the end of this Section

Drug Evaluation & Classification Training

IX-OB

Alcohol - The Most Familiar CNS Depressant





Drug Evaluation & Classification Training

IX-1

Chloral hydrate ("Mickey Finn"):

The first non-alcohol CNS depressant

Drug Evaluation & Classification Training

1X-2

Major Types of Non-alcohol CNS Depressants

- Barbiturates
- Non-barbiturates
- Anti-anxiety tranquilizers
- · Anti-depressants
- Anti-psychotic tranquilizers
- Combinations

Drug Evaluation & Classification Training

1X-3

Specific Barbiturates Examples			
Drug	Brand Name	Street Names	
Secobarbital	Seconal	Reds, Red Devils, RDs, Fender Benders, F-40's	
Pentobarbital	Nembutal	Yellows, Yellow Jackets	
Amobarbital	Amytal	Blues, Blue Heavens	
Amosecobarbital	Tuinal	Rainbows, Christmas Trees	
Phenobarbital	Luminal	Pink Ladies	
Drug Evaluation & Classification Trai	nîng	IX-4A	

Drug	Brand Name	Street Name
Chloral hydrate	Felsule, Noctec	Knock Out Drops, Mickey Finn
Glutethimide	Doriden	1
Methyprylon	Noludar	
Methaqualone	Parest, Quaalude, Sopor, Optimil, Mandra	Ludes
Ethelorvynol	Placidyl	
Diphenhydramine Hydrochlori	de Benadryl, Sominex	1
Ethinamate	Valmid	1
Paraldehyde	Paral	
Carisoprodol	Soma	
Diphenhylhydantoin Sodium	Dilantin	
Gamma Hydroxybutyrate	-	GHB, Liquid X

Specific Anti-Anxiety Tranquilizers Examples		
Drug	Brand Names	Street Names
Chlordiazepoxide	Librium	
Diazepam	Valium	
Alprazolam	Xanax	
Lorazepam	Ativan	
Triazolam	Halcion	
Flurazepam	Dalmane	
Estazolam	Prosom	
Temazepam	Restoril	
Oxazepam	Serax	
Clonazepam	Clonopin	7.4
Flunitrazepam	Rohypnol	Roofies, Roches
Drug Evaluation & Classification Training	IX-4G	

Drug	Trade Name
Phenelzine Sulfate	Nardil
Amitriptyline hydrochloride	Elavil, Endep
Desipramine hydrochloride	Norpramin, Pertofrane
Doxepin hydrochloride	Adapin, Sinequan
Fluoxetine	Prozac
mipramine	Tofranil

Specific Anti-Psychotic Tranquilizers Examples

Drug	Trade Name
Lithium Carbonate	Lithane
Lithium Citrate	
Droperidol	Inapsine, Innovar
Haloperidol	Haldol
Chlorpromazine	Thorazine

Drug Evaluation & Classification Training IX-4

Specific Combinations of Depressants

- Chlordiazepoxide in combination with Amitriptyline
 - Trade name: "Limbitrol"
- Perphenazine in combination with Amitriptyline Hydrochloride

Trade name: "Triavil"

 Chlordiazepoxide Hydrochloride in combination with Clidinium Bromide

Trade name: "Librax"

Drug Evaluation & Classification Training

IX-4F

Methods of Administering **CNS Depressants**







Injection

Drug Evaluation & Classification Training

Possible Effects of **CNS Depressants**

- Reduced inhibitions
- · Lack of coordination
- Inability to divide attention
- · Slurred and incoherent speech
- Slowed reflexes
- · Emotional instability
- Poor Judgment and impaired concentration
- Impaired vision

rag Evaluation & Classification Training

D-X3

Onset and Duration Classes

- Ultrashort
- Very fast acting, very brief effects
- Fairly fast acting, effects last several hours
- Intermediate
 - Relatively slow acting but prolonged effects
- Long
 - Delayed but long-lasting effects

Drug Eyelsation & Classification Training

Examples of Short-to-Intermediate CNS Depressants

- Barbiturates
 - Seconal
 - Nembutal
 - Tuinal
 - Amytal
- · Anti-anxiety tranquilizers
 - Valium
 - Librium
 - Xanax
 - Serax
- raluation & Classification Trainin

- Non-barbiturates
 - Noctec or Felsule
 - Doriden
 - Noludar
 - Quaalude

 - Placidyl
 - Valmid Equanil or Miltown
 - Soma

IX-8

Evaluation of Suspects Under the Influence of CNS Depressants

SFST Evidence

- Horizontal Gaze Nystagmus will be present
- · Vertical Gaze Nystagmus may be present (with high doses for that individual)
- · Impaired performance will be evident on Walk and Turn and One Leg Stand
- · Impaired performance will be evident on Romberg and Finger to Nose

Drug Evaluation & Classification Training

IX-9A

Evaluation of Suspects Under the Influence of CNS Depressants

General Indicators

- Drowsiness
- · Droopy eyelids (ptosis)
- · Thick, slurred speech
- Uncoordinated
- Fumbling
- · Slow, sluggish reactions

Drug Evaluation & Classification Training

Evaluation of Suspects Under the Influence of CNS Depressants

Eye Examinations

- · Lack of Convergence present
- · Pupil size will be normal*
- · Pupillary reaction to light will be slow
 - * Methaqualone and Soma will cause pupil dilation

Drug Evaluation & Classification Training

1X-9

Evaluation of Suspects Under the Influence of CNS Depressants

Vital Signs

- · Blood pressure will be down
- · Pulse will be down*
- Body temperature will be normal
 - * Quaaludes and ETOH may elevate

e Evaluation & Classification Training

(X-9D

CNS Depressant Symptomatology Chart

HGN	Present
Vertical Gaze Nystagmus	Present (High dose for that individual)
Lack of Convergence	Present
Pupil Size	Normai*
Reaction to Light	Slow
Pulse Rate	Down**
Blood Pressure	Down
Temperature	Normal
Muscle Tone	Flaccid

· Soma and Quaaludes usually dilate pupils

** Quaaludes and ETOH may elevate

Drug Evaluation & Classification Training

IX-10



000426	Drug Influence			
Evaluator Alley, Vern	DRE No 0323	Rolling Log No. 0301		
Recorder/Witness Tom	Crash: KNone □ Fatal □ Injury	□ Property		
coree's Name (Last, First, MD) OCKVOFT, Carolyn	9-9-60 F	Race Arresting Officer (I) Woodu	sard, Tom	MSP
Date Examined/Fime/Location August 6, 2000 0045		eath Results: 000 Refused strument # 1234	□ Refused	Urine Blood
Miranda Warning Given: Yes DNo Wha	t have you eaten today? When ICKENSOUP 806	lock Nothing?	How much?	Time of last drink?
Time now? When did you last sleep? How long! NidNight Last night 6 hos	Are you sick or injured?	□ Yes WNo	Are you diabetic or ep □ Yes % No	ileptic?
Do you take insulin?	Do you have any physical def	ects? □ Yes PNo	Are you under the car a doctor or dentist?	e of □ Yes #No
Are you taking any medication or drugs? • Yes		PONSIVE	Face	7 7 77
None of your business!	Breath Normal		Norma	
Speech	Eyes: O Reddened Conjun O Normal O Bloodshot	Watery ▼None □ L	Eye □REye	'racking: 7Equal □ Unequal
	il Size: *Equal nequal (explain)	Able to follow		Eyelids: Normal Droopy
Pulse & Time HGN		rtical Nystagmus? □ Yes V No	One Leg Stand	
1. 60 / 4050 Lack of Smooth Pursui	: Yes Yes co	nvergence Right Eye Left Eye		
2. 58 / 0105 Max. Deviation	Yes Yes	Right Bye	U C	1 9
3. 60 / 0//7 Angle of Onset	35° 35°		<u>_</u>	\\
To I To I Walle and Manual		Cannot Keep Balance		
Approx. Approx.		1 st Nine	2nd Nine Swa	ys While Balancing
1 1 1 1 Carried	فاستنصفهم	Stops Walking		
	,	Misses Heel-Toe	T & Use	s Arms to Balance
		Steps Off Line	б о ног	ping
	Section in	Italses Arms	Put	s Foot Down
	m m	Actual Steps Taken 9		
Internal Clock Describe Turn 46 Estimated At 30 Sec.		Cannot Do Test (Explain)	Type of Footw	
Staggered	to the right.	Room Light Darkness	Direct Nasal Area	1
○ Right △ Left			<u> </u>	
Draw lines from spot touched	Left Eye	4.0 6.0	3.5 Oral Cavity	1000
W.	Right Eye	1 ·		itai
U. Otto.		Yes Rebound Dilation I No Yes No	Reaction To Light 5/0 W	
(aux)		нент аям	LEFT ARM	
2			no RS	
4	\frac{\sqrt{3}}	A.	· ble	
(5)	26	NO VIS	ble Mark	\searrow
	mp 98.5.			7
Muscle Tone: □ Normal Flaccid □ Rigid Comments:			resh Puncture Marks	5
hat Medicine or Drug Have You Been Using?	How Much? Time of Use			
prother gave me. Kn	on't what don't rem	Eval Start Time	S house. Time Complete	d ,
AUG. 6, 2000 0015	10035	0045		7/25
Member Signature (Include Rayk)	ID No. 89		atkins	
Opinion of Evaluator: Rule Out ☐ Medical	☐ Alcohol ☐ Stimula ■ Depressant ☐ Hallucir	nt □ PCP	□ Inhalant □ Cannabis	

		DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>
LOG NO.		DRE: Vern Alley	ARRESTEE: Carolyn A. Cockroft
5. INITIAL OBSE	RVATIO	ESS 3. BREATH TEST 4. NOTIFICATION / DNS 6. MEDICAL PROBLEMS 7. PSYCHO I 10. SUSPECTS STATEMENTS 11. OPINIO	PHYSICAL 8. CLINICAL INDICATORS
1. LOCATION:	Examir	nation of Carolyn A. Cockroft took place in	the Intoxilyzer Room, 8th District Hqtrs, PhoenixPD
2. WITNESS: A	rresting	Officer - Tom Woodward #4532 Phoenix	PD
3. BREATH TI	EST: Of	ff. Woodward administered Intoxilyzer bre	ath test to Cockroft, the result was 0.00%
4. NOTIFICAT	ION/I	INTERVIEW of ARRESTING OFFICER	R: Writer was notified by Woodward that he
had arrested s	ubject f	or DUI, and suspected that she was "high o	n something". Off. Woodward further stated that the
subject had be	een driv	ing at 10 mph on the LaCienda Expressway	, and appeared dazed and stuporous.
She performe	d the S	FSTs poorly but exhibited no odor of an al	coholic beverage.
5. INITIAL O	BSERV	ATIONS: Writer observed subject in the I	ntoxilyzer Room, she was quiet, withdrawn and slow
to respond to	questio	ns. When walking towards the Intoxilyzer	she stumbled and nearly fell.
6. MEDICAL I	PROBL	EMS: None observed or stated.	
7. РЅҮСНОРН	YSICA	L TESTS: Romberg Balance: Subject ha	d approximately a 2" circular sway and estimated
46 seconds as	30 seco	onds. Walk and Turn: Subject lost balance	during the instructions, started to soon,
stepped off th	e line, n	nissed heel to toe, raised her arms, staggere	ed while turning and took (11) steps instead
of (9). One L	eg Stan	d: Subject swayed, raised her arms, hopped	and put her foot down. Finger to
Nose: Subject	missed	tip of his nose on each attempt.	
8. CLINICAL I	INDICA	ATORS: Subject exhibited HGN and lack of	of convergence. Pulse was below the normal
range. Systol	ic blood	l pressure was below the normal range. Pu	pils reacted slowly to light.
9. SIGNS of IN	GESTI	ON: None were evident	
10. STATEME	NTS: St	ubject admitted to taking "some medicine"	her brother gave her. She stated that she did not
know what t	he medi	cine was.	
11. OPINION o	f EVAI	LUATOR: In my opinion Carolyn Cockro	ft is under the influence of a CNS Depressant and
unable to op	erate a v	vehicle safely.	
12. TOXICOLO	OGICA	L SAMPLE: Subject provided a urine sam	ple.
13. MISCELLA	NEOU	S:	

Drug Influence Evaluation 000428 Rolling Log No. DRE No Evaluator 42,5 482 1X-2 SERGEANT DEB SCHOOLER None Recorder/Witness Crash: □ Property □ Fatal □ Injury R.C. STUDDARD, JACP TAP estee's Name (Last, First, MI) DOB Race Arresting Officer (Name, ID No.) C.D.# 8825 ال LAIRD 0311.59 APELLIZZERI MICHAEL Date Examined/Time/Location □ Refused Chemical Test Breath Results: □ Urine | Blood Instrument# □ Refused .05 DRE ROOM; A.I.B., SAN JOSE P.D., Miranda Warning Given: XYes No What have you eate 17833 What have you eaten today? When? Have you been drinking? How much? Time of last drink? "CHEESEBURGER" "LUNCH TIME" "A GLASS OF WINE ダノブロ Are you sick or injured? Are you diabetic or epileptic? Time now? When did you last sleep? How long? □Yes XNo □ Yes XNo 9 O'CLOCK" "LAST NIGHT" THRS Are you under the care of Do you take insulin? Do you have any physical defects? 🗆 Yes 💢 No a doctor or dentist? XYes □ No FOR STRESS" Coordination Are you taking any medication or drugs? Yes Do Attitude POUR, STAGGERING COOPERATIVE Breath Face BEVERAGE:
Blindness: NORMAL ATIMES A DAY ODOR OF ALCOHOLIC VALIUM ☐ Reddened Conjunctiva Tracking: □ Normal MBloodshot □ Watery None □ L Eye □ R Eye Equal □ Unequal SLULLED None Pu Corrective Lens: Able to follow stimulus: XYes □ No Pupil Size: KEqual Eyelids: □ Glasses □ Contacts, if so □ Hard □ Soft □ Unequal (explain) □ Normal Normal One Leg Stand HGN Left Eye Vertical Nystagmus? □ Yes X No Pulse & Time Right Eye 21/30 29/30 1, 60 /2130 Lack of Smooth Pursuit Convergence YES YES Right Eye Left Eye 2 60 12145 Max. Deviation YES YES 56,2157 30 30° Angle of Onset Walk and Turn Test "RUBBER LEGGED WALK" Cannot Keep Balance 🗸 🗸 Romberg Balance Starts Too Soon_ Approx. Арргох. 1st Nine 2nd Nine / Sways While Balancing Stops Walking /5/ Uses Arms to Balance Misses Heel-Toe Steps Off Line □ Hopping Raises Arms 1/2 COUSTANT // Puts Foot Down Actual Steps Taken Describe Turn Cannot Do Test (Explain) Type of Footwear Internal Clock LOST BALANCE GUA _Estimated At 30 Sec. NA RUNNING SHOES STAGGERED Pupil Size Room Light Darkness Direct Nasal Area ○ Right △ Left CLEAR Left Eye 6.5 3.5 Draw lines from spot touched 4.5 Oral Cavity Right Eye 4.5 6.5 3.5 CLEAC Rebound Dilation Hippus □ Yes Reaction To Light X No □ Yes 🕱 No Swi LEFT ARM /1` No VisiBLE MARKS 98.6. Blood Pressure__ | O 6 66 Temp_ Muscle Tone: X Normal ☐ Flaccid ☐ Rigid Attach Photos Of Fresh Puncture Marks Comments: Time of Use? (PIVI) Where Were The Drugs Used? (Location) at Medicine or Drug Have You Been Using? How Much? Juē's TAVERN Time DRE Notified ه 0<u>۰ د د د د د</u> MULIÄV - A COUPLE OF Date/Time of Arrest Time Completed Eval Start Time 2115 シンぐら 2045 Member Signature (Include Rank) Reviewed By: STUDDARD ID No. 17838 □ Inhalant Opinion of Evaluator: □ Rule Out Alcohol □ Stimulant Depressant □ Narcotic Analgesic □ Medical ## Hallucinogen □ Cannabis

	DRUG INFLUENCE EVALUATION	Page_2_of _2						
LOG NO.	DRE: Sergeant Deb Schroder	ARRESTEE: Impellizzeri, Michael T						
5. INITIAL OBSERVATION	1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.							
1. LOCATION: Exami	nation of Michael T. Impellizzeri, took plac	e in the DRE room San Jose PD Hdqtrs.						
2. WITNESS: Arresting	g Officer - C.D. Laird # 8825, Virgina Beach	PD. R.C. Studdard, IACP/TAP Representative						
3. BREATH TEST: W	riter observed Officer Laird administer GCI	breath test to Impellizzeri, the result was 0.05%						
4. NOTIFICATION / 1	INTERVIEW of ARRESTING OFFICER	: Writer was conducting DRE certification training						
at VBPD Hqtr. Offic	er Laird stated that he and Mr. Studdard had	d come upon the subject slumped in the driver's						
seat of a vehicle stop	oped in W/B traffic lane of S.R. #175, near t	he intersection with Snowden River Pkwy. Officer						
Laird further stated s	subject appeared to be very drunk and perfo	rmed poorly on the field sobriety tests.						
5. INITIAL OBSERVA	ATIONS: Writer observed subject seated in	a slumped position in a chair next to the GCI. Subj.						
was mumbling, sway	ing, and was slow to respond to my initial q	uestions.						
6. MEDICAL PROBL	EMS: Subject stated he was under the care	of a doctor for stress.						
7. PSYCHOPHYSICA	L TESTS: Romberg Balance: Subject swa	yed approximately 3" font to back and						
estimated 50 seconds	s as 30 seconds. Walk and Turn: Subject lo	st balance twice during the instructions, stepped off						
the line, missed heel	to toe, raised arms for balance, and staggere	ed while turning. One Leg Stand: Subject swayed,						
raised arms, and put	his foot down. Finger to Nose: Subject mis	sed tip of his nose on each attempt.						
8. CLINICAL INDIC	ATORS: Subject exhibited HGN and lack	of convergence. One of the pulse reading was below						
the normal range. B	Blood pressure was below the normal range.							
9. SIGNS of INGEST	TON: There was an odor of alcoholic bever	age on the subjects breath.						
10. STATEMENTS: St	ubject admitted to drinking wine and taking	some Valium pills. He stated that he takes Valium						
(4) times per day for	r stress.							
11. OPINION of EVAL	LUATOR: In my opinion Michael Impelliz	zer is under the influence of Alcohol and another						
CNS Depressant an	d unable to operate a vehicle safely							
12. TOXICOLOGICA	L SAMPLE: Subject agreed to provide a b	ood sample.						
13. MISCELLANEOU	S: Subject voluntarily produced a vial conta	nining which he identified as containing his Valium						
pills. He further state	ed that he had filled the prescription for (50	pills two days earlier. There were only 22 pills						
remaining.								

One Hour and Forty-Five Minutes

SESSION X

CENTRAL NERVOUS SYSTEM STIMULANTS

SESSION X CENTRAL NERVOUS SYSTEM STIMULANTS

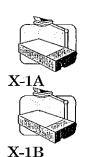
Upon successfully completing this session, the participant will be able to:

- o Explain a brief history of the CNS Stimulant category of drugs.
- o Identify common drug names and terms associated with this category.
- o Identify common methods of administration for this category.
- o Explain the symptoms, observable signs and other effects associated with this category.
- Explain the typical time parameters, i.e., onset and duration of effects, associated with this category.
- O State the clues that are likely to emerge when the Drug Evaluation and Classification Process is conducted for a person under the influence of this category of drugs.
- o Correctly answer the "topics for study" questions at the end of this Section.

Content Segments		Learning Activities	
A.	Overview of the Category	0	Instructor Led Presentations
В.	Possible Effects	0 .	Review of Drug Evaluation and Classification Exemplars
C.	Onset and Duration of Effects	0	Reading Assignments
D.	Overdose Signs and Symptoms	O	Video Presentations
E.	Expected Results of the Evaluation	0	Slide Presentations

Aides	Lesson Plan	Instructor Notes
	CENTRAL NERVOUS SYSTEM STIMULANTS	Total Lesson Time: Approximately 105 Minutes
CONTRACTOR OF THE PROPERTY OF		Session title on wall chart.
X-0A&B		Briefly review the objectives, content and activities of this session.
(Session Objectives)		
	A. Overview of the Category	
25 Minutes		
	CNS Stimulants speed up the operation of the Central Nervous System.	
	a. "Speed Up" does <u>not</u> mean "improve".	Emphasize that abuse of CNS Stimulants does not make the brain work "better" or "smarter". Rather, they induce the brain to cause many of the body's organs to work harder, but not better.
	b. The "speeding up" results in increased heartbeat, pulse, respiration, blood pressure and temperature.	However: Robert Louis Stevenson wrote "The Strange Case of Dr. Jekyll and Mr. Hyde" while under the influence of cocaine. He wrote sixty thousand words in six days.
	c. All of these effects can lead to physical harm to the stimulant user.	
	d. The "speeding up" also produces nervousness, irritability and an inability to concentrate or think clearly.	

Instructor Notes



X-1C

- e. These psychological effects can lead to unpredictable and bizarre behavior by the stimulant user.
- 2. There are three major subcategories of Central Nervous System Stimulants.
 - a. <u>Cocaine</u>

b. The Amphetamines

Examples:

- o Methamphetamine
- o Amphetamine Sulfate
- o Biphetamine
- o Desoxyn

c. Others

- o Ritalin (methylphenidate hydrochloride)
- o Preludin (phenmetrazine hydrochloride)
- o Cylert (pemoline)
- o Ephedrine
- o Caffeine

Point out that the Amphetamines include a large number of individual drugs, only a few of which are listed on Visual X-1.

Point out that there are many "other" CNS Stimulants (i.e., non-Cocaine and non-Amphetamines); the ones listed on the visual are only a few of those.

Point out that we will focus on Cocaine and the Amphetamines, because they are the most widely abused CNS Stimulants. But, the students should be aware that there are other stimulant drugs.

Coca plant: Scientific name "Erythroxylon Coca".



X-2 ("Coca Plant")

- 3. Cocaine derives from the <u>coca</u> plant.
 - a. The plant is native to South America.

		-		
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Instructor Notes

- b. Cocaine is made from the leaves of the coca plant.
 - plant. | NOTE:
 not be co
- c. Archaeological evidence indicates that natives of Peru chewed coca leaves 5,000 years ago.
- d. Sigmund Freud personally experimented with Cocaine for approximately 3 years.
- e. Small quantities of cocaine originally were included in the formula for Coca Cola.
- 4. Amphetamines were first synthesized near the end of the 19th Century.
 - a. The first use of
 Amphetamines for medical
 purposes began in the
 1920's.
 - b. Initial medical application was to treat colds.
 - Amphetamines cause the nasal membranes to shrink.
 - o This gives temporary relief from stuffy nasal passages.

NOTE: the coca plant should not be confused with the cocoa plant, from which chocolate is made.

Use of Cocaine in products such as Coca Cola was outlawed by the Pure Food and Drug Law of 1906.

Point out that much more effective drugs have been developed to treat cold symptoms. Amphetamines are no longer prescribed as cold remedies.



X-3 ("Sample Amphetamin es and their Medical Purposes")

Aides	Lesson Pla	Instructor Notes
	c. Present day m purposes for a include:	
	o control syn narcolepsy o control cer	disorder that causes the individual to fall asleep compulsively, often several
		e behavioral Example: Ritalin or Cylert are commonly prescribed for children diagnosed with ADD or similar disorders.
	to perform	allow persons Force previously gave pilots
	o treat mild	depression
	o control app	Many over the counter appetite control products contain CNS Stimulants as their active ingredient.
	o antagonize Depressant	the effects of drugs are antagonistic when the signs and symptoms of one are opposite to the signs and symptoms of the other.
	o prevent and surgical sh	.
	o maintain b during surg	lood pressure gery
	o treat Parki Disease	nson's Parkinson's Disease: a form of paralysis characterized by muscular rigidity, tremor and weakness.

Aides	Lesson Plan	Instructor Notes
	o enhance the action of certain analgesic (pain killer) drugs	·
	d. Numerous pharmaceutical companies manufacture Amphetamines for these purposes.	
X-4 ("Pharmaceutical Amphe-	e. Examples of common pharmaceutical Amphetamines.	
tamines")	, , , , , , , , , , , , , , , , , , ,	
	o <u>Dexedrine</u> (dextroamphetamine sulfate) used to treat narcolepsy and hyperkinetic behavior, and for weight control. (Street names "Dexies",	Manufactured by Smith, Kline and French. NOTE: Dexedrine probably is the most commonly prescribed Amphetamine.
_	"Hearts")	
	o Benzedrine (Amphetamine sulfate) used to treat narcolepsy, hyperkinetic behavior and weight problems. (Street names "Bennies", "Whites", "Cartwheels")	Manufactured by Smith, Kline and French.
	o Biphetamine (combination of dextroamphetamine and Amphetamine) used in weight reduction. (Street name "Black Beauty")	Manufactured by Pennwalt Pharmaceutical Company.

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Instructor Notes

- Desoxyn (Methamphetamine hydrochloride, also known as desoxyephedrine) used in weight reduction.
- Pharmaceutical combinations of Amphetamines and CNS Depressants.
 - Dexamyl (combines dextroamphetamine sulfate and amobarbital)

- Eskatrol (combines dextroamphetamine sulfate and prochlorperazine)
- 5. Large quantities of Amphetamines are also illegally manufactured in this country.
 - The two most commonly abused illicit Amphetamines are Methamphetamine and Amphetamine sulfate.

Manufactured by Abbott Pharmaceutical Company.

If available: display 35mm slides of these various drugs.

Point out that one of the most undesirable side effects of pharmaceutical Amphetamines is insomnia. Certain manufacturers offer drugs that combine CNS Depressants with Amphetamines to offset the insomnia.

Manufactured by Smith, Kline and French.

Remind students that amobarbital is a derivative of barbituric acid.

Manufactured by Smith, Kline and French.

Prochlorperazine is a nonbarbiturate CNS Depressant.

If available: display 35mm slides of these various drugs.



("Pharmaceut ical Combinations of Amphetamines and CNS Depressants")



X-6 ("Illicit Amphetamin es")

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Instructor Notes



- b. Methamphetamine is also known as Methedrine.
- c. Its more common "street names" are "speed"; "crank"; "ice"; "crystal"; "meth"; and, "water".
- 6. There are some other CNS Stimulants, apart from Cocaine or the Amphetamines.
 - a. Preludin is a licitly manufactured CNS
 Stimulant that is not an Amphetamine:
 - o generic name phenmetrazine hydrochloride
 - o used in weight control
 - o has all of the basic effects of amphetamine
 - b. Ritalin is another licitly manufactured, non-Amphetamine CNS Stimulant:
 - o generic name <u>methylphenidate</u> hydrochloride
 - o used to treat mild
 depression, hyperkinetic
 behavior, narcolepsy
 and drug induced
 lethargy produced by
 CNS Depressants.

If available: display 35 mm slides of illicitly manufactured methamphetamine and amphetamine sulfate.

Manufactured by Boehringer Ingelheim.

Manufactured by Ciba Pharmaceutical Company.

Ask students if they know of any children for whom ritalin has been prescribed.

- o has many of the basic clinical effects of Amphetamine.
- c. <u>Cylert</u> is a third licitly manufactured, non-Cocaine and non-Amphetamine CNS Stimulant:
 - o generic name Pemoline.
 - o used to treat Attention Deficit Disorder (ADD), also known as "hyperactivity".
 - o has many of the basic clinical effects of Amphetamine.
- d. Ephedrine is a licitly manufactured stimulant used in diet aides, body building supplements. It can also be found in herbal teas and preparations.
- e. Cathine and Cathinone are the two psychoactive chemicals derived from the Khat plant. It originates from the sub-Sahara regions of Africa.
- f. Methcathinone is illicitly manufactured from common household chemicals.
 Effects are very similar to methamphetamine.
- 7. Methods of ingestion of CNS Stimulants.

If available: display 35mm slides of Preludin and Ritalin.

Manufactured by Abbott Laboratories.

Remind the students that we will focus on Cocaine and the Amphetamines for our discussion of CNS Stimulants and their effects.

Aides

Lesson Plan

Instructor Notes



X-8A (Methods of Ingestion)

- a. There are a variety of ways in which the different CNS Stimulants may be ingested.
- b. <u>Cocaine</u> is commonly insufflated (snorted), smoked, injected and taken orally.
- c. In order to be smoked, a pure form of Cocaine is required.
 - o Much of the Cocaine sold in this country is mixed with other materials, or chemically bonded to other elements.
 - o Various chemical processes can be used to "free" the Cocaine from other elements and impurities.
 - o One such process produces pure Cocaine in the form of small chunks.
 - o These chunks are known as "Crack" or "Rock Cocaine".
- d. Licitly manufactured

 <u>Amphetamines</u> are taken
 orally, in the form of
 tablets, capsules and liquid
 elixirs.

NOTE: the term "Crack" derives from the cracking sound produced when the chunks are burned for smoking.



- Illicitly manufactured Methamphetamine most commonly is injected or smoked but sometimes may be snorted or taken orally.
- f. Another crystalline. smokeable form of Methamphetamine is known as "Crystal Meth"; it is very similar to "Ice".
- g. Illicitly manufactured Amphetamine sulfate usually is produced in tablet form (called "Mini bennies") and is taken orally.
- В. Possible Effects
 - 1. Both Cocaine and the Amphetamines produce euphoria, a feeling that there are no problems.
 - a. A feeling of super strength, and of absolute self confidence may also be present.
 - b. With Cocaine, but not with Amphetamines, there is an anesthetic effect, and the dulling of pain may contribute to the euphoria.
 - 2. Stimulant users tend to become hyperactive, indicated by a nervousness, extreme talkativeness, and an inability to sit still.

Point out that bruising often will be seen around a Methamphetamine injection site.

Point out that "Ice" tends to be clear in appearance, while "crystal meth" is less pure and has a cloudy appearance.

Solicit students' questions and comments about the overview of CNS Stimulants.



Aides	Lesson Plan	Instructor Notes
	3. CNS Stimulants tend to release inhibitions, allowing users to commit acts that they normally would avoid.	
	4. Stimulant users <u>misperceive</u> <u>time and distance</u> .	Example: To the subject, time seems to be speeded up, so that 2 hours may seem like 2 minutes.
	5. Persons under the influence of CNS Stimulants become easily confused, and lose the <u>ability to concentrate</u> or to think clearly for any length of time.	Point out that this lack of concentration makes it very difficult for the user to perform divided attention tests successfully.
		Solicit students' questions and comments concerning possible effects of CNS Stimulants.
	C. Onset and Duration of Effects	
10 Minutes		
	 The onset and duration of effects are quite different for Cocaine as compared to the Amphetamines. Generally speaking, Cocaine's effects are much briefer than are 	
	Amphetamine's.	
	b. The time parameters of Cocaine vary with the method of ingestion.	Note: Subjects that have ingested both Cocaine and Alcohol will produce a metabolite known as "Cocaethylene". Which has a half-life of four hours possibly extending the effects of Cocaine longer than the norm.

Instructor Notes



X-10 ("Cocaine Time Factors")

- 2. When Cocaine is <u>smoked</u>, or "freebased", the drug goes immediately to the lungs, and is absorbed into the blood stream very rapidly.
 - a. The smoker begins to feel the effects of the Cocaine virtually immediately.
 - b. The "rush", or euphoria, is reported to be very intense.
 - However, the euphoric effects only last 5-10 minutes after the Cocaine is smoked.
- 3. When Cocaine is <u>injected</u>, the drug is passed directly to the blood stream, where it is carried swiftly to the brain.
 - a. The effects are felt within seconds.
 - b. The onset of effects is very intense.
 - c. The effects usually continue to be felt for 45-90 minutes.
- 4. When Cocaine is <u>snorted</u> (insufflated), the onset of effects is not quite as rapid as with smoking or injecting.
 - a. The user typically feels the onset of effects within 30 seconds after snorting the drug.
 - b. Although the "rush" occurs, it is not quite as intense as it is when the Cocaine is smoked or injected.

Note: Injection sites will be discussed in Session XVII (Narcotic Analgesics).

Point out that snorting remains a very popular method of ingesting Cocaine.

- c. The effects from snorting usually last from 30-90 minutes.
- 5. <u>Oral</u> ingestion of Cocaine usually is the least preferred method.
 - a. The user generally does not begin to feel the effects for 3-5 minutes.
 - The effects are not as intense as they are with other methods of ingestion.
 - c. However, the effects may last 15-30 minutes longer than with other methods.
- 6. With all methods of ingestion, the duration of Cocaine's effects tend to be briefer than the effects of most other drugs.
 - a. As the effects wear off, it becomes very difficult to observe evidence of impairment.
 - b. If the suspect is not evaluated by a Drug Recognition Expert fairly soon after the suspect has been apprehended, the DRE may not uncover evidence of the CNS Stimulant.
- When Methamphetamine is <u>injected</u>, the initial effects are very similar to the injection of Cocaine.
 - a. The user begins to feel the effects within a few seconds.

<u>Clarification</u>: the effects of Cocaine taken orally may last from 45-120 minutes.

Point out that it is very possible that a Cocaine user may not be examined by a DRE until at least 30 minutes following the suspect's use of Cocaine. Often, much more time will have elapsed. For this reason, Cocaine use can be difficult to ascertain from a drug evaluation and classification examination.



X-11 ("Methamphe tamine Time Factors")

A	٠	7		
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Instructor Notes

- b. The "rush" is very intense, and lasts at a high level of intensity for 5-30 seconds.
- c. Unlike Cocaine, Methamphetamine's effects are long lasting: the user stays "high" or "wired" for 4-8 hours following injection.
- 8. When Methamphetamine is smoked, the rush is very intense, and the effects are long lasting, i.e., up to 8 hours or longer.
- When Methamphetamine is snorted or taken orally, the onset takes longer, the rush is much less intense, and the effects are much briefer.
- D. Overdose Signs and Symptoms
 - 1. Overdoses of Cocaine or Amphetamines can cause the pleasurable effects to turn into panic and often violent behavior.

 If the overdose is caused by Cocaine, it is commonly referred to as Cocaine Psychosis or Cocaine Delirium.
 - a. Subject may become very confused and aggressive.
 - b. Subject may suffer convulsions and faint or pass into a coma.
 - Heartbeat (pulse) will increase, possibly dramatically.
 - d. Hallucinations may occur.

Solicit students' comments and questions concerning time parameters of Cocaine and Methamphetamine.

Write on Flipchart "Cocaine Psychosis or Cocaine Delirium".



5 Minutes



Aides

Instructor Notes

- 2. Death can occur from sudden respiratory failure, or from heart arrhythmia, leading to cardiac arrest.
- 3. Another danger is that subjects may attempt to treat CNS Stimulant overdose with Barbiturates, possibly leading to overdose of CNS Depressants.

Example: The feeling that bugs are crawling under the skin is also known as "Coke Bugs".

Note: It is important that officers are aware of this to avoid in custody deaths.

Solicit students' comments and questions concerning overdoses of CNS Stimulants.



60 Minutes



X-12A ("SFST Evidence")

- E. Expected Results of the Evaluation
 - 1. Observable evidence of impairment.
 - a. Standardized Field Sobriety Tests.
 - Horizontal Gaze Nystagmus will not be present with suspects under the influence of CNS Stimulants.
 - Vertical Gaze Nystagmus will not be present.
 - Performance on Walk and Turn and One Leg Stand may be impaired due to the suspect's hyperactivity and inability to concentrate.

Example: suspect may start too soon on Walk and Turn, and may tend to walk fast, thus losing balance or missing heel to toe. Suspect may also count very rapidly on the one leg stand test.

Instructor Notes



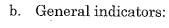
("General Indicators")

Performance on Nose tests will be impaired.

Romberg and Finger to

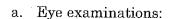
Point out that CNS Stimulants impair the user's perception of time, so that the subject's estimate of 30 seconds, on the Romberg test, may be speeded up.

Also, his or her finger movements may be abrupt, jerky and inaccurate.



- restlessness O
- 0 anxiety
- euphoria o
- talkativeness 0
- irritability
- runny nose 0
- 0 redness to nasal area
- grinding teeth, bruxism
- 0 leg tremors
- eyelid tremors

Evidence associated with the physiologic examinations.



- Lack of Convergence will not be evident
- pupils generally will be dilated
- pupil reaction to light generally will be slow
- Vital signs examinations:
 - blood pressure generally will be elevated
 - pulse generally will be increased

NOTE: Indicators associated with the nasal area may be evident if the subject is in the habit of snorting Cocaine.

Point out that the technical term for "dilated pupils" is Mydriasis.



X-12C ("Eye Examinations")



X-12D ("Vital Signs Examinations")

Aides	Lesson Plan	Instructor Notes
	o body temperature generally will be elevated	
X-13 ("CNS Stimulant Symptoma- tology Chart")	3. Summary	
	4. Demonstrations	
	a. Video tape demonstrations	Show video tape of subject(s) under the influence of CNS Stimulants. Relate behavior/observations to the CNS Stimulant Symptomatology Chart.
	b. Drug Evaluation and Classification exemplar demonstrations.	Refer students to the exemplars found at the end of Section X in their student manuals.
	·	Relate the items on the exemplars to the CNS Stimulant Symptomatology Chart.
		Solicit students' questions or comments concerning expected results of the evaluation of subjects under the influence or CNS Stimulants.

Session X

Central Nervous System Stimulants





Central Nervous System Stimulants

Upon successfully completing this session, the participant will be able to:

- Explain a brief history of the CNS Stimulant category of drugs
- Identify common drug names and terms associated with this category
- Identify common methods of administration for this category
- Explain the symptoms, observable signs and other effects associated with this category

Central Nervous System Stimulants (continued)

- Explain the typical time parameters, i.e., on-set and duration of effects, associated with this category
- State the clues that are likely to emerge when the drug evaluation and classification process is conducted for a person under the influence of this category of drugs
- Correctly answer the "Topics for Study" questions at the end of this section

Drug Evaluation & Classification Training

Types of CNS Stimulants

Cocaine









Types of CNS Stimulants (continued)



- The Amphetamines
 - Methamphetamine
- Amphetamine Sulfate
- Biphetamine
- Desoxyn









Types of CNS Stimulants (continued)

- Others
 - Ritalin
 - Preludin
 - Cylert
 - Ephedrine
 - -- Caffeine

Coca Plant





"Erythroxylon Coca"

Date Featurion & Classification Training

X-2

Medical Uses of Amphetamines

- · Control symptoms of narcolepsy
- · Control hyperactivity in children
- · Relieve or prevent fatigue
- · Treat mild depression
- Control appetite
- · Antagonize effects of depressants
- · Prevent and treat surgical shock
- · Maintain blood pressure during surgery
- · Treat Parkinson's disease
- Enhance the action of analgesic drugs

Drug Evaluation & Classification Training

¥.3

Commonly Prescribed Pharmaceutical Amphetamines

- · Dexedrine
- · Biphetamine
- Dextroamphetamine Sulfate
- Combination of Amphetamine and Dextroamphetamine
- Smith, Kilne and French
- Pennwalt Pharmaceutical
- Benzedrine
- Desoxyn
- Amphetamine Sulfate
- Methamphetamine Hydrochloride
- Smith, Kline and French
- Abbott Pharmaceutical

Drug Evaluation & Classification Training

Pharmaceutical Combinations of Amphetamines and CNS Depressants

Dexamyl

Dextroamphetamine Sulfate and Amobarbital

Eskatrol

Dextroamphetamine Sulfate and Prochlorperazine

Drug Evaluation & Classification Training

X-5

Commonly Abused Illicit Amphetamines

Methamphetamine





Amphetamine Sulfate

Drug Evaluation & Classification Training

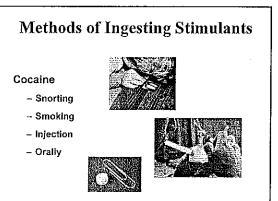
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Other CNS Stimulants (Besides Cocaine or the Amphetamines)

- · Preludin
 - Phenmetrazine Hydrochloride
 - Boehringer Ingelheim Co.
- Ditalia
- Methylphenidate Hydrochloride
- Ciba Pharmaceutical
- Cylert
 - -- Pemoline
- Abbott Laboratories

Drug Evaluation & Classification Training

X-7



Methods of Ingesting Stimulants (continued)

- Methamphetamine
 - Injection
 - Snorting
 - Orally

 - Smoking





- · Other Amphetamines
 - Orally (tablets, capsules, etc.)



Possible Effects of **CNS Stimulants**

- · Euphoria
- Hyperactivity
- · Release of inhibitions
- · Misperception of time and distance
- · Inability to concentrate

Cocaine Time Factors

- · Smoked (freebase)
 - Virtually immediate
- Very intense "rush"
- Effects last 5-10 minutes
- Intense "rush"

Snorted

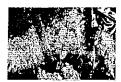
→ Effects last 30-90 minutes

- Effects are felt within 30

- Injected
 - Effects are felt within seconds
 - Very intense "rush"
- Effects last 45-90 minutes
- Orally
 - Effects begin within 3-5 minutes
 - Effects are less intense
- Effects last 45-120 minutes

Methamphetamine Time Factors

- · Effects are felt within seconds
- · "Rush" is very intense for 5-30 seconds
- · Effects can last up to 12 hours



Evaluation of Suspects Under the Influence of CNS Stimulants

SFST Evidence:

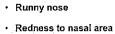
- · HGN or VGN none
- · Impaired performance will be evident on Walk and Turn and One Leg Stand
- · Impaired performance will be evident on Romberg and Finger-to-Nose

Evaluation of Suspects Under the Influence of CNS Stimulants

General indicators:

If subject snorts cocaine:

- Restlessness
- Anxiety
- Euphoria
- Talkativeness
- · Irritability
- Bruxism
- · Eyelid and leg tremors





Evaluation of Suspects Under the Influence of CNS Stimulants

Eye Examinations:

- · Lack of Convergence none
- · Pupils will be dilated (Mydriasis)
- · Pupillary reaction to light will be

Drug Evaluation & Classification Training

Evaluation of Suspects Under the Influence of CNS Stimulants

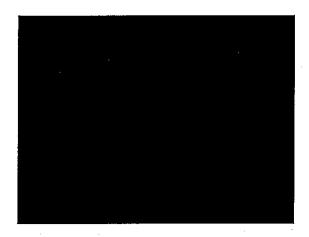
Vital Signs:

- Blood pressure will be <u>up</u>
- · Pulse will be up
- · Body temperature will be up

CNS Stimulant Symptomatology Chart

HGN	None
VGN	None
Lack of Convergence	None
Pupil Size	Dilated (mydriasis)
Reaction to Light	Slow
Pulse Rate	Up
Blood Pressure	Up
Temperature	Up
Muscle Tone	Possibly rigid

Drug Evaluation & Classification Training



	Drug Influence Evaluation									
	Evaluator Recorder/Witness	14	DRE No		Rollin	g Log No.				X -1
	Recorder/Witness		Crash: □ Fatal) D'Inj	one jury	□ Pro	perty			
ĺ	'estee's Name (Last, First, MI)		DOB	Sex	Race		resting Office			
	Date Examined/Time/Location	JES R	7.10.0	~3 M	<u>w</u>	\subseteq	NALE	2. 1	4927	UST Blood
			PAL TES	ST1 201-5	Breath Res	+ #	□ Refused′	C	hemical Test □ Refuse	□ Urine)ズ Blood d
	7-8-96 223c Miranda Warning Given: AYes By:	s No What	have you eaten	today? V	Vhen?	Have you	ı been drinki	ing?	How much?	Time of last drink?
	By: Time now? When did you last	sleep? How long?	Are you sic	<u>مرن کو</u> k or injured?	<u> 10 vscy (</u>	ت ت الاes الا	フナ・・ン(コ No		Are you diabetic o	r enilentic?
	Bo'crock LAST NICH	HT 3HRS	7.70			77 554			□ Yes XNo	
				e any physica	il defects?	⊺Yes ≱X	No		Are you under the a doctor or dentist	
	Are you taking any medication or	drugs? □ Yes 🕱 No		PATIVE					Coordination Poor≥ STU	mer wa
			Breath NORM]	Face NORMAL	
	Speech		Eyes; □	Reddened Co			Blindness:			Tracking: X Equal D Unequal
	PAPID NEWOUS Corrective Lens: None □ Glasses □ Contacts, if so □ H:	Pupil	Size: X Equa equal (explain)	1					ilus: X Yes □ No	Eyelids:
	Pulse & Time HG	IN	Left Eye	Right Eye	Vertical Ny	stagmus?	□ Yes XN	lo	One Leg Sta	nd
	1. 112 / 2240 Lac	k of Smooth Pursuit	No	NO .	Convergence	ce			(3)	(a)
	2. 108 / 2253 Max	x. Deviation	NO	NO	Right 1	Eye	Left Eye		(A)	1 ()
	3. 100 / 2305 Ang	de of Onset	NONE	riorje	-				0	, 🖣
	Romberg Balance	Walk and Turn Te					ep Balance	//	7	30 45/30
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(3, 03, 04, 00,					tops Walk George Woo	·			ses Arms to Balance
					Steps Off Line Raises Arms Hopping					
		3 4	a Dirticoc		Α.	ctual Step			- d d b	uts Foot Down
	Internal Clock 15 Estimated At 30 Sec.	Describe Turn て のいこと のい			1)o Test (E	xplain)		Type of Foot	
	Estimated At 30 Sec.	COICE MICH	.646.61		A/A			T: .	COM BA	•
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	4	三	3	NO						
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	(5)	Talenta and the second of the	<u> </u>	MAR	KS			_		
h	Blood Pressure 142	GL. Temp	99.9.					-		
	Muscle Tone: Normal Flacci Comments:			1 _		Attach :	Photos Of	Fresh I	Puncture Mark	s
. . (t Medicine or Drug Have You	Been Using? F	How Much?	Time of U	Jse?	Where V	Vere The Dru	igs Used?	(Location)	
1	NOTHING DUX			12/		13/4		J		
l	Date/Time of Arrest	Time	DRE Notified	حجج	c,	Eval Sta		 23©	Time Complet	ed 2310
Ì	Member Signature (Include Rank)	o	۷	I II) No	3321	Reviewe	d By:	الاد (LITEK	
ľ	Opinion of Evaluator:	Rule Out ∪∈	Alcohol Depressant	≰Stimı □ Hallu	ılant	D PCP	tic Analgesic		□ Inhalant □ Cannabis	
- [_		- opromant	~ rrantu		- 114100	" v mear Repire		- Jannavia	i

LOG NO.	5.25 0.27 0.34	TION Page 2 of 2
1. LOCATION 2. WIT	DRE: Officer Ron Moen	ARRESTEE: James R. Hedlund
5. INITIAL OBSERVA	TNESS 3. BREATH TEST 4. NOTIFICAT TIONS 6. MEDICAL PROBLEMS 7. PS ON 10. SUSPECTS STATEMENTS 11. (TION / INTERVIEW ARRESTING OFCR. SYCHOPHYSICAL 8. CLINICAL INDICATORS OPINION 12. TOXICOLOGY SAMPLE 13. MISC.
1. LOCATION: Exa	mination of James R. Hedlund took plac	ce in the DRE Room, 3rd Precinct, Tucson PD
2. WITNESS: Arrest	ing Officer - Officer R. Engle, #2309 Tu	icson PD
3. BREATH TEST:	Officer Engle administered Intoxilyzer	breath test to Hedlund, the result was 0.00%
4. NOTIFICATION	/ INTERVIEW of ARRESTING OF	FICER: Writer was notified by Officer Engle immediately
upon completion o	f the breath test. Officer Engle stated st	ubject had been apprehended for driving 110/65 zone and
driving without hea	adlights.	
5. INITIAL OBSER	VATIONS: Writer observed subject in	the DRE room sitting next to Officer Engle. Subject rocked
rocked back and fo	orth while seated on the bench.	
6. MEDICAL PROE	BLEMS: None observed or stated.	
7. PSYCHOPHYSIC	CAL TESTS: Romberg Balance: Subje	ect swayed approximately 3" front to back and estimated 15
seconds as 30 secon	nds. Walk and Turn: Subject started to	soon, lost balance during instructions, raised his arms,
and turned in an ab	rupt swivel. One Leg Stand: Subject sw	vayed, raised his arms, hopped and put his foot down.
Finger to Nose: Sub	pject missed tip of his nose on each atten	npt with his right hand.
8. CLINICAL INDIC	CATORS: Subject's pulse, blood press	sure and temperature were above the normal range. His
pupils were dilate	d and reacted slowly to light.	
9. SIGNS of INGEST	FION: Subjects nostrils were found to	contain a residue of white powder.
10. STATEMENTS:	Subject denied taking any medicine or	drugs. When asked, "how much coke did you snort
tonight?" Subjec	t stated "I won't answer that"	
11. OPINION of EV	ALUATOR: In my opinion James R. F	Hedlund is under the influence of a CNS Stimulant and
unable to operate a	a vehicle safely.	
2. TOXICOLOGIC	CAL SAMPLE: Subject agreed to provi	ded a blood sample.

	000455		Drug	Influe	nce Eva				
	Evaluator JOHN, C		DRE No		Rollin	ig Log No.			
_	Recorder/Witness		Crash: □ Fatal	□ N □ In	one ijury	□ Pro	pperty		
	restee's Name (Last, First, MI)	سد. و ه	DOB	Sex	Race	- 1	resting Officer (Nar	ne, ID No.)	
*	KOHLHEPP KIN		03/24/	73 F			ROBERTS.	R # 20	168 CPD
	Date Examined/Time/Location		•		т.	sult s).e v	o neruseu - r	Chemical Test ☐ Refus	O Orme Blood
	OCT 10, 1996 - 2313 Miranda Warning Given: ■Yes □ No	What	have you eate	n today?	When?	Have you	u been drinking?	How much?	Time of last drink?
						/	ATHING.		NIA
	Time now? When did you last sleep?	How long?	Are you sid	k or injured?	- 	O Yes 🙀	No	Are you diabetic o	r epileptic?
	MIDNIGIT YESTER DAY Do you take insulin?	AHRS						□ Yes 🌞 No	
	Do you take insulin?	#No	Do you hav	ve any physic	al defects?	□Yes 🗳	No	Are you under the a doctor or dentist	
	Are you taking any medication or drugs?	Yes •• No			 		· <u>- · </u>	Coordination	
		а.	Breath	_	Bist	Les	1655	Face JII	TERY, STUMBLU
	I DONT DO DRUG Speech VERY TALKATIVE	<u> 5</u>		NORM	44			NOR	MAL
				Reddened Co Bloodshot			Blindness: Mone □ L Eye	e □R Eye	Tracking:
i	RAPID " TRIPPING" OVER CU Corrective Lens:	DVZDC Punit S	Size: € Equa	1)				mulus :≉ Yes □ No	Eyelids:
	□ Glasses □ Contacts, if so □ Hard □ S		qual (explain)				Able to lollow sti	midius## Tes 140	Eyends; Normal □ Droopy
	Pulse & Time HGN		Left Eye	Right Eye	Vertical Ny	stagmus?	□ Yes ÞNo	One Leg Sta	ind (22)
	1. 100 / 2320 Lack of Smo	oth Pursuit	NO	NO	Convergence Right		Left Eye]	7
	2. 103 2331 Max. Deviat	ion	NO	NO	Itight,	Laye .	Left Eye		9_
-	3. 104 / 2343 Angle of Ons	et	NOVE	Nout				₩	(A)
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								LEFT ARM	
	Blood Pressure 144 / 104	Temp_	99.8		2				2
ľ	Muscle Tone: ♦Normal □ Flaccid □ Ri Comments:	gid				Attach l	Photos Of Fresh	ı Puncture Mark	ıs
	t Medicine or Drug Have You Been Usi	ng? Ho	ow Much?	Time of I	Jse?	Where W	ere The Drugs Use	d? (Location)	
1	I DOUT USE DAUGS	ANYANO	21.21.		USC D		REFUSE		
	Date/Time of Arrest	Time I	DRE Notified	1 101	· · / · · · · ·	Eval Sta	rt Time	Time Complet	ęd.
+	Member Signature (Inclyde Rank)		<u> </u>	ID No	フラミ	Reviewe	45/5 d By:	<u>  659</u>	5
L	Opinion of Evaluator:   ORUE Out	40h	Alcohol	<b>⊅</b> Stimu	1766	□ PCP	-166	O Inhalant	
	□ Medical		Depressant		cinogen		ic Analgesic	□ Cannabis	

000456		
	DRUG INFLUENCE EVALUA	TION Page 2_of 2
LOG NO.	DRE: Officer Clark John	ARRESTEE: Kim J. Kohlhepp (m)
5. INITIAL OBSERVATION	ONS 6. MEDICAL PROBLEMS 7. PS	TION / INTERVIEW ARRESTING OFCR. LYCHOPHYSICAL 8. CLINICAL INDICATORS OPINION 12. TOXICOLOGY SAMPLE 13. MISC.
1. LOCATION: Exami	nation of Kim J. Kohlhepp took place	e in the DRE Room, 3rd Precinct, Alburqueque PD
2. WITNESS: Arresting	Officer - Officer R. Roberts, #8712	Alburqueque PD
3. BREATH TEST: Of	fficer Roberts administered Intoxilyze	er breath test to Kohlhepp, the result was 0.00%
4. NOTIFICATION / I	INTERVIEW of ARRESTING OF	FICER: Writer was notified by Officer Roberts immediately
upon completion of the	ne breath test. Officer Roberts stated	subject had been apprehended for driving 65/30 zone,
failure to stop for a tr	affic signal and driving without headl	ights.
5. INITIAL OBSERVA	ATIONS: Writer observed subject in	the DRE room standing next to Officer Roberts.
When told to sit dow	n, subject stood up again within seven	ral seconds and fidgeted from foot to foot.
6. MEDICAL PROBL	EMS: None observed or stated.	
7. PSYCHOPHYSICA	L TESTS: Romberg Balance: Subje	ect swayed approximately 2" side to side and estimated 12
seconds as 30 seconds	s. Walk and Turn: Subject stepped of	ff the line, raised his arms, and turned in an abrupt (about
face) One Leg Stand:	Subject swayed, raised his arms, hop	ped and put his foot down. Finger to Nose: Subject missed
tip of his nose on each	ı attempt.	
8. CLINICAL INDICA	TORS: Subject's pulses, blood pres	sure and temperature were above the normal range. His
pupils were dilated an	d reacted slowly to light.	
9. SIGNS of INGESTIC	ON: Subjects nostrils were found to	be red and ulcerated.
10. STATEMENTS: Su	bject denied ever using drugs. Subse	equently stated "I don't use drugs anymore"
11. OPINION of EVAL	UATOR: In my opinion Kim Kohlh	epp is under the influence of a CNS Stimulant and unable
to operate a vehicle s	afely.	
12. TOXICOLOGICAL	SAMPLE: Subject agreed to provi	ded a blood sample.
13. MISCELLANEOUS	S: There is an outstanding bench was	rrant on the subject Kim J. Kohlhepp, for failure to appear
on a charge of posses	sion of methamphetamine.	
		/

One Hour

## SESSION XI

PRACTICE: EYE EXAMINATIONS

## SESSION XI PRACTICE: EYE EXAMINATIONS

Upon successfully completing this session, the participant will be able to:

- o Conduct examinations of pupil size and reaction to light, under both lighted room and darkened room conditions.
- o Articulate the eye examination procedures.
- o Document the results of the examinations of pupil size and reaction to light.

Content Segments			<u>Learning Activities</u>		
A.	Procedures For This Session	0	Instructor Led Presentations		
B.	Room Light Examinations	O	Students' Hands On Practice		
C.	Dark Room Examinations	0	Instructor Led Coaching		
D.	Session Wrap Up	0	Student Led Coaching		





XI-0 (Session Objectives)



10 Minutes

PRACTICE: EYE EXAMINATIONS

- A. Procedures For This Session
  - 1. Participants will work in three or four member teams.
    - a. At any given time, one member of the team will be engaged in conducting and recording eye examinations of another member.
    - The remaining member(s)
       will help coach and critique
       the student who is
       conducting the
       examinations.
  - 2. Participants will take turns serving as test administrator, test subject and coach.
  - 3. Teams initially will practice under lighted room conditions.
    - a. Check pupil size under normal room light.
    - b. Check reaction to light and pupil size using a pen light in a lighted room.

Total Lesson Time: Approximately 60 Minutes

Point out "Practice Sessions" wall chart.

Briefly review the objectives, content and activities of this session.

Make team assignments.

Emphasize that students can help each other learn by pointing out errors of omission or commission.

<u>Clarification</u>: students will shine a pen light directly into the subject's eye. <u>Demonstrate</u> this, using a student subject.

Aides	Lesson Plan	Instructor Notes
	4. Teams subsequently will practice under darkened room conditions.	
	a. Check pupil size in near total darkness.	
	b. Check reaction to light and pupil size under direct pen light.	
	5. Students will record their estimations using Eye Examinations Data Sheet.	Point out the copies of the Eye Examination Data Sheet in the Student's Manual.
		Solicit students' questions concerning procedures for this practice session.
	B. Room Light Examinations	
20 Minutes	Pupil size estimation, under room light.	Monitor teams and coach students as necessary and appropriate.
	2. Pupil reaction and size estimation, under direct pen light.	When the first student completes the two estimations, have the team members exchange roles. Continue this process.
		Sequence of roles should be as follows:  1. Test administrator 2. Test subject 3. Coach 4. Test administrator (continue cycle)
		Terminate this segment after 20 minutes, or after each student has twice served as a test administrator (whichever comes first).

Aides	Lesson Plan	Instructor Notes
25 Minutes	C. Dark Room Examinations	Allow students approximately 90 seconds for their eyes to adapt to the darkened conditions.
	Pupil size estimation, under near total darkness.	Monitor teams and coach students as necessary and appropriate.
	2. Pupil reaction and size estimation, under direct pen light.	When the first student completes the two checks, have the team members exchange roles. Continue this process.
5 Minutes	D. Session Wrap Up	Sequence of roles should be as follows:  1. Test administrator 2. Test subject 3. Coach 4. Test administrator (continue cycle)  Terminate this segment after 25 minutes, or after each student has twice served as a test administrator (whichever comes first).  Offer appropriate comments and observations about the students' performance.  Solicit students' comments concerning the practice session.

## **Session XI**

Practice: Eye Examinations



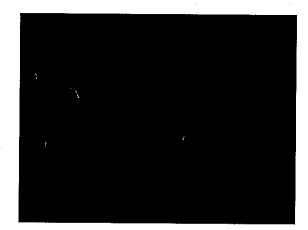
## Practice: Eye Examinations

Upon successfully completing this session, the participant will be able to:

- Conduct examinations of pupil size and reaction to light, under both lighted room and darkened room conditions
- · Articulate the eye examination procedures
- Document the results of the examinations of pupil size and reaction to light

Drug Evoluttion & Classification Training

Xi-O



One Hour and Forty-Five Minutes

SESSION XII

ALCOHOL WORKSHOP

#### SESSION XII ALCOHOL WORKSHOP

Upon successfully completing this session, the participant will be able to:

- o Correctly administer the preliminary clinical examinations and psychophysical tests used in the drug evaluation procedure.
- Observe and record the suspect's performance on the preliminary clinical examinations and psychophysical tests.
- o Determine the level of impairment based on the results of the suspect's preliminary clinical examinations and psychophysical tests.

Content Segments			<u>Learning Activities</u>		
A.	Procedures	O	Instructor Led Presentations		
В.	Hands-On Practice	О	Student Led Practice		
C.	Session Wrap Up	0	Instructor Led Discussion		

Aides	Lesson Plan	Instructor Notes
	ALCOHOL WORKSHOP	Total Lesson Time: Approximately 105 Minutes Session title on wall chart.
XII-0 (Session Objectives)		Briefly review the objectives, content and activities of this session.
10 Minutes	A. Procedures	
XII-1 (Examinations and tests)	<ol> <li>Students will work in three or four member teams during this session.</li> <li>Each team will administer a battery of tests to each volunteer.</li> <li>a. The preliminary clinical examinations and psychophysical tests include:         <ul> <li>o Pupil Size (Room Light)</li> <li>o Horizontal Gaze Nystagmus</li> <li>o Vertical Gaze Nystagmus</li> <li>o Lack of Convergence</li> <li>o Romberg</li> <li>o Walk and Turn</li> <li>o One Leg Stand (both legs)</li> <li>o Finger to Nose</li> <li>o Pulse</li> </ul> </li> </ol>	Make team assignments.  Point out that for the Drug Evaluation and Classification Examination, it is helpful to estimate angle of onset for HGN, and to relate it to BAC.
; ;	b. Results/observations of all tests will be recorded on the Standard Drug Evaluation Report Form.	Point out that copies of the report form are in the Student's Manual. Each team will need one report form for each volunteer.

Aides		Lesson Plan	Instructor Notes
	3.	For each volunteer, team members should perform the following duties:	
		a. One team member will administer the tests to the volunteer.	
		b. One team member will record the results on the report form.	
	7/200	c. The other team member(s) will assist the test administrator in observing the volunteer's performance on the tests.	Emphasize that team members will take turns performing the various duties, as they deal with the different volunteers.
	4.	Some volunteers will have BACs above 0.10, others will have lower BACs.	
	5.	The following safety precautions will be strictly enforced:	·
		a. No weapons will be present.	
		b. Volunteers will not be left unattended at any time.	Solicit students' questions concerning the procedures for the Alcohol Workshop.
	В.	Hands On Practice	
75 Minutes	1.	Test administration	Monitor teams as they test the volunteers.
70 minutos	2.	Test recording	Make sure that each student takes at least one turn as a test administrator.
			Coach students, as necessary, to improve their performance as test administrators.
	'	•	

Aides	Lesson Plan	Instructor Notes
		Terminate the hands on practice after 75 minutes, or after each team has tested 5 volunteers (whichever occurs first).
	C. Session Wrap Up	
20 Minutes		
	1. Feedback of teams' assessments	Record teams' assessments of each volunteer's probable BAC status on the chalkboard or flip chart (see next page for a sample chalkboard array). If a chalkboard or flipchart is not available, an overhead has been made.
		Ask each team <u>briefly</u> to describe the evidence that led the members to their conclusions about a particular volunteer's BAC.
	2. Feedback of volunteers' BACs.	Record each volunteer's actual BAC on the chalkboard array.
3 22 3	3. Discussion	Make appropriate comments concerning teams' assessment of the volunteers' BACs. These comments should take into account such factors as absorption and elimination rates, differences in tolerance to alcohol, volunteers' medical conditions, etc.  Solicit students' comments or questions concerning the alcohol workshop.

# SAMPLE CHALKBOARD ARRAY FOR RECORDING TEAMS' ASSESSMENTS.

## TEAMS' ESTIMATES OF BAC

Volunteer	0.05 or less	.0607	.0809	.10- .11	.12- .13	.14- .15	.16 or more	Actual BAC
								,
				ı				
							·	

(TABLE ENTRIES REPRESENT TEAMS' "VOTES")

## **Session XII**

## Alcohol Workshop



## **Alcohol Workshop**

Upon successfully completing this session, the participant will be able to:

- Correctly administer the preliminary clinical examinations psychophysical tests used in the Drug Evaluation procedure
- Observe and record the suspect's performance on the preliminary clinical examinations and psychophysical tests
- Determine the level of impairment based on the results of the suspect's preliminary clinical examinations and psychophysical tests

Drug Evaluation & Classification Training

XII-O

#### **Examinations And Tests**

- · Pupil Size (Room Light)
- · Horizontal Gaze Nystagmus
- · Vertical Gaze Nystagmus
- · Lack of Convergence
- Romberg
- · Walk and Turn
- · One Leg Stand (Both Legs)
- · Finger to Nose
- Pulse

Drug Evaluation & Classification Training

XII-



## SESSION XIII

PHYSICIAN'S DESK REFERENCE (PDR) AND OTHER REFERENCE SOURCES

## SESSION XIII PHYSICIAN'S DESK REFERENCE (PDR) AND OTHER REFERENCE SOURCES

Upon successfully completing this session, the participant will be able to:

- o Explain how the various sections of the PDR can provide information that will:
  - * aid in the drug influence evaluation;
  - * aid in courtroom testimony.
- o Use the PDR; that is, in a practical exercise, when presented with color photographs of typical prescription drugs encountered in law enforcement contacts, the student will correctly identify and classify those drugs, and list the signs and symptoms that can be caused by them and observed and documented during a drug influence examination.

### Content Segments

- A. Physician's Desk Reference as a resource
- B. Practical Exercise
- C. Other Resource Material

### Learning Activities

- o Instructor Led Presentations
- o Small Group Exercise



XIII-0 (Session Objectives) PHYSICIAN'S DESK REFERENCE (PDR)

10 Minutes

- A. Physician's Desk Reference as a Resource
  - 1. PDR is published annually.
    - a. Many versions are published:
      - o PDR for prescription drugs
      - o PDR for nonprescription drugs
      - o PDR for ophthalmology
    - b. PDR supplements are published periodically as new products are introduced during the year.
    - c. Function of the publisher is compilation, organization and distribution of information.
    - d. Product descriptions are prepared by the manufacturer, and edited and approved by their respective medical directors.

Total Lesson Time: Approximately 30 Minutes

Briefly review the content, objectives and activities of this session.

Instructors Note: Due to the unique nature of this session, instructors teaching this session should strive to develop innovative and interactive creative learning activities.

Point out that the PDR has been admitted as a "learned treatise" in previous court cases.

Point out that we will use the PDR for prescription drugs.

There are other PDR publications in addition to these.

Exhibit copy of a PDR.

00473		•	
Aides	Lesson	ı Plan	Instr
XIII-1	2. Sections of a I	PDR.	Point out th
Allt-1	a. Manufacti (section 1)	urers Index	List of manu phone numb vided prescr
		ame Index and ued Products	Alphabetica available an discontinued
			Note: New e will have a r 2 and 4.
	c. Product Ca (section 3).	ategory Index	Products list appropriate
		nd Chemical ex (section 4).	Products list and chemica according to ingredient(s)
	e. Product Ide Section (se	entification ction 5).	Point out the contains actured reproduction
	f. Product In	formation Section	Point out the

#### **Instructor Notes**

Point out that the sections are color coded for easy use.

List of manufacturers (with phone numbers) who have provided prescribing information.

Alphabetical listing of products available and a listing of discontinued products.

Note: New editions PDR 1996 will have a merging of Sections 2 and 4.

Products listed according to appropriate category.

Products listed under generic and chemical name headings according to the principal ingredient(s).

Point out that this section contains actual size, full color reproductions.

Point out that this section describes composition, action, uses, administration, dosage, contraindications, precautions, side effects, the form in which supplied and other information concerning use. Also includes common names, generic compositions or chemical names.

Diagnostic product descriptions.

List of centers and emergency telephone numbers.

g. Diagnostic Product Information (section 7)

(section 6).

h. Poison Control Centers

Aides	Lesson Plan	Instructor Notes	
	i. Guide to Management of Drug Overdose.	Information concerning drug overdosage.	
	3. Use of PDR in Drug Evaluation and Classification	This information is contained in the product identification section.	
	a. To identify prescription drugs.		
	b. To identify the effects of prescription drugs for comparison with observed effects.	This information is contained in the product information section.	
	4. How to use the PDR.		
	a. Identification of an unknown product.	Demonstrate how to identify a tablet, capsule, etc. using the product identification section.	
VIII 2	b. Identification of drug pharmacology.	Demonstrate how to use the product information section.	
AIII-2		Example: Nembutal sodium capsules (pentobarbital sodium capsules)	
	5. Location and acquisition of agency's PDR(s).	Point out that PDRs can be obtained from physicians, hospitals, etc. It is not essential to have the current version for typical enforcement uses.	
	B. Practical Exercise	Assign students to small groups and provide color slides	
15 Minutes	1. Small group exercise	or photographs of typical prescription drugs encountered	
	2. Group reports	during enforcement contacts.	
·		Have the group identify the drugs and describe typical "actions" or symptoms that can be observed and documented during a Drug Evaluation.	
	1	Each group must have a PDR.	





XIII-3A and 3B

- C. Other Resources
  - 1. National Highway Traffic Safety Administration, Traffic Law Enforcement Division
  - 2. State Drug Evaluation and Classification Program Coordinator.
  - 3. "The DRE" Newsletter
  - 4. Traffic Law Center
  - 5. Local Poison Control Center
  - 6. Medical Dictionaries
  - 7. The Pill Book, The Drug Identification Bible, and other consumer's guides to drugs
  - 8. Other texts
  - 9. Newspaper and magazine articles on drugs and drug impaired driving, including counter-culture magazines such as "High Times".
  - Software programs such as Pharmacists, Body Works, Mosbey's Medical Dictionary and other programs are available on disks and CDs.
  - 11. Various resources are available through Online services and the Internet.

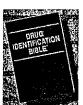
Point out that names and addresses of the State DEC Coordinators are included in the Student's Manual.

Published by the Prosecutor's Office, Phoenix, Arizona.

Instructor: Discuss some other useful texts known to you.

## **Session XIII**

Physician's Desk Reference (PDR) and Other Reference Sources



#### Physician's Desk Reference (PDR) and Other Reference Sources

Upon successfully completing this session, the participant will be able to:

- Explain how the various sections of the PDR can provide information that will:
  - aid in the drug influence evaluation
  - aid in courtroom testimony
- Use the PDR in a practical exercise, when presented with color photographs of typical prescriptions drugs encountered in law enforcement contacts. The student will correctly identify and classify those drugs and list the signs and symptoms that can be caused by them and observed and documented during a drug influence examination

Drug Evaluation & Classification Training

XIII-C

#### Sections of a Physician's Desk Reference

- · Manufacturers' index
- Product name index and discontinued products
- Product category index
- Generic and chemical name index
- Product identification
  section
- Product information section
- Diagnostic product information
- · Poison control centers
- Guide to management of drug overdose

Drug Evaluation & Classification Training

XIII-

## Product Information Section Example

Nembutal sodium capsules (pentobarbital sodium capsules)

- Description
- Clinical pharmacology
- Indications and usage
- Warnings
- Precautions
- Dosage and administration
- Drug abuse and dependence
- How supplied

Drug Evaluation & Classification Training

XIA-

## **Continuing Information Sources**

- National Highway Traffic Safety Administration, Traffic Law Enforcement Division
- · State DEC Program Coordinator
- "DRE" Newsletter Phoenix City Prosecutor's Office 455 North 5th Street Suite 400 Phoenix, AZ 85004

Drug Evaluation & Classification Training

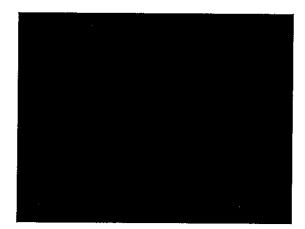
XIII-3A

# Continuing Information Sources (continued)

- · Local poison control center
- · Medical dictionary
- Pill Book
- Other text books (physiology, pharmacology, toxicology, etc.)
- · Newspaper and magazine articles

Drug Evaluation & Classification Training

BE-fitX



One Hour and Forty-Five Minutes

SESSION XIV
HALLUCINOGENS

### SESSION XIV HALLUCINOGENS

Upon successfully completing this session, the participant will be able to:

- o Explain a brief history of the Hallucinogen category of drugs.
- o Identify common drug names and terms associated with this category.
- o Identify common methods of administration for this category.
- o Explain the symptoms, observable signs and other effects associated with this category.
- o Explain the typical time parameters, i.e., onset and duration of effects, associated with this category.
- o State the clues that are likely to emerge when the Drug Evaluation and Classification Process is conducted for a person under the influence of this category of drugs.
- o Correctly answer the "topics for study" questions at the end of this Section.

Content Segments		Learning Activities	
A.	Overview of the Category	O	Instructor Led Presentations
В.	Possible Effects	0	Review of Drug Evaluation and Classification Exemplars
C.	Onset and Duration of Effects	0	Reading Assignments
D	Overdose Signs and Symptoms	0	Video Presentations (If Available)
E.	Expected Results of the Evaluation	O	Slide Presentations

Aides	Lesson Plan	Instructor Notes		
* 10 4	HALLUCINOGENS	Total Lesson Time: Approximately 105 Minutes		
XIV-0A&B (Session Objectives)	A. Overview of the Category	Session title on wall chart.		
20 Minutes				
	<ol> <li>Hallucinogens are drugs that affect a person's perceptions, sensations, thinking, self awareness and emotions.</li> <li>The word "Hallucinogen" means something that causes hallucinations.</li> <li>An hallucination is a sensory experience of something that does not exist outside the mind.</li> </ol>	Briefly review the objectives, content and activities of this session.  Definition from The Random House College Dictionary (Revised Edition, 1980).		
	<ul> <li>o Seeing, hearing, smelling, tasting or feeling something that isn't really there.</li> <li>o Having distorted sensory perceptions, so that things look, sound, smell, etc. differently from the way they really are.</li> </ul>			



- c. Hallucinogenic drugs usually produce what are called pseudo-hallucinations: i.e., the user typically is aware that what he or she is seeing, hearing, smelling, etc. isn't real, but is a product of the drug.
- d. One common type of hallucination produced by these drugs is called <u>Synesthesia</u>, which means a transposing of sensory modes.
  - o Sounds for example, may be transposed into sights.
  - o Sights may be transposed into odors.
- e. The illusions and distorted perceptions produced by hallucinogenic drugs may be very alarming, even terrifying.
  - o They may produce panic and uncontrolled excitement.
  - o The user may be unable to cope with the terror, and may attempt to flee wildly.
  - A user who is emotionally or mentally unstable may become psychotic in response to this frightening experience.

But emphasize that the fact that the user knows the hallucinations aren't real doesn't make those hallucinations any less dangerous if they occur while driving.

Note: Synesthesia may occur naturally in an insignificant percentage of the population.

Examples: The user may "see" a flash of color, or some other sight, when the telephone rings.

The user may "smell" a particular fragrance when he or she looks at something painted red.

<u>Point out</u> that the expression "bad trip" refers principally to these panic filled reactions to Hallucinogens.



XIV-2 ("Flashback")

- f. A terrifying "bad trip" sometimes may be reexperienced as a <u>flashback</u>.
  - o In simple terms, a flashback is a vivid recollection of a portion of an hallucinogenic experience.
  - o A flashback does <u>not</u> occur because of a residual quantity of drug in the user's body.
  - o Instead, a flashback essentially is a very intense daydream.
- g. There are three types of flashback:
  - o Emotional: Feelings of panic, fear, etc; the sensations of a "bad trip".
  - o Somatic: Altered body sensations, tremors, weakness, dizziness, crawly, tingly feelings on the skin.
  - o Perceptual: Distortions of vision, hearing, smell and/ or other senses.
    These distortions are "re-runs" of the original "trip".

But point out that subsequent use of the drug may precipitate a flashback, by causing the user to reexperience the frightening illusions of the previous "bad trip".



XIV-2A (Types of Flashback)

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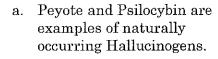
#### Lesson Plan

#### **Instructor Notes**



XIV-2B (Illusions and Delusions)

- h. Remember that hallucinogens produce <u>illusions</u>, <u>delusions</u> or both.
  - o An illusion is a false perception, i.e., a misrepresentation of what the senses are receiving.
  - o A delusion is a false belief.
- Because they often make the user appear to be insane, Hallucinogens sometimes are called psychotomimetic drugs.
- 2. Some Hallucinogens come from natural sources, while others are synthetically manufactured.



b. LSD, MDA, MDMA, DMT, STP and TMA are examples of synthetically manufactured Hallucinogens. Example of an illusion: "I see an Elephant".

Example of a delusion: "I am an Elephant".

Print "PSYCHOTOMIMETIC" on the chalkboard or flip chart.

"Psychotomimetic" means "something that mimics psychosis". A psychosis is a major mental disorder. It implies a loss of touch with reality.



XIV-3A (Hallucinogens)

Instructor, for your information: Other naturally occurring Hallucinogens include nutmeg; jimpson weed; morning glory seeds; and, bufotenine, a substance found in the glands

Note: Some regional or local Hallucinogens may be discussed in more detail.

LSD: Lysergic Acid Diethylamide

of certain toads.



Aides	Lesson Plan		Instructor Notes		
			MDA: Methylene Dioxyamphetamine		
			MDMA: Methylene DioxyMethamphetamine (also known as "Ecstasy").		
			STP: (also known as DOM) Dimethoxylamphetamine		
			TMA: Trimethoxyamphetamine		
			DMT: Dimethyltryptamine		
			Instructor, for your information: Drugs such as MDA, MDMA, STP and TMA all contain amphetamine based compounds. The are for this reason sometimes called "psychedelic amphetamines". In essence, they are high powered CNS Stimulants that cause hallucinations.		
		yote is a small, spineless etus.	·		
	a.	The active, hallucinogenic ingredient in peyote is mescaline.	If available, show 35 mm slides of the peyote cactus and of dried "buttons" or crowns from the cactus.		
	Ъ.	Peyote use by certain Indian tribes for religious rituals pre-dates Columbus' discovery of America by many centuries.	Mescaline is a chemical relative of adrenalin. Its effects on the body may be similar to those that would result from a massive rush of adrenalin.		
	c.	Peyote is used legally in religious ceremonies of the Native American Church.	Mescaline was first isolated from Peyote in 1856. It was named after the Mescalero Apaches.		

Aides	Lesson Plan	Instructor Notes	
		Persons who are not American Indians cannot be members of the Native American Church.	
	4. Psilocybin is a drug found in a number of different species of mushrooms of the genus Psilocybe.	Eighty-one species of these mushrooms have been identified as hallucinogenic.	
	a. These mushrooms also have been used in Indian religious ceremonies for thousands of years.	If available, show 35 mm slides of Psilocybin Mushrooms.	
	b. An unstable derivative of Psilocybin, called <u>Psilocin</u> , also is found in these mushrooms and also has hallucinogenic properties.	Psilocybin is chemically very similar to serotonin, a neurotransmitter that is found in the brain.  The effects of Psilocybin may be similar to what would happen if the brain were suddenly flooded with Serotonin.	
	<ol> <li>LSD is perhaps the most famous of the synthetically manufactured Hallucinogens.</li> </ol>		
	a. "LSD" is an abbreviation of Lysergic Acid Diethylamide.		
	b. It was first produced in 1938, although its halluci- nogenic properties were not discovered until 1943.		
	c. LSD was used in psychotherapy during the 1940's and early '50's.	Example: It was occasionally used in the treatment of alcoholism.	
		If available, show 35 mm slides of various forms of LSD.	

#### Lesson Plan

## **Instructor Notes**



- d. Although LSD is a synthetic drug, it was first derived from Ergot, a fungus that grows on rye and other grains.
- e. In the Middle Ages, when people accidentally ate this fungus, their resulting bizarre behavior was thought to stem from possession by the Devil.
- f. The trials and subsequent burning of "witches" in Salem, Massachusetts in 1692 probably was due to accidental Ergot consumption by those women.
- g. Ergot is still used medically to treat migraine headaches.
- 6. MDA, STP and TMA are synthetically manufactured Hallucinogens that sometimes are called "Psychedelic Amphetamines".
  - a. They are chemically related to Amphetamines and produce many effects similar to those of CNS Stimulants.
  - b. They are also chemically related to Mescaline.
  - c. MDA is an abbreviation for 3, 4-Methylenedioxy-amphetamine
  - d. Among users, MDA sometimes is referred to as the "Mellow Drug of America".

Write "LSD derived from Ergot, a fungus" on the chalkboard or flip chart.

Sandoz Laboratories markets a combination of caffeine and Ergot called Cafergot.

The small "buttons" or crowns of the Peyote Cactus are dried and eaten, or may be brewed into a beverage for

drinking.

Aides	Lesson Plan	Instructor Notes
	o Similarly, the Psilocybin Mushrooms are dried and eaten, or may be brewed into a beverage for drinking.	
	<ul> <li>b. Some Hallucinogens can also be <u>smoked</u> (example: LSD impregnated on Marijuana or tobacco cigarettes).</li> </ul>	Point out that some Hallucinogens such as LSD can be absorbed through the skin. Officers should make it a practice to wear latex gloves when handling any suspected drugs.
	c. Some users <u>inject</u> LSD.	
	d. MDA can also be <u>insufflated</u> , or "snorted".	Solicit students' comments or questions on this overview of Hallucinogens.
	B. Possible Effects	
5 Minutes		
,	1. The effects of Hallucinogens vary widely, and are affected by the user's personality, mood and expectations, and by the surroundings in which the drug is taken.	
	a. Generally, Hallucinogens intensify whatever mood the user is in at the time the drug is taken.	
	o If the user is depressed, the drug will deepen the depression.	
	o If the user is feeling pleasant, the drug will heighten that feeling.	

Aides

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		b. If the user expects that the drug will help him or her achieve new insights or an expanded consciousness, the "trip" will seem to have that effect.	
	2.	However, Hallucinogens also often uncover mental or emotional flaws that the user was unaware of possessing.	
	3.	Therefore, many users who expect a positive experience with the drug will encounter instead the panic of a "bad trip".	
	4.	The most common effect of the Hallucinogen is hallucination: the distorted perception of reality, often with a mixing of senses that makes it virtually impossible for the drug influenced user to function in the real world.	Solicit students' comments or questions on this overview of Hallucinogens.
	C.	Onset and Duration of Effects	
15 Minutes XIV-4A ("Time	1.	The time parameters associated with Hallucinogens vary from drug to drug.	
Factors of Peyote")	2.	The effects of Peyote (Mescaline) begin to be felt within approximately one-half hour after eating the cactus "buttons".	
		a. 30 minutes: nausea, possibly leading to vomiting; mild rise in blood pressure, pulse, temperature and heart rate; pupils dilate.	

Lesson Plan

**Instructor Notes** 

#### Lesson Plan

#### **Instructor Notes**

- b. One hour: sensory changes begin; visual distortions accompanied by rich colors; objects take on new forms and begin to move; shapes "come alive".
- c. <u>3-4 hours</u>: sensory changes reach their peak; synesthesia (mixing of senses) commonly occurs.
- d. 10 hours: gradual decline in effects.
- e. 12 hours: nearly total recovery from effects.
- 24 hours: approximately 87% of the Mescaline has been excreted from the body.
- 3. Psilocybin also begins to exert its effects within one-half hour.
  - a. 1-30 minutes: dizziness, light headed feeling, giddiness; the extremities (hands, feet, etc.) may feel very light or very heavy.
  - b. 30-60 minutes: vision blurs; colors become brighter, leave longer lasting after images; objects take on sharp visual definition; hearing becomes more acute.



Factors of Psilocybin")



- c. 60-90 minutes: color patterns and shapes start to develop; the surfaces of objects appear to develop waves and wave-like patterns; distance perception becomes impaired; feelings of euphoria develop.
- d. 90-100 minutes: body sensations increase, along with mental perceptions; user commonly becomes introspective.
- e. <u>120-180 minutes</u>: effects start to diminish.
- 4. LSD's effects begin to be felt within 30-45 minutes.
  - a. 30-45 minutes: blood pressure, pulse and temperature rise; pupils dilate; hair starts to stand on end (Piloerection); nausea, dizziness and headache develop.
  - b. <u>4-6 hours</u>: effects reach their peak.
  - c. 7-9 hours: effects diminish.
  - d. <u>10-12 hours</u>: user feels normal.
- 5. Onset and duration of effects of other Hallucinogens vary widely from about 2 hours to about 24 hours.



#### 5 Minutes

- D. Overdose Signs and Symptoms
  - 1. Death from overdose of LSD or Mescaline is not common.
    - a. It is unlikely that other
      Hallucinogens would
      directly result in death from
      overdoses.
    - b. However, an overdose can be extremely dangerous and indirectly result in death.
      - o The extreme panic and agitation of a "bad trip" have been known to result in suicide, or in accidental death as the user attempts to flee the hallucinations.
      - o Sometimes Hallucinogens induce a
        perception of
        invulnerability in the
        user, leading to bizarre
        and very dangerous
        behavior, and death.
  - 2. The most common danger of an overdose of Hallucinogen is an intense "bad trip", which can result in severe and sometimes permanent psychosis.
  - 3. Some evidence also suggests that prolonged use of LSD may produce organic brain damage, leading to impaired memory, reduced attention span, mental confusion and impaired ability to deal with abstract concepts.

Example: At least one LSD user was killed when he attempted to stop a train. Others have died from jumping off buildings believing they can fly.

Solicit students' comments and questions concerning time factors.

#### Aides

#### Lesson Plan

#### **Instructor Notes**

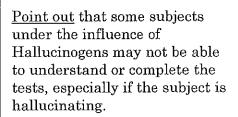


60 Minutes



XIV-5A ("SFST Evidence")

- E. Expected Results of the Evaluation
  - 1. Observable evidence of impairment.
    - a. Standardized Field Sobriety Tests.
      - o Neither Horizontal nor Vertical Gaze Nystagmus will be present.
      - o Performance on the Romberg balance test will be impaired, particularly in the subject's estimation of the passage of 30 seconds.
      - o Performance on the
        Walk and Turn, One
        Leg Stand and Finger to
        Nose tests will be
        markedly impaired due
        to the subject's severe
        visual distortion,
        impaired perception of
        distance and decreased
        muscle coordination.
    - b. General indicators
      - o dazed appearance
      - o body tremors
      - o perspiring
      - o uncoordinated movements
      - o muscle tone normal/rigid



Emphasize that DRE officers conducting evaluations on subjects under the influence of hallucinogens should be especially careful due to the bizarre and unpredictable behavior of these subjects.



XIV-5B ("General Indicators")



XIV-5C ("Eye Examinations")



XIV-5D ("Vital Signs Examinations")



XIV-6 ("Hallucinogens Symptomatology Chart")

- o difficulty with speech
- o statements/utterances suggesting hallucinations or distorted sensory perceptions
- 2. Evidence associated with the physiologic examinations.
  - a. Eye examinations
    - o Lack of Convergence will not be evident.
    - o pupils generally will be dilated.
    - o Reaction to light will usually be normal.
      Certain Psychedelic
      Amphetamines usually will slow the pupils' reaction to light
  - b. Vital signs examinations
    - o blood pressure generally will be elevated.
    - o pulse generally will be up.
    - o body temperature generally will be up.
- 3. Summary

4. Demonstrations

Aides	Lesson Plan	Instructor Notes
	a. Video tape demonstrations (if available)	Show video tape of subject(s) under the influence of Hallucinogens. Relate behavior and observations to the symptomology chart.
	b. Drug Evaluations and Classification exemplar demonstrations	Refer students to the exemplars found at the end of Section XIV of their student manuals.
		Relate the items noted on the exemplars to the symptomatology chart.
		Solicit students' questions or comments concerning expected results of the evaluation of subjects under the influence of Hallucinogens.

# **Session XIV**

## Hallucinogens



## Hallucinogens

Upon successfully completing this session, the participant will be able to:

- Explain a brief history of the Hallucinogen category of drugs
- Identify common drug names and terms associated with this category
- Identify common methods of administration for this category
- Explain the symptoms, observable signs and other effects associated with this category

Drug Evaluation & Classification Training

XIV-OA

# Hallucinogens (continued)

- Explain the typical time parameters, i.e., onset and duration of effects, associated with this category
- State the clues that are likely to emerge when the drug evaluation and classification process is conducted for a person under the influence of this category of drugs
- Correctly answer the "topics for study" questions at the end of this section

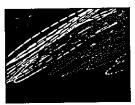
Drug Evelvation & Classification Training

NV-OB

## Synesthesia:

A transposition of senses

- "Seeing sounds"
- "Hearing colors"



Drug Evaluation & Classification Training

VII.

#### Flashback:

A vivid recollection of a hallucinogenic experience

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XIV-2

## **Types of Flashbacks**

Emotional

Most dangerous, feelings of panic, fear, etc., sensation of "bad trip"

Somatic

Altered bodily sensations, tremors, weakness, dizziness, crawly, tingly feeling on the skin

Perceptual

Distortions of vision, hearing, smell, taste and touch (associated with original "trip" least harmful, unless driving a motor vehicle)

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XIV-2A

#### Illusion:

A false perception

#### **Delusion:**

A false belief

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## Common Hallucinogens





Peyote (Mescaline)





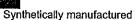
Psilocybin

· Both occur naturally

XIV-3A

## Common Hallucinogens (continued)





- LSD (Lysergic Acid Diethylamide)
- MDA (3,4-Methylenedioxyamphetamine)
- STP (or DOM) (2 Methyl-2,5 Dimethoxylamphetamine)

TMA (3,4,5-Trimethoxyamphetamine)





#### **Time Factors of Peyote**

30 minutes: Onset

Nausea; elevated blood pressure, pulse and temperature; ditated

60 minutes: Development of hallucinogenic effects

Visual distortions, rich colors, changing forms and 3-4 hours: Peak effects

Synesthesia

10 hours: Gradual decline of effects

12 hours: Nearly total recovery

24 hours: Elimination nearly completed

#### Time Factors of Psilocybin

- 1-30 minutes: Onset Dizziness; giddiness; lightness or heaviness of extremities
- 30-60 minutes: Beginning of sensory effects Blurred vision, sharpness of color, increased aculty
- 60-90 minutes: Sensory effects intensify Patterns and shapes develop and move; distance perception is impaired; euphoria develops
- 90-100 minutes: Peak effects Subject becomes introspective
- 120-180 minutes: Effects begin to diminish

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#### Time Factors of LSD



30-45 minutes: Onset

· 4-6 hours: Peak effects

· 7-9 hours: Effects diminish

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10-12 hours: Subject feels normal



XIV-4C

## **Evaluation of Suspects Under** the Influence of Hallucinogens

#### SFST Evidence:

- · HGN None
- Impaired performance will be evident on Walk and Turn and One Leg Stand
- Impaired performance will be evident on Romberg and Finger To Nose

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XIV-SA

## **Evaluation of Suspects Under** the Influence of Hallucinogens

General Indicators:

- · Dazed appearance
- · Body tremors
- · Perspiring
- · Uncoordinated movements
- · Muscle tone normal/rigid .
- · Difficulty with speech
- · Statements suggesting hallucinations

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XIV-58

## **Evaluation of Suspects Under the Influence of Hallucinogens**

#### Eye Examinations:

- · Lack of Convergence none
- · Pupils will be dilated (Mydriasis)
- · Reaction to light will be normal*

*Certain psychedelic amphetamines cause slowing

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XIV-50

## **Evaluation of Suspects Under** the Influence of Hallucinogens

#### Vital Signs:

- · Blood pressure will be up
- · Pulse will be up
- · Body temperature will be up

Drug Evaluation & Classification Training

XIV-50

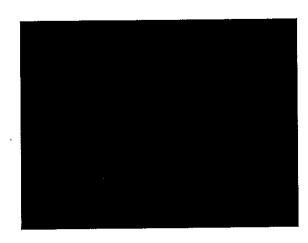
## Hallucinogen Symptomatology Chart

HGN	None
VGN	None
Lack of Convergence	None
Pupil Size	Dilated (mydriasis)
Reaction to Light	Normal*
Pulse Rate	Uр
Blood Pressure	Up
Temperature	Up
Muscle Tone	Possibly rigid

* Certain psychedelic amphetamines cause slowing

Drug Evaluation & Classification Trainin

XIV-6



	Drug Influence Evaluation							
Γ	Evaluator PAGE	T	DRE No / L	0	Rolling Log	005		
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		last sleep? How long?	Are you sick o	r injured?	<b>⊗</b> Yes □]	No	Are you diabetic or	epileptic?
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	Opinion of Evaluator:	☑ Rule Out □ Medical	□ Alcohol □ Depressant	<ul><li>Stimular</li><li>Hallucin</li></ul>		otic Analgesic	□ Innaiant □ Cannabis	

00000				
	DRUG INFLUENCE EVALUATION	N Page <u> 2</u> of <u>2</u>		
LOG NO.	DRE: Sgt. Tom Page	ARRESTEE: Rebecca S. Hoeckle		
5. INITIAL OB	1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.			
1. LOCATIO	ON: Examination of Rebecca S. Hoeckle took place in	the Central Testing Unit. Nassau County PD		
2. WITNESS	S: Arresting Officer - Officer R. Buoneto, Nassau Cou	unty PD and ADA Edward Bracken, Suffolk County		
3. BREATH	I TEST: Officer Buoneto administered an Intoxilyzer	breath test to Hoeckle, the result was 0.00%		
4. NOTIFIC	CATION / INTERVIEW of ARRESTING OFFICE	R: Writer was notified by Officer Buoneto and		
requested t	to conduct a DRE evaluation. Officer Buoneto states	d the subject had been operating her 1994 Chevrolet		
(NY127 N	VCQ) and was stopped in the S/B traffic lane of Island	d Drive, at the intersection with Hauppage Drive for a		
green light.	t. Upon approaching the vehicle, subject turned to hir	n, pointed to the traffic light and said "God is light		
and the ligh	ht is of God"			
5. INITIAL	OBSERVATIONS: Subject was seated next to the I	ntoxilyzer table and staring fixedly ahead. She		
slowely tu	urned towards me and asked "are you of God?" I rep	lied that my name was Tom, and that I would like to		
examine h	her. She nodded and said," God sent you therefore yo	u must be good." Her speech was rapid and she		
stuttered s	stuttered slightly.			
6. MEDICA	L PROBLEMS: Subject indicated she was experience	ing a mildly upset stomach. At the end of the DRE		
examinati	ion, Dr. J. P. Mooney was summoned to examine her	. (		
7. PSYCHO	PHYSICAL TESTS: Subject was unable to stand w	ithout assistance, and it was necessary to terminate the		
Romberg	Romberg Balance, Walk and Turn, and the One Leg Stand Tests virtually immediately for the subjects own safety.			
Finger to 1	Nose was conducted while the subject was in the seat	ed position she missed tip of her nose on each		
attempt.				
8. CLINICA	AL INDICATORS: Subject's pulse, blood pressure a	and temperature were above the normal range, and her		
pupils we	ere dilated.			
9. SIGNS of	INGESTION: Subjects breath had a sour and ranci	d odor.		
10. STATEM	MENTS: Subject stated that she was fasting for religion	ous reasons, and that her religion forbids the ingestion		
of alcoho	olic beverages. She also stated that her medium doesn	't allow her to use drugs. She further indicated that		
her mediu	am is her religious leader a man "whose body is of fir	e and air, and whose spirit is of light.		
which is o	which is of God" She indicated she had just attended a service conducted by the medium.			
11. OPINIO	ON of EVALUATOR: In my opinion Rebecca S. Hoo	eckle is under the influence of a Hallucinogen and		
unable to	unable to operate a vehicle safely.			
12. TOXICO	12. TOXICOLOGICAL SAMPLE: Subject agreed to provided a blood sample.			
13. MISCELLANEOUS:				
		· · ·		

000501	Drug Influen		on		
Evaluator HAVERSAT, A	DRE No	Rolling Log No.			
Recorder/Witness	Crash: □ Non □ Fatal □ Inju	ry O Pro			
estee's Name (Last, First, MI)	DOB Sex		esting Officer (Name		
JARBURTON, CINDY T Date Examined/Time/Location	07/18/70 P	u	JACKSON	1 F #6	3/0 //PPD
1	·	Inchusmont #	· I	Chemical Test □ Refused	□ Urine 🎓 Blood
APRIL Z5 1996 - 2300 - Miranda Warning Given: Yes O No Wha	ZND DISTRICT	134		How much?	
Miranda Warning Given: Yes □ No Wha By: What was a second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second of the second	t have you eaten today? W	, ,	t been drinking?	How much?	Time of last drink?
JACKSON, F S	PAGHETTI	LUNCH /L	No THINE	Are you diabetic or	enilentic?
By: JACKSON, F  Time now? When did you last sleep? How long?  7 PM YESTERDAY 6 HK  Do you take insulin?	I DENT THE	Nk so Itey	110	□ Yes •No	
Do you take insulin? ☐ Yes ♠No	Do you have any physical	defects? □ Yes •	No	Are you under the o a doctor or dentist?	
Are you taking any medication or drugs? □ Yes •••••••••••••••••••••••••••••••••••	o Attitude LOOPERATIVE SE	AT FEARFAL	I DISTRATES	Coordination	TAGGERING
	[ D	JOHNAL	Z 2001117-1 C. y	Face PERSPI	
Speech RAMBILIAN	Eyes:   Reddened Cor		Blindness:		Tracking:
13/14/10/04/15/0	Mormal □ Bloodshot	□ Watery	None □ L Eye	OR Eye	<b>●</b> Equal ○ Unequal
Corrective Lens: None Pupi	l Size: ÆEqual		Able to follow stim		Eyelids:
□ Glasses □ Contacts, if so □ Hard □ Soft □ Ur	nequal (explain)				Normal □ Droopy
Pulse & Time HGN	Left Eye Right Eye	Vertical Nystagmus?	'□Yes 👂 No	One Leg Star	nd LEG SHAFING
1. 1/2 / 23/0 Lack of Smooth Pursuit	100 10	Convergence	Left Eye	9	) <u>(</u> )
2. //6 / 2325 Max. Deviation	NO NO	Right Eye	Leit Lye	Ü	6
3. //6 / 1340 Angle of Onset	NONE LEW			<b>Q</b>	
Romberg Balance Walk and Turn		Cannot Ke	eep Balance		
Approx. Approx.		Starts Too	1 st Nine 2 nd	Nine I R	, ways While Balancing
2503503		Stops Wall	king VV V	/	ses Arms to Balance
	,	Misses Hee	1/VV V	7 <u>777</u>	
	فالقلامليلين	Steps Off I Raises Arn	<del>  _   '</del>		opping
	n s m s :	M Actual Ste	CONSTRAI CO.	S D P	uts Foot Down
Internal Clock Describe Turn	LOST BALANCE	Cannot Do Test (I	Explain)	Type of Foot	wear
// Potimated At 30 Sec	-	1//		LOP	AFER'S
SIGNISE	D NEARLY FEL		Darkness Dire	ect Nasal Area	,
O Right \( \triangle \text{Left} \)  If CAM FEEL MY FACE  Draw lines from spot touched	Left Eye			<u> </u>	LEAR
Draw lines from spot touched	3.1	6,0	7.5 5.	5 Oral Cavity	
A A	Right Ey	6.0	7.5 5	$5 \mid CC$	EAK
N. 040.	Hippus		nd Dilation React	ion To Light	
	111	A110 1162	•	NORM	AL
		NGHT ARM		LEFT ARM	/
2				~	
X	/3				
	(3)		- Min	E .	
(5)	8		WO 11247	<b>秋わり</b>	\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\
	<u> </u>				
Riond Praceura 150 / 107 Ter	np 99,8 °				7
Musels Breast C Mormal C Floorid & Rigid	up		D1 ( 0°F )	1 To 74 T	1
Commonto		Attach	n Photos Of Fres	h Puncture Mar.	KS
at Medicine or Drug Have You Been Using?	How Much? Time of	Use? Where	Were The Drugs Us	ed? (Location)	
MOTHINE	NO A	NSWER	NO A	WEWER _	
Date/Time of Arrest Ti	me DRE Notified 2240	Eval S	Start Time	Time Comple	eted 45
7 1 1 7 1 1 1	ID Noy		wed By:	1.0	1~
GRI TTIVOREN	□ Alcohol □ Stim		n heland	□ Inhalant	
Opinion of Evaluator:			cotic Analgesic	□ Cannabis	

DRUG INFLUENCE EVALUATION Page 2_of 2
LOG NO. DRE: Sgt. Art Haversat ARRESTEE: Cindy T. Warburton
1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.
1. LOCATION: Examination of Cindy T. Warburton, took place in the DRE room, 2nd District Hdqtrs. Capitol PD
2. WITNESS: Arresting Officer - F. Jackson # 6310 Capitol PD and R.C. Studdard, IACP/TAP Representative
3. BREATH TEST: Writer observed Officer Jackson administer GCI breath test to Warburton, the result was 0.00%
4. NOTIFICATION / INTERVIEW of ARRESTING OFFICER: Writer was serving as on-duty DRE for 2nd
District when informed by dispatch that Officer Jackson was enroute with a subject and was requesting a drug
evaluation. Upon arrival Officer Jackson stated the subject had been arrested driving N/B along the gravel shoulder of
the S/B lane Higgenbotham Ave. Jackson further stated the subject pointed to the police baton and shouted "My God
there's a terrible big snake hanging from your belt. Subsequently, she shouted that the blue and red emergency lights
on his of cruiser were bleeding into her eyes and skin.
5. INITIAL OBSERVATIONS: Writer observed subject seated next to the GCI. Subject was very frightened and
disoriented. She pointed to the clock on the wall and shouted "Keep that off me, keep it away!" At the time the clock
indicated 2245 hours. Minutes later in response to my question "What time is it now?" Subject stated it was "7
o'clock"
6. MEDICAL PROBLEMS: None observed or stated.
7. PSYCHOPHYSICAL TESTS: Romberg Balance: Subject swayed approximately 3" side to side and estimated 10
seconds as 30 seconds. Walk and Turn: Subject started walking to soon, lost her balance during the instructions,
missed heel to toe, stopped walking, stepped off the line, raised her arms, staggered while turning, and only took
(8) steps on the way back. One Leg Stand: Subject swayed, raised arms, hopped, and put her foot down. Finger to
Nose: Subject missed tip of her nose on each attempt. She opened her eyes and shouted "I can't feel my face!
My face is missing!"
8. CLINICAL INDICATORS: Subject had dilated pupils. Blood pressure, pulse, and temperature were above the
normal range.
9. SIGNS of INGESTION: None were evident
10. STATEMENTS: Subject stated that she felt hot, and denied any drug use.
11. OPINION of EVALUATOR: In my opinion Cindy T. Warburton is under the influence of a Hallucinogen, and
unable to operate a vehicle safely
12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a blood sample.
13. MISCELLANEOUS: At the time of the evaluation, subject was wearing a T-shirt bearing the words "Legalize
Acid"

	DRUG INFLUENCE EVALUATION	Page_2_of_2		
LOG NO.	DRE: Sgt. Bob Hohn	ARRESTEE: Lew B. Buchanan		
1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.				
1. LOCATION: Exami	nation of Lew B. Buchanan, took place in t	ne DRE room, Central Testing Unit Nassau County		
2. WITNESS: Arresting	Officer - D. Gregory, Nassau County PD			
3. BREATH TEST: W	riter observed Officer Gregory administer (	GCI breath test to Buchanan, the result was 0.05%.		
Subject later admitte	ed to consuming "a couple of beers"			
4. NOTIFICATION /	INTERVIEW of ARRESTING OFFICER	: Writer was summoned to Central Testing to		
conduct a DRE evalu	ation. Officer Gregory stated he had observe	ed subject driving at 10/55 zone on the		
Cross Island Parkway	y, drifting from lane to lane. Subject perform	ned poorly on the SFSTs.		
5. INITIAL OBSERVA	ATIONS: Writer observed subject in the bre	ath testing room, he was swaying slightly as he stood,		
and appeared dazed a	nd disoriented. He responded slowly to my	greeting, but was generally cooperative and		
responsive to question	s. In response to my question "What time is	it now?" Subject stated it was "about 10 o'clock"		
6. MEDICAL PROBL	EMS: Subject indicated some nausea			
7. PSYCHOPHYSICA	L TESTS: Romberg Balance: Subject swa	yed approximately 3" in a circular motion and		
estimated 35 seconds	s as 30 seconds. Walk and Turn and One Le	g Stand: Subject was unable to perform tests. Tests		
were terminated for	subject's safety. Finger to Nose: Subject mi	ssed tip of his nose on each attempt.		
8. CLINICAL INDICA	ATORS: Subject exhibited lack of smooth p	ursuit and dilated pupils. Blood pressure, pulse, and		
temperature were ab	ove the normal range.			
9. SIGNS of INGESTI	ON: None were evident			
10. STATEMENTS: S	ubject stated that he did not used any drugs.	·		
11. OPINION of EVAL	LUATOR: In my opinion Lew B. Buchana	n is under the influence of Alcohol and a		
Hallucinogen, and u	Hallucinogen, and unable to operate a vehicle safely			
12. TOXICOLOGICA	L SAMPLE: Subject agreed to provide a bl	ood sample.		
13. MISCELLANEOU	S:			
	CONTRACTOR OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF THE PROPERTY OF			

Forty-Five Minutes

## SESSION XV

PRACTICE: TEST INTERPRETATION

#### SESSION XV PRACTICE: TEST INTERPRETATION

Upon successfully completing this session, the participant will be able to:

- o Analyze the results of a complete Drug Evaluation and Classification Examination and identify the category or categories of drugs affecting the individual examined.
- o Articulate the basis for the drug category identification.

## Content Segments

- A. Interpretation Demonstrations
- B. Interpretation Practice

### Learning Activities

- o Instructor Led Demonstrations
- o Small Group Practice
- o Participant Led Presentations

Aides	Lesson Plan	Instructor Notes
, 1	PRACTICE: TEST INTERPRETATION	Total Lesson Time: Approximately 45 Minutes
Mo e Color		Point out the "Test Interpretation" wall chart.
XV-0 (Session		Briefly review the objectives, content and activities of this session.
Objectives)	A. Interpretation Demonstrations	
20 Minutes		
	1. Case #1: "Subject Adams"	Direct students to review to the "Subject Adams" exemplar in Section XV of their manuals.
	a. Preliminary examination.	Review the results of the Preliminary Examination of Subject Adams.
		Ask students: "What category or categories of drugs would produce preliminary examination results consistent with this exemplar?" Probe to draw out the bases for students' responses.
	b. Eye examinations.	Review the results of the Eye Examinations of Subject Adams.
		Ask students to discuss the category or categories of drugs that would cause these eye examination results.
	c. Psychophysical tests.	Review the results of the Psychophysical Tests of Subject Adams.

Aides	Lesson Plan	Instructor Notes
		Ask students to discuss the category or categories of drugs that would produce these psychophysical test results.
	d. Vital Signs examinations.	Review the results of the Vital Signs Examinations of Subject Adams.
		Ask students to discuss the category or categories of drugs that would produce these results.
	e. Dark room examinations.	Review the results of the Dark Room Examinations of Subject Adams.
		Ask students to discuss the category or categories of drugs that would produce these results.
	f. Other evidence and additional observations.	Review the results of the examinations for injection sites and muscle rigidity, and of the final interview of Subject Adams.
	g. Narrative report.	Briefly review the narrative report on the reverse side of the "Adams" exemplar. Point out that the DRE's opinion is missing from this sample report.
		Ask students to comment on the category or categories of drugs that would be consistent with all of the evidence on this exemplar.
	h. Opinions of evaluator.	Point out that the evidence indicates that Subject Adams is under the influence of CNS Depressants.

Aides	Lesson Plan	Instructor Notes	
		Solicit students' questions concerning this demonstration.	
	2. Case #2: "Subject Baker".	Direct students to review to the "Subject Baker" exemplar.	
·	a. Preliminary examination.	Review the results of the Preliminary Examination of Subject Baker.	
		Ask students: "What category or categories of drugs would produce preliminary examination results consistent with this exemplar?" Probe to draw out the bases for students' responses.	
	b. Eye examinations.	Review the results of the Eye Examinations of Subject Baker.	
		Ask students to discuss the category or categories of drugs that would cause these eye examination results.	
	c. Psychophysical tests.	Review the results of the Psychophysical Tests of Subject Baker.	
		Ask students to discuss the category or categories of drugs that would produce these psychophysical test results.	
	d. Vital signs examinations.	Review the results of the Vital Signs Examinations of Subject Baker.	
		Ask students to discuss the category or categories of drugs that would produce these results.	

Aides	Lesson Plan	Review the results of the Dark Room Examinations of Subject Baker.			
	e. Dark room examinations.				
		Ask students to discuss the category or categories of drugs that would produce these results.			
	f. Other evidence and additional observations	Review the results of the examinations for injection sites and muscle rigidity, and of the final interview of Subject Baker.			
	g. Narrative report.	Briefly review the narrative report on the reverse side of the "Baker" exemplar. Point out that the DRE's opinion is missing from this sample report.			
		Ask students to comment on the category or categories of drugs that would be consistent with all of the evidence on this exemplar.			
	h. Opinions of evaluator.	Point out that the evidence indicates that Subject Baker is under the influence of CNS Stimulants.			
		Solicit students' questions concerning this demonstration.			
	B. Interpretation Practice				
25 Minutes	1. Team practice.	Assign students to work in teams of three or four members.			

Aides	Lesson Plan	Instructor Notes			
		Tell teams that they are to review three exemplars (Subjects Charles, Dodge and Edwards). Team members are to discuss the evidence among themselves and reach a conclusion concerning the category or categories of drugs, if any.			
		Teams will present their conclusions to the entire class.			
	a. Review and discussion of exemplars by teams.	Allow teams approximately 15 minutes to review the three exemplars and reach their conclusions.			
	b. Feedback of results.	Poll the teams to determine their conclusions concerning the category or categories of			
	o Subject Charles	drugs present in each subject.			
	o Subject Dodge	Offer appropriate comments			
	o Subject Edwards	concerning the teams performance.			
	2. Session wrap-up.	Solicit students' comments and questions concerning this practice session.			
		·			
	;				

## DRUG CATEGORIES FOR INTERPRETATION PRACTICE

SUBJECT

CATEGORY(IES)

Adams

CNS Depressant

Baker

**CNS Stimulant** 

Charles

Alcohol only (CNS Depressant)

 ${\bf Dodge}$ 

CNS Stimulant

Edwards

Hallucinogen

# **Session XV**

Practice: Test Interpretation



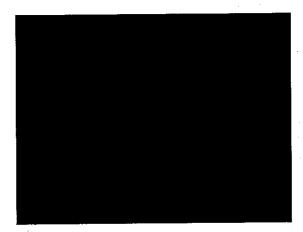
## **Practice: Test Interpretation**

Upon successfully completing this session, the participant will be able to:

- Analyze the results of a complete drug evaluation and classification examination and identify the category or categories of drugs affecting the individual examined
- Articulate the basis for the drug category identification

Drug Evaluation & Classification Training

xv.o



**Drug Influence Evaluation** Rolling Log No. PO-05 Crash: None □ Injury □ Property D Fatal Arresting Officer (Name, ID No.) DOB Sex MOHN. Breath Results: , OO Refused □ Urine Instrument# □ Refused Have you been drinking? Time of last drink? How much? How long? Are you diabetic or epileptic? When did you last sleep? Time now? □ Yes ■No Do you have any physical defects? Yes No Are you under the care of Are you taking any medication or drugs? 

Yes Tracking: □ Reddened Conjunctiva Eyes: Blindness: Speech & Equal □ Unequal ■ Normal □ Bloodshot □ Watery ■ None □LEye □REye Slow SLURRED, THICK TONGO Able to follow stimulus: 

Yes □ No Eyelids: 🗷 Equal □ Glasses □ Contacts, if so □ Hard □ Soft □ Unequal (explain) One Leg Stand Left Eye Right Eye Vertical Nystagmus? □ Yes □ No Pulse & Time HGN 405 Lack of Smooth Pursuit Convergence Right Eye Left Eye Max. Deviation Angle of Onset Cannot Keep Balance Romberg Balance Walk and Turn Test Starts Too Soon R Sways While Balancing Approx. 2nd Nine Stops Walking Uses Arms to Balance Misses Heel-Toe Steps Off Line □ Hopping Raises Arms Z Puts Foot Down Actual Steps Taker Type of Footwear Cannot Do Test (Explain) Describe Turn Internal Clock TURNED BACKWARDS .55 Estimated At 30 Sec. Hich Heelst Pupil Size Room Light Darkness Direct ○ Right △ Left Left Eye 3.0 Draw lines from spot touched Oral Cavity Right Eye Hippus O Yes Reaction To Light Rebound Dilation **⊘**No 🗆 Yes ₿No Temp_ Muscle Tone: ☐ Normal Attach Photos Of Fresh Puncture Marks at Medicine or Drug Have You Been Using? Where Were The Drugs Used? (Location) How Much? CFUSEd Time Completed □ Inhalant Opinion of Evaluator: Rule Out □ Alcohol □ Stimulant □ Medical □ Hallucinogen □ Narcotic Analgesic □ Cannabis Depressant

	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>				
LOG NO.	DRE: Officer Jim Brown	ARRESTEE: Frances A. Adams (f)				
5. INITIAL OBSERVATION	1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.					
1. LOCATION: DRE e	xamination room 4th District, Arizona Depa	urtment Public Safety				
2. WITNESS: Arresting	g Officer - Sgt. R. Hohn #2345 Arizona De	epartment of Public Safety				
3. BREATH TEST: W	riter administered GCI breath test to Adam	is, the result was 0.00%				
4. NOTIFICATION / I	INTERVIEW of ARRESTING OFFICER	:				
5. INITIAL OBSERVA	ATIONS: Writer observed subject seated ne	ext to the breath test instrument, her head was tilted				
forward, her eyes wer	e closed, her breathing was deep but slow.	She responded slowly to my questions and her speech				
was slow and slurred						
6. MEDICAL PROBL	EMS: None noted or stated					
7. PSYCHOPHYSICA	L TESTS:					
8. CLINICAL INDICA	ATORS:					
9. SIGNS of INGESTION: None were evident						
10. STATEMENTS: Subject stated that she was very sleepy, and denied taking any medicine or drugs.						
11. OPINION of EVALUATOR: In my opinion Frances A. Adams is under the influence of						
and unable to operate a vehicle safely						
12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a blood sample.						
13. MISCELLANEOUS:						

	Drug Influence Evaluation						
I	Evaluator 3005/6 C.	DRE No OO	19	Rolling Log No.	- 326		
Ì	Recorder/Witness Dyas, Sat.	Crash:	<b>⋉</b> None □ Injury	□ Prop			
( (	"restee's Name (Last, First, MI) Baker, Sam B.	DOB/5/197	'2 Sex	Race Arr	esting Officer (N	ame, ID No.) 7. W. *3	210 CTD
•	July 19, 2000 2230	3 ro Oistri	Et Bre Inst	ath Results: 0.00 rument#/23		Chemical Test  □ Refuse	□ Urine 🐞 Blood
		at have you eaten to			been drinking?	How much?	Time of last drink?
	Time now? When did you last sleep? How long		or injured?	□ Yes 🛒	No.	Are you diabetic o	or epileptic?
	Do you take insulin?	Do you have a	ny physical defe	cts? □ Yes 📽 l	No	Are you under the a doctor or dentis	
	Are you taking any medication or drugs?   Yes	~	poper	ative		Coordination Face	tumbling
	~	720	zncia eddened Conjunc	tivo	Blindness:	Normal	Sweaty Tracking:
	Rapid		Bloodshot DW		₩ None □LE	Cye □ R Eye stimulus: ♥ Yes □ No	Equal □ Unequal  Eyelids:
		Jnequal (explain)			Able to lonow	· · · · · · · · · · · · · · · · · · ·	Normal Droopy
	Pulse & Time HGN	Left Eye R	<u> </u>	cical Nystagmus?	□ Yes ■ No	One Leg St	and 40/04(27)
	1. 108 , 2235 Lack of Smooth Pursu	it <b>//</b> 0	<del></del>	vergence Right Eye	Left Eye	IN 30 S	ccons
	2. 1/2 / 2246 Max. Deviation	No	No			ິ່	
	3. /00 / 2253 Angle of Onset		None			<b>A</b>	I ¥
	Romberg Balance Walk and Turn Approx. Approx.	Walked val	oidly	Cannot Ke Starts Too		L_/ R/	
	3 Approx. 3 O Approx. 0			Stops Walk	ring 1st Nine	1 1/////	Sways While Balancing
	$\mathcal{L}$		)	Misses Hee		6/ 16/	Uses Arms to Balance
		menter		/ Steps Off L Raises Arm			Hopping
		5	-	Actual Step		9 0	Puts Foot Down
	Internal Clock Describe Turn	tructed	C	annot Do Test (E	Explain)	Type of Fo	otwear CEIS
			Pupil Size	Room Light	<u> </u>	Direct Nasal Area	Redness
	○ Right △ Left  Draw lines from spot touched	5 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 10 to 1	Left Eye	6.5	8.0	6.0 Oral Cavit	y nosc
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	1 0 HE	· K)	Hippus O	Yes Reboun		eaction To Light	w
			AkG	HT ARM		LEFT ARM	15
		3	<b>#</b>			/a mar	
					No VISI	ble	
	(5)	<del></del>	_		Mo		
	Muscle Tone: 《Normal □ Flaccid □ Rigid	emp 99.7.		Attach	Photos Of Fr	esh Puncture Ma	rks
	'at Medicine or Drug Have You Been Using?	How Much?	Time of Use?			Used? (Location)	
		1swer	No Ans		10 Ans	uier	
		TO TO TO A TO A TO A TO A TO A TO A TO	2200		tart Time 22	Time Comp	leted 2310
	Member Signification (Include Rank)		ID No. 888	Review		1101	*
	Opinion of Evaluator:   Rule Out	□ Alcohol □ Depressant	□ Stimulant □ Hallucino	□ PCP	otic Analgesic	□ Inhalant □ Cannabis	<u> </u>
	□ Medical	n pehtessaut	- manucino	Por - Hard	ATTO TITIOTE CORE	- Julianto	

	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>			
LOG NO.	DRE: Sgt. Clark John	ARRESTEE: Sam B. Baker			
5. INITIAL OBSERVATION	1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.				
1. LOCATION: DREE	Examination room 3rd District Capitol PD				
2. WITNESS: Arresting	GOfficer - Sgt. T. W. Tower # 3210 Capitol	PD and Sgt. Toby Dyas, Tempe Police Department			
3. BREATH TEST: W	riter observed Sgt. T. W. Tower administer	a breath test to Baker, the result was 0.00%			
4. NOTIFICATION / I	INTERVIEW of ARRESTING OFFICER	:			
5. INITIAL OBSERVA	ATIONS: Writer observed subject standing	next to the breath test instrument. He repeatedly			
shifted his weight fror	n foot to foot, and scratched his face and her	ad. He was perspiring heavily, and appeared			
nervous, anxious and j	ittery				
6. MEDICAL PROBLE	EMS: None noted or stated				
7. PSYCHOPHYSICAL TESTS:					
8. CLINICAL INDICA	ATORS:				
9. SIGNS of INGESTION: Reddened nasal area.					
10. STATEMENTS: Subject denied taking any medicine or drugs.					
11. OPINION of EVALUATOR: In my opinion Sam B. Baker is under the influence of					
and unable to operate a vehicle safely					
12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a blood sample.					
13. MISCELLANEOUS:					

**Drug Influence Evaluation** Rolling Log No. Evaluator 000518 Recorder/Witness Crash. □ None □ Fatal □ Property □ Injury rrestee's Name (Last, First, MI) Sex Arresting Officer (Name, ID No.) 5083 WPD CHARLES, MARY C 6/3/1972 F Examined/Time/Location EXAMINATION ROOM AR 17,1996 0045 4" DISTRICT W SHELMAN, S Breath Results: Chemical Test □ Urine Instrument# □ Refused Miranda Warning Given: What have you eaten today? Have you been drinking? How much? Time of last drink? P122A SHERMAN, S CASTNIGHT , NOT A COUPLE OF BEERS Are you sick or injured? When did you last sleep? How long? Are you diabetic or epileptic? □ Yes ¥No Do you have any physical defects? UYes XNo Are you under the care of a doctor or dentist? □ Yes XNo Coordination / STALLERING-Are you taking any medication or drugs? ♥Yes □ No Attitude COOPERATIVE BIRTH CONTROL PILLS AL FLUSHED Tracking: Eyes: ☐ Reddened Conjunctiva Blindness: Speech SLURRED K Equal □ Unequal □ Normal A Bloodshot Watery % None □LEye □REye Pupil Size: KEqual Able to follow stimulus: A Yes □ No Eyelids: □ Normal ADroopy □ Glasses □ Contacts, if so □ Hard □ Soft □ Unequal (explain) One Leg Stand HGN Pulse & Time Left Eye Right Eye Vertical Nystagmus? □ Yes > No 10050 Convergence Right Eye Lack of Smooth Pursuit YES Left Eye Max. Deviation Angle of Onset LIBBEL, Letter Cannot Keep Balance Romberg Balance Starts Too Soon_ Sways While Balancing 1st Nine 2nd Nine Stops Walking Uses Arms to Balance Misses Heel-Toe Steps Off Line □ ,Hopping Raises Arms CONSTANT CONSTA Puts Foot Down Actual Steps Taken Internal Clock Describe Turn Cannot Do Test (Explain) Type of Footwear Estimated At 30 Sec. LOST BALANCED/STALLERED TENNIS Shoe S Pupil Size Room Light Darkness Direct ○ Right △ Left Left Eye Draw lines from spot touched Oral Cavity Right Eye Rebound Dilation Hippus □ Yes Reaction To Light **⋉**No □ Yes XNo SZON Temp 98.0 Blood Pressure //O Muscle Tone: □ Normal 🏋 Flaccid □ Rigid Attach Photos Of Fresh Puncture Marks Comments: at Medicine or Drug Have You Been Using? Where Were The Drugs Used? (Location) How Much? Time of Use? INE, JUST MX PILL NO ANSWER ANSWER NOANSWER 110 Time DRE Notified Eval Start Time Time Completed 0025 Member Signature (Include Rank) 70*0* Opinion of Evaluator: □ Rule Out □ Alcohol □ Stimulant □ Inhalant □ Cannabis □ Medical □ Hallucinogen □ Narcotic Analgesic □ Depressant

	DRUG INFLUENCE EVALUATION	Page_2_of_2				
LOG NO.	DRE: Sgt. Michael Hayes	ARRESTEE: Mary C. Charles				
5. INITIAL OBSERVATI	LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR.     INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS     SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.					
1. LOCATION: DRE	Examination room 4 th District Washington	State Patrol				
2. WITNESS: Arresting	g Officer - S. Shermann # 5083 Washington	n State Patrol and Sandy Richardson, NHTSA				
3. BREATH TEST: W	riter observed Officer Shermann administer	a breath test to Charles, the result was 0.09%				
4. NOTIFICATION /	INTERVIEW of ARRESTING OFFICER	<b>k</b> :				
5. INITIAL OBSERV	ATIONS: Writer observed subject in the ho	lding area of central booking, she was staggering				
and stumbling, she sv	wayed and repeatedly blinked her eyes and h	er speech was very slurred				
6. MEDICAĻ PROBL	LEMS: None noted or stated					
7. PSYCHOPHYSICA	AL TESTS:					
		•				
8. CLINICAL INDIC	ATORS:					
9. SIGNS of INGEST	TON: Subject had an odor of alcoholic beve	rage on her breath.				
10. STATEMENTS: Subject admitted she had been drinking. However, she denied taking any medicine or using any						
drugs other than bi	rth control pills.					
11. OPINION of EVALUATOR: In my opinion Mary C. Charles is under the influence of						
and unable to operate a vehicle safely						
12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a blood sample.						
13. MISCELLANEOUS:						

000500	Drug In	nuen			<u> </u>	<del></del>	<del></del>	
Evaluator 000520 11 DWELL, JERRY	DRE No			g Log No.				
Recorder/Witness	Crash: □ Fatal	□ Non □ Inju		□ Prop				
stee's Name (Last, First, MI)	DOB	Sex	Race			(Name, ID No.)	CU HTA	
SODE, FRED D	10/13/65	1 / ^ /	Brooth Pos	uite: 🗘 🗘	AIRO 1	C.D. 76	Urine KBlood	
nstee's Name (Last, First, MI)  JOLE, FRED  Date Examined/Time/Location 5th 015TR  FEB 22, 1997 23(11)	51000	٥٨	Instrument	120	Refused	□ Refi	Time of last drink?	
Miranda Warning Given: XYes No What By: (ARD, C.)	have you eaten toda $IACOS$ 2	19? W. 2-3 HR	hen? ?s Ato	/	NoniNC	- N/A	NIT	
Time now? When did you last sleep? How long?  //O'CNCK YESTERDAY OR SHRS	Are you sick or i	njured?		Yes 🕍	Vo	Are you diabeti □ Yes 🕱 N		
Do you take insulin?	Do you have any	y physical	defects? □	Yes 🔌	Vo	Are you under t a doctor or dent	he care of ist?   Yes No	
Are you taking any medication or drugs?   Yes  No	CAR	FFREE, COOPERATIVE			a TI UK	Coordination	Coordination J. 178RY, STUMBLING	
		<i>y</i>	MAL	-		Face NC	Face NORMAL	
Speech RAFIN	_, _, _, _, _, _, _, _, _, _, _, _, _, _	ldened Conjunctiva Blindness: Bloodshot □ Watery None □ I			⊠None □	LEye □REye	Tracking:	
COLLEGIA POLICE	Size: # Equal equal (explain)				Able to follo	w stimulus: 🗯 Yes □ N	Eyelids: XQ Normal □ Droopy	
Pulse & Time HGN		ht Eye	Vertical Ny	stagmus?	XYes □ No	One Leg	Stand	
1. 100 12305 Lack of Smooth Pursuit	NO N		Convergen Right		Left Eye		<b>പ</b>	
2./04 /23/( Max. Deviation	NO N		0	$\ni$		Q	0 0	
3/00 123 26 Angle of Onset  Romberg Balance Walk and Turn To	1 0/ 0	WE			ep Balance			
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Steps Off Line Raises Arms								
Raises Arms Actual Steps Taken								
Internal Clock	USTRUCTE	^		Do Test (E	xplain)		Footwear ReeT Shoe S	
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Draw lines from spot touched	,	Left Eye	5.	5	8.5	5.5 Oral Ca	vitv	
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3	<u>×6</u>	FOUR	/炒			-		
Blood Pressure 190 / 96 Temp 99.5.								
Muscle Tone: Normal   Flaccid   Rigid   Attach Photos Of Fresh Puncture Marks Comments:								
at Medicine or Drug Have You Been Using?	How Much?	Time of	Use? NSWE/Z	Where	Were The Dru ∕\{∧	igs Used? (Location)  ANSWER		
Date/Time of Agrest 997/22/5 Tim	ne DRE Notified	15		Eval St	art Time	Time Con	npleted 3.3 O	
Member Signature (Include Rank)		11) 199	<i>7</i> 2	Review	red By:			
Opinion of Evaluator:   Rule Out  Medical	□ Alcohol □ Depressant	□ Stim □ Hall	ulant ucinogen	□ PCP □ Narc	otic Analgesic	□ Inhalan □ Cannab		

	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>			
LOG NO.	DRE: Lt. Jerry Tidwell	ARRESTEE: Fred D. Dodge			
1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.					
1. LOCATION: DRE	Examination room 5th District HTD				
2. WITNESS: Arresting	g Officer - C. D. Laird #7654 HTD				
3. BREATH TEST: O	fficer Laird administer a breath test to Fred I	Dodge, the result was 0.00%			
4. NOTIFICATION /	INTERVIEW of ARRESTING OFFICER	:			
5. INITIAL OBSERV	ATIONS: Writer observed subject at 2255 h	us. In the breathalyzer room. He was smiling and			
joking with officer I	Laird. Dodge's speech was rapid and loud.	He seemed boisterous and unconcerned about being			
under arrest.					
6. MEDICAL PROBL	EMS: None noted or stated				
7. PSYCHOPHYSICA	AL TESTS:				
8. CLINICAL INDICA	ATORS:				
	•				
9. SIGNS of INGEST	ION: Subject had four (4) fresh puncture wo	ounds on the underside of his left forearm.			
10. STATEMENTS: S	ubject denied taking any medicine or using	g any drugs. When questioned about the punsture			
marks he grinned ar	nd stated "Gee, I guess those must be mosqu	ito bites", then laughed.			
11. OPINION of EVALUATOR: In my opinion Fred D. Dodge is under the influence of					
and unable to operate a vehicle safely					
12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a blood sample.					
13. MISCELLANEOU	JS:				

000522	Drug In	fluence E	olling Log No.	on		· ·
Evaluator UNSWORTH, J		□ None				
Recorder/Witness	Crash: □ Fatal	□ None □ Injury	□ Prop	erty		
estee's Name (Last, First, MI)	ров			esting Officer (Nan	ne, ID No.)	000
WARDS JOAN E	01/11/69	, .	i		± 2456	CRD
Date Examined/Time/Location	0.00		h Results: O.C.		Chemical Test  ☐ Refused	□ Urine <b>▼</b> Blood d
APRIL 1, 1996 2300	CPD at have you eaten too		17:	been drinking?	How much?	Time of last drink?
By: HALL, I	VOTHING	N/A	Νo	THING		NA
Time now? When did you last sleep? How long? Are you sick or injured? Yes a No Are you diabetic or epileptic?  I DON'T TO YES ARE YOU THEN THE STORMACH						
KNEW I UDNI NEMEMBER I TELL SIZE IO III I Vas SNO Are you under the care of						
Do you take insum:	1 '	r THINK	_		a doctor or dentist	? DYes \$No
Coordination				RIENTED STAGGER		
	D 414		121(17 1 10			ZED APPEARANCE
Speech	Eyes: □ Rec	JokmAL ddened Conjunction	va	Blindness:		Tracking:
DIFFICULTY IN SPEAKING AT TIMES	In James Nati	Bloodshot □ Wat	ery	None □LEy		E Equal □ Unequal
Corrective Lens: None Pu	pil Size: Equal			Able to follow sti	mulus: 🗗 Yes 🗆 No	Eyelids: Normal © Droopy
□ Glasses □ Contacts, if so □ Hard □ Soft □ □	Jnequal (explain)					- Marian - Droopj
Pulse & Time HGN	Left Eye Ri	ght Eye Vertic	al Nystagmus?	□ Yes #No	One Leg Sta	and O
Lack of Smooth Pursu	it NO	NO Conve	ergence		NO!	D PA
100 3326			Light Eye	Left Eye		
151 2227		\	***		9-TE	
3. 104 / 2337 Angle of Onset  Romberg Balance Walk and Turn		ONE	Cannot Ke	ep Balance		STOPPED
Approx. Approx.	uTest M M M	5 M KI	Starts Too		L R	Sways While Balancing
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$\mathcal{A}$	MMMMW	# 4 W	Misses Hee	1444444	MILL MEN GAN	Jses Arms to Balance
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90 Estimated At 30 Sec. TURNED	BACKWAR	Pupil Size	Room Light		irect Nasal Area	- A
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Muscle Tone:   Normal   Flaccid   Rigid   Comments:   A a see   A			Attach	Photos Of Fre	sh Puncture Mai	cks
Comments: ARMS YEAY RIBIA  at Medicine or Drug Have You Been Using?	How Much?	Time of Use?	Where	Were The Drugs U	Jséd? (Location)	
at Medicine or Drug Have You Been Using!	No ANSWER	No ANSW	ا ایا	Vo ANSWE		
	Fime DRE Notified	2245	Eval St	tart Time 300	Time Compl	93.45
Date/Time of Arrest 996 9-7-35  Member Signature (Include Hank)		ID No. 54		red By:		<del></del>
Member Signature (Include Hank) Oninion of Evaluator:	□ Alcohol	□ Stimulant	□ PCP		□ Inhalant	
Opinion of Evaluator:   Rule Out  Medical	□ Depressant	□ Hallucinog		otic Analgesic	□ Cannabis	

LOG NO. DRE: Officer J. Unsworth ARRESTEE: Joan E. Edwards  1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.  1. LOCATION: DRE Examination room 5th District CTD  2. WITNESS: Arresting Officer - Ian Hall # 3456 CTD  3. BREATH TEST: Officer Hall administer a breath test to Joan E. Edwards, the result was 0.00%  4. NOTIFICATION / INTERVIEW of ARRESTING OFFICER: Writer was contacted by Officer Hall at 2255 hrs. Officer Hall stated he had just arrested a "very weird" woman. He further stated "she's either on drugs or crazy." Her vehicle was stopped in the intersection of Studdard Ave. and Haversat Dr., she was standing on the hood of her car waving her arms and screaming incoherently at passing traffic.  5. INITIAL OBSERVATIONS:  6. MEDICAL PROBLEMS: Subject stated indicated some nausea.
<ol> <li>LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR.</li> <li>INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS</li> <li>SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.</li> <li>LOCATION: DRE Examination room 5th District CTD</li> <li>WITNESS: Arresting Officer - Ian Hall # 3456 CTD</li> <li>BREATH TEST: Officer Hall administer a breath test to Joan E. Edwards, the result was 0.00%</li> <li>NOTIFICATION / INTERVIEW of ARRESTING OFFICER: Writer was contacted by Officer Hall at 2255 hrs. Officer Hall stated he had just arrested a "very weird" woman. He further stated "she's either on drugs or crazy." Her vehicle was stopped in the intersection of Studdard Ave. and Haversat Dr., she was standing on the hood of her car waving her arms and screaming incoherently at passing traffic.</li> <li>INITIAL OBSERVATIONS:</li> </ol>
<ol> <li>5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.</li> <li>1. LOCATION: DRE Examination room 5th District CTD</li> <li>2. WITNESS: Arresting Officer - Ian Hall # 3456 CTD</li> <li>3. BREATH TEST: Officer Hall administer a breath test to Joan E. Edwards, the result was 0.00%</li> <li>4. NOTIFICATION / INTERVIEW of ARRESTING OFFICER: Writer was contacted by Officer Hall at 2255 hrs. Officer Hall stated he had just arrested a "very weird" woman. He further stated "she's either on drugs or crazy." Her vehicle was stopped in the intersection of Studdard Ave. and Haversat Dr., she was standing on the hood of her car waving her arms and screaming incoherently at passing traffic.</li> <li>5. INITIAL OBSERVATIONS:</li> </ol>
<ol> <li>WITNESS: Arresting Officer - Ian Hall # 3456 CTD</li> <li>BREATH TEST: Officer Hall administer a breath test to Joan E. Edwards, the result was 0.00%</li> <li>NOTIFICATION / INTERVIEW of ARRESTING OFFICER: Writer was contacted by Officer Hall at 2255 hrs. Officer Hall stated he had just arrested a "very weird" woman. He further stated "she's either on drugs or crazy." Her vehicle was stopped in the intersection of Studdard Ave. and Haversat Dr., she was standing on the hood of her car waving her arms and screaming incoherently at passing traffic.</li> <li>INITIAL OBSERVATIONS:</li> </ol>
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5. INITIAL OBSERVATIONS:
6. MEDICAL PROBLEMS: Subject stated indicated some nausea.
7. PSYCHOPHYSICAL TESTS:
8. CLINICAL INDICATORS:
9. SIGNS of INGESTION: None were evident.
10. STATEMENTS: Subject denied taking any medicine or using any drugs.
11. OPINION of EVALUATOR: In my opinion Joan E. Edwards is under the influence of
and unable to operate a vehicle safely
12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a blood sample.
13. MISCELLANEOUS: Subject was transported to the psychiatric ward, at the county jail, for continued monitoring.

One Hour and Forty Minutes

SESSION XVI
PHENCYCLIDINE (PCP)

# SESSION XVI PHENCYCLIDINE (PCP)

Upon successfully completing this session, the participant will be able to:

- o Explain a brief history of PCP.
- o Identify common drug names and terms associated with PCP.
- o Identify common methods of administration for PCP.
- o Explain the symptoms, observable signs and other effects associated with PCP.
- o Explain the typical time parameters, i.e., onset and duration of effects, associated with PCP.
- o State the clues that are likely to emerge when the drug evaluation and classification process is conducted for a person under the influence of PCP.
- o Correctly answer the "topics for study" questions at the end of this Section.

Content Segments		Learning Activities	
A.	Overview of the Category	0	Instructor Led Presentations
B.	Possible Effects	o	Review of Drug Evaluation and Classification Exemplars
C.	Onset and Duration of Effects	0	Reading Assignments
D.	Overdose Signs and Symptoms	O	Video Presentations
E.	Expected Results of the Evaluation	0	Slide Presentations

#### **Instructor Notes**





XVI-0A&B (Session Objectives)



25 Minutes



PHENCYCLIDINE (PCP)

Total Lesson Time: Approximately 100 Minutes

Session title on wall chart.

Briefly review the objectives, content and activities of this session.

A. Overview of the Category

The chemical name for PCP is PhenylCyclohexyl Piperidine.

Write the chemical name on the chalkboard or flip chart, underlining the first "P", the first "C" and the last "P".

Point out that "Phencyclidine" is a contraction, or shortened form of the chemical name.

Point out that an "analog" is a chemical that is very similar to the drug in terms of molecular structure or in psychoactive

effects.

- 1. Phencyclidine or PCP, is a drug that, along with its <u>analogs</u>, forms a distinct category.
- 2. PCP shares some characteristics with each of the three categories of drugs previously covered in this training.
  - a. It produces some effects that are similar to the effects of CNS Depressants.
    - b. It produces some effects that are similar to those of CNS Stimulants.

Examples of effects PCP shares with Depressants: Nystagmus, slurred speech, slowed responses.

Examples of effects PCP shares with CNS Stimulants: elevated vital signs, frenzied behavior.

#### **Instructor Notes**

c. In some respects it acts like a Hallucinogen.

Point out that in many medical texts and other reference documents, PCP may be classified as a Hallucinogen. However, for purposes of the Drug Evaluation and Classification procedure, it is treated as a separate category.

Developed by Parke-Davis and Company, a leading pharmaceutical firm.

- 3. Phencyclidine was first developed in the late 1950s.
  - a. The developers were searching for a drug that would serve as an efficient intravenous anesthetic.
  - b. PCP proved to be a very effective anesthetic.
  - c. It was patented and marketed in 1963 under the trade name <u>Sernyl</u>.
  - d. It was used in the treatment of mental and psychological disorders, including schizophrenia and alcoholism.
  - e. Many adverse side effects were experienced by persons who had been treated with PCP.
  - f. In 1967, use of
    Phencyclidine as an
    anesthetic for humans was
    discontinued.

An <u>anesthetic</u> is an agent that reduces or abolishes sensation.

<u>Sernyl</u> derives from the word <u>serene</u>, the apparent mood induced by PCP. In fact, however, the PCP user often is very far from "serene".

<u>Point out</u> that some of these side effects will be discussed later.



#### **Instructor Notes**

g. In 1968, Parke-Davis repatented PCP under the trade name <u>Sernylan</u>, which was restricted to use as a veterinary anesthetic.

 $\underline{Sernyl}$  for  $\underline{an}$ imals =  $\underline{Sernylan}$ .

Point out that this is why PCP sometimes goes by the "street" names "Monkey Dust"; "Elephant Tranquilizer"; "Horse Tranquilizer"; etc.

Print Ketamine on the

chalkboard or flip chart.

Some brand names of

and Vetalar.

Ketamine: Ketalar, Ketaject

- h. However, Sernylan was often illicitly diverted to "street" use, so most legitimate manufacturing of PCP was stopped in 1978.
- i. Another drug in this category is called **Ketamine**. It continues to be manufactured and sold legitimately.
- Ketamine is used as a surgical anesthetic, both for animals and humans, especially children.
- k. Ketamine is also used for burn victims.
- 4. Continuing research demonstrated that PCP consistently produced adverse side effects.
- Delirium: confusion, disordered speech, frenzied excitement, hallucinations.



XVI-2 ("Adverse Side Effects")

- a. delirium
- b. visual disturbances, hallucinations
- c. agitation, anxiety
- d. rigid muscle tone



#### **Instructor Notes**

- e. elevated blood pressure
- f. convulsions
- g. difficulty in speech
- h. violent reactions
- 5. Some lingering and long term effects were also noted.
  - a. Some patients complained of dizziness for several hours after their attention and consciousness appeared to be cleared of PCP's effects.
  - b. Some patients reported memory disorders and other psychological disorders resembling schizophrenia for several months and even years afterwards.
- 6. Cases of terribly bizarre, self destructive behavior have been reported with persons under the influence of PCP.
  - a. One young man methodically pulled his own teeth out, using a pair of pliers.

Convulsion: involuntary contortion of the muscles, producing contortion of the body and limbs.

PCP has sometimes been called a <u>psychotomimetic</u> drug; i.e., it produces effects that mimic psychosis, or "craziness". When the craziness remains long after the drug has dissipated, we say that its effects were <u>psychotogenic</u>, i.e., it didn't simply mimic craziness, it caused craziness.

<u>Point out</u> that PCP can render the user impervious to pain. It anesthetizes the central nervous system to the extent that surgery could be performed on the user while he or she is wide awake.

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#### **Instructor Notes**

- b. Another individual suffered hallucinations of unbelievably grotesque monsters, and gouged out his own eyes to avoid seeing the monsters.
- c. Another young man drank rat poison, attempting to kill rats that he imagined were inhabiting his body.
- 7. PCP is relatively easy to manufacture.
  - The chemicals required to produce it are readily available commercially.
  - b. The formula for producing PCP has been widely publicized.
  - c. The hardware needed to combine the chemicals is very basic.

NOTE: Instructors should feel free to replace or supplement these examples with others known personally to them.

Emphasize, however, that there is some danger present in the manufacturing process. Illicit PCP laboratories frequently explode and burn.

Note that PCP labs commonly contain potassium cyanide and hydrochloric acid. If combined, those two chemicals produce the same lethal gas used in gas chambers designed for executions.

Emphasize that officers should exercise great caution when they discover an illicit PCP lab.

Review the policy and procedures of the students' department for dealing with PCP labs and materials.

#### Aides

#### Lesson Plan

#### Instructor Notes



XVI-4 ("Methods of Ingestion")

- 8. Numerous "street names" are used to designate PCP.
- 9. Methods of ingestion of PCP.
  - a. Many users ingest PCP by smoking.
  - b. PCP can be applied in either powder or liquid form to a variety of vegetable or leafy substances, which can then be smoked in a pipe or home made cigarette.
  - Popular substances include mint leaves, parsley, oregano, tobacco or Marijuana.
  - d. Commercially prepared cigarettes can also be dipped in liquid PCP, allowed to dry and then smoked.
  - e. Some users prefer to dip a string in liquid PCP, and then insert the string into a tobacco cigarette.
  - f. PCP can also be <u>insufflated</u> or "snorted".
  - g. It can also be taken <u>orally</u>, in capsule or tablet form.

Briefly review the more common street names in vogue in the students' communities.

If available, display 35 mm slides of the various PCP ingestion paraphernalia.

NOTE: Liquid PCP is especially dangerous because it can be absorbed through the skin. Hence, it could be used as a weapon.

<u>Point out</u> that PCP smoke is very hot and can irritate the mouth and tongue. Mint leaves and similar material help to cool the smoke.

<u>NOTE</u>: PCP adulterated cigarettes usually will be wrapped in metal foil to be preserved.

Point out that "Kool" and "Sherman" brand cigarettes are popular for this, because they are mentholated. PCP-adulterated cigarettes are sometimes called "Super Kools" or "Sherms".

NOTE: White cigarette paper will be stained brown if adulterated with PCP. Brown cigarette paper will show white crystals, when adulterated.

# **Instructor Notes**

- h. Some users <u>inject</u> liquid PCP, either directly into a vein or under the skin or into a muscle.
- Some users have administered PCP to themselves by dropping liquid PCP onto their eyes, using an eyedropper.
- j. Transdermal absorption of PCP has also been reported (i.e., when applied to the skin, especially as a liquid, PCP can penetrate directly into the body and bloodstream).

Reemphasize the danger to officers handling suspected drugs without proper protective gloves.

Solicit students' questions and comments about the overview of PCP.



- B. Possible Effects
  - 1. PCP produces impairments and other observable effects on the human mind and body that are a combination of some of the effects associated with CNS Depressants, CNS Stimulants and Hallucinogens.
    - a. Slow, slurred speech.
    - b. Disorientation.
    - c. Loss of memory.
    - d. Agitation, excitement.
    - e. Blank stare.

<u>Point out</u> that these effects will not necessarily appear in a predictable sequence as dose increases.

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	A	id	es	3

#### **Instructor Notes**

- Passivity, but may abruptly turn violent if confronted with a situation perceived to be threatening.
- Muscle tone rigid.
- h. Noncommunicative.
- Depersonalization, loss of the sense of personal identity.
- Sensory distortions, hallucinations.
- k. Perspiring.
- Increased pain threshold
- Regular users of PCP develop a tolerance that may mask many of these observable effects.
- PCP has been called a Dissociative Anesthetic, i.e., it cuts off the brain's perceptions of the senses.
  - a. PCP users often feel that their heads are physically separated from their bodies.
  - b. They sometimes report feeling they are dead, and that their heads are floating away.
- C. On-set and Duration of Effects

Studies show that between 3-30% of PCP subjects will exhibit violence.

#### Clarification:

"Depersonalization" is a syndrome characterized by a feeling of unreality and of estrangement from one's self, body or surroundings.

Note: Especially auditory hallucinations.

Solicit students' questions and comments concerning possible effects of PCP.

Write "Dissociative Anesthetic" on the chalkboard or flip chart.





5 Minutes

# **Instructor Notes**



XVI-6

- 1. The onset of PCP's effects varies somewhat with the method of ingestion.
  - a. When PCP is smoked or injected, onset occurs within 1-5 minutes.
  - b. When inhaled ("snorted") onset occurs in 2-3 minutes.
  - c. Onset is considerably slower when PCP is taken orally: 30-60 minutes.
- 2. The effects reach their peak in about 15-30 minutes, assuming the PCP was smoked, injected or snorted.
- 3. The effects generally last 4-6 hours, but they can go somewhat longer.
- 4. The user usually, but not always returns to normal within 24-48 hours.
- D. Overdose Signs and Symptoms
  - In addition to the bizarre, violent and self destructive behavior discussed previously, persons severely intoxicated by PCP may exhibit definite and extreme symptoms signifying a medically dangerous condition.
    - a. A deep coma, lasting for up to 12 hours.
    - b. Seizures and convulsions.
    - c. Eyes generally open and staring blankly.

Solicit students' questions and comments concerning onset and duration factors.



5 Minutes

#### Aides

#### Lesson Plan

#### **Instructor Notes**

- 2. A danger associated with severe PCP intoxication is that the person may die due to respiratory depression.
- 3. There is also some evidence that PCP may trigger a heart attack, if the user had some preexisting condition disposing him or her to possible cardiac problems.
- 4. There is also some evidence that prolonged use of PCP can lead to psychosis, which can be permanent.
- E. Expected Results of the Evaluation
  - 1. Observable evidence of impairment.
    - Standardized Field Sobriety Tests.
      - o Horizontal Gaze
        Nystagmus generally
        will be present with a
        very early angle of
        onset.
      - Vertical Gaze Nystagmus usually will be present.
      - Performance on Romberg and Finger to Nose will be impaired: muscle tone will usually be rigid. Internal clock may be slowed.

A good practice (followed by LAPD) is to take all PCP suspects to medical treatment.

Solicit students questions and comments concerning signs and symptoms of PCP overdose.

NOTE: So-called "Resting Nystagmus" may be evident, especially with high doses. That is a distinct jerking of the eyeballs even as the suspect stares straight ahead.



60 Minutes



XVI-7A ("SFST evidence")



continued)

#### **Instructor Notes**

- o Performance on Walk and Turn and One Leg Stand will be impaired: muscle tone will be rigid.
- o Suspect may neglect to estimate the passage of 30 seconds when performing the Romberg test.
- b. General indicators:
  - o blank stare
  - o warm to the touch
  - o perspiring
  - o muscle tone rigid
  - o speech slurred and slow repetitive speech
  - o non-responsive, slow responses
  - o confused
  - o agitated
  - o may become suddenly violent
  - o chemical odor (of Ether, used in preparation of PCP)
  - o self-reported hallucinations



Suspect may exhibit a "high gait ataxia" or "moon walking", i.e., taking abnormally high and slow steps, as though he or she were trying to step over obstacles in his or her path.

Aides	Lesson Plan	Instructor Notes
	o cyclic behavior	NOTE: "Cyclic behaviors" mean that the signs and symptoms tend to increase and decrease cyclically.
	2. Evidence associated with the physiologic examinations.	
	a. Eye examinations:	
XVI-7D ("Eye Exam- inations")	o Lack of Convergence generally will be present.	
	o pupil size generally will be normal	
	o reaction to light generally will be normal	
	b. Vital signs examinations:	
XVI-7E ("Vital Signs Examina-	o blood pressure will be up	
tions")	o pulse will be up	-
<b>5</b>	o body temperature will be up	Point out that people under the influence of PCP often feel so hot that they remove all of their clothing.
	3. Summary	

XVI-8 ("PCP Symptomatol ogy Chart")

4. Demonstrations

Point out that tolerance may reduce some PCP symptoms.

Aides	Lesson Plan	Instructor Notes
	a. Video tape demonstrations	Show video tape of suspect(s) under the influence of PCP. Relate behavior and observations to the PCP symptomatology chart.
	b. Drug Evaluation and Classification exemplars demonstrations.	Refer students to the exemplars found at the end of Section XVI of their student manuals.
	·	Relate the items noted on the exemplars to the PCP Symptomatology Chart.
		Solicit students' questions or comments concerning Expected Results of the Evaluation of PCP suspects.
·		

# **Session XVI**

PCP and Its Analogs



# Phencyclidine (PCP)

Upon successfully completing this session, the participant will be able to:

- · Explain a brief history of PCP
- Identify common drug names and terms associated with PCP
- Identify common methods of administration for PCP
- Explain the symptoms, observable signs and other effects associated with PCP

Drug Evaluation & Classification Training

XVI-O/

# Phencyclidine (PCP) (continued)

- Explain the typical time parameters, i.e., on-set and duration of effects, associated with PCP
- State the clues that are likely to emerge when the drug evaluation and classification process is conducted for a person under the influence of PCP
- Correctly answer the "topics for study" questions at the end of this section

Drug Evaluation & Classification Training

XVI-O

### **Brief History of PCP**

- · Developed in the late 1950's
- · An effective intravenous anesthetic
- · Patented in 1963; trade name "Semyl"
- · Used in treating mental and psychological disorders
- · Very undesirable side effects were noted
- · Use as an anesthetic for humans
- · Was discontinued in 1967
- Re-patented in 1968 as an animal tranquilizer: trade name "Sernylan"

Drug Evaluation & Classification Training

XVL-1

#### Some Adverse Side Effects of PCP

- Delirium
- Blood pressure -- up
- · Hallucinations
- Convulsions
- · Agitation, anxiety
- Difficulty in speech
- · Muscle tone -- rigid
- · Violent reactions

Drug Evaluation & Classification Treining

XVI-2

#### Some "Street Names" for PCP

- Ace
- Krystal
- Amoeba
- KJ (Or CJ)
- Trank
- Devil Dust
- Jet Fuel
- KJ Krystal
- Juice
- Angel Dust
- Dust
- Angerbust
- Dust
- Krystal Joints
- Magic Dust
- Embalming FluidMonkey Tranquilizer
- Monkey DustCrystal Joints
- Lovely
- Crystal

Drug Evaluation & Classification Training

XVI-3A

#### More "Street Names" for PCP

- · Peace Pill
- Paz
- Green
- · Elephant Tranquilizer
- · Horse Tranquilizer
- Animal Tranquilizer

- Peace Weed · Mint Weed

Super Kools

Super Grass

Super Weed

Zombie Weed

- Green Leaves
- Killer Weed
- Tic Tac
- Sherms

Dove Evaluation & Classification Tesicine

# Methods of Ingesting PCP

- Smoking
- Orally
- · Injection
- Eyedropper
- · Insufflation (inhaling, "snorting")



#### **Possible Effects of PCP**

- · Slow, slurred speech
- Disorientation
- · Loss of memory · Agitation, excitement
- Blank stare
- Passivity...but possibly abruptly turning violent
- · Muscle tone rigid
- Noncommunicative
- Depersonalization
- Sensory distortion, hallucinations
- Excessive perspiration
- Increased pain threshold

# **On-set and Duration** of PCP's Effects

On-set

Smoked: 1-5 minutes Injected: 1-5 minutes Snorted: 2-3 minutes Orally: 30-60 minutes

> Peak effects Generally in 15-30 minutes

> > <u>Duration</u> 4-6 hours

# **Evaluation of Suspects** Under the Influence of PCP

#### SFST Evidence:

- · Horizontal Gaze Nystagmus will be present with a very early angle of onset (maybe "immediate" or even "Resting" Nystagmus)
- Vertical Gaze Nystagmus will be present
- Impaired performance will be evident on Walk and Turn and One Leg Stand Tests

Drug Evaluation & Classification Training

# **Evaluation of Suspects** Under the Influence of PCP

SFST Evidence (continued):

- · Impaired performance will be evident on Romberg and Finger To Nose
- · Muscle tone will be rigid

XV1-78

# **Evaluation of Suspects Under the Influence of PCP**

#### **General Indicators:**

- Blank stare
- · Warm to the touch
- Perspiring
- Muscle tone rigid
- Speech slurred and repetitive
- Non-responsive, slow responses

- Confused
- Agitated
- · May become suddenly violent
- · Chemical odor
- · Self-reported hallucinations
- · Cyclic behavior

XVI-7C

# **Evaluation of Suspects Under the Influence of PCP**

#### Eye Examinations:

- · Lack of Convergence present
- · Pupil size will be normal
- · Pupillary reaction to light will be normal

# **Evaluation of Suspects** Under the Influence of PCP

#### Vital Signs Examination:

- · Blood pressure will be up
- · Pulse will be up
- · Body temperature will be up

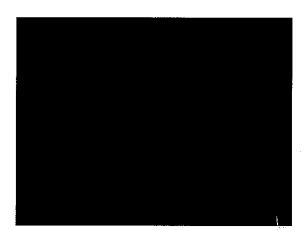
Drug Evaluation & Classification Training

XVI-7E

# **PCP Symptomatology Chart**

HGN	Present
VGN	Present
Lack of Convergence	Present
Pupil Size	Normal
Reaction to Light	Normal
Pulse Rate	Up
Blood Pressure	Up
Temperature	Up
Muscle Tone	Rigid

Drug Evaluation & Classification Training



Drug Influence Evaluation Rolling Log No. 284 Evaluator 000542 GEORGE, 4265 MARK Recorder/Witness Crash: M None  $\square$  Property □ Fatal □ Injury - ^ restee's Name (Last, First, MI) DOB Sex Race Arresting Officer (Name, ID No.) 9.6.59 BROWN A # 1832 MYSP Н OSS ROBERT .ate Examined/Time/Location Chemical Test □ Refused □ Urine XBlood Breath Results: NYSP Instrument# □ Refused DEC 8 2000 2145 HES 0.00 DISTRICT 17831 Miranda Warning Given: XYes □ No What have you eaten today? Have you been drinking? When? How much? Time of last drink? LAM "DOTHING" FRIED N/A 7137 CHICKEN When did you last sleep? How long? Are you diabetic or epileptic? Time now? Are you sick or injured? □ Yes 🕱 No □ Yes KNo 8 O'CLIEK YESTERDAY 6 HRS Do you take insulin? □ Yes 🛪 No Do you have any physical defects? Tes X No Are you under the care of a doctor or dentist? □ Yes No Are you taking any medication or drugs? ☐ Yes X No Attitude Coordination PASSIVE STAGGERING BUT COOPERATIVE POOR. Breath Face FLUSHED \$ 8UNDTY CHEMICAL ODOIS □ Reddened Conjunctiva Speech Blindness: Tracking: Normal □ Bloodshot □ Watery None □LEye □REye XEqual □ Unequal S-NELED, SLOW & LOW Corrective Lens: None Pupil Size: KEqual Able to follow stimulus: XYes □ No Eyelids: □ Glasses □ Contacts, if so □ Hard □ Soft □ Unequal (explain) ×Normal □ Droopy Right Eye Vertical Nystagmus? XYes □ No One Leg Stand Pulse & Time HGN Left Eye 1. 100 / 2150 Lack of Smooth Pursuit Convergence Y=5 YES Right Eye Left Eye 2 108 / 2204 Max. Deviation YE5 ye s 1/26 100/3217 Angle of Onset MIMEDIATE TEST STOPPED Walk and Turn Test TEST STOPPED Romberg Balance Cannot Keep Balance -Approx. Approx. Starts Too Soon 1st Nine 2nd Nine Sways While Balancing Stops Walking 3 Uses Arms to Balance Misses Heel-Toe ALL ALL Steps Off Line □ Hopping Raises Arms IJJ 炉 Puts Foot Down Actual Steps Taken I SWAY CIRCULAR Internal Clock Describe Turn SWINGLED IN ONE Cannot Do Test (Explain) Type of Footwear ABRUPT MOTION. LOST 45 __Estimated At 30 Sec. N/A BALANCE TENNIS SHOES Room Light Direct Pupil Size Darkness Nasal Area ○ Right ∧ Left CLEAR Left Eye Draw lines from spot touched 4.0 3.5 6.0 Oral Cavity CHEMICAL Right Eye HTRAJISI CIO 2000 4.0 6.0 3.5 Hippus □ Yes Reaction To Light Rebound Dilation ΧNο □ Yes 🐧 No NORMAL (2 NO VISIBLE MARKS Temp <u>99.8</u> Blood Pressure 146 1 100 Muscle Tone: □ Normal □ Flaccid A Rigid Attach Photos Of Fresh Puncture Marks Comments: ARMS VERY RIGID at Medicine or Drug Have You Been Using? How Much? Time of Use? Where Were The Drugs Used? (Location) Sinzing. 2724 NG 20 م*نجرن<u>د ۵</u>* Time DRE Notified Date/Time of Arrest Eval Start Time Time Completed ブブンロ 2 JED0 2100 2120 ID No. 7654 Member Signature (Include Rank) Reviewed By: FARMAR # 10938 Opinion of Evaluator: □ Rule Out □ Alcohol PCP □ Stimulant □ Inhalant □ Medical □ Depressant □ Hallucinogen □ Narcotic Analgesic □ Cannabis

#### DRUG INFLUENCE EVALUATION

Page 2_of 2

LOG NO.

DRE: Sgt. Mark George

ARRESTEE: Robert H. Ross

- 1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR.
- INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS
- 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.
- 1. LOCATION: Examination of Robert H. Ross, took place in the DRE room, NYSP Tarrytown
- 2. WITNESS: Arresting Officer Trooper Alan D. Brown
- 3. BREATH TEST: Troper Brown administer a breath test to Ross at 2135 hours, the result was 0.00%
- 4. NOTIFICATION / INTERVIEW of ARRESTING OFFICER: Writer was contacted by radio at 2120 hrs. and advised to return to the station to conduct a DRE evaluation. Tpr. Brown informed me that he had observed Ross driving S/B in the median of the NYS Thruway, at approximately 10 mph. Brown stated that the subject appeared dazed and could not state where he was or where he had come from.
- 5. INITIAL OBSERVATIONS: Writer observed subject at 2140 hrs. He appeared dazed and disoriented, he had a fixed stare and responded very slowly (approx. 5 10 seconds delay) to all my questions and instructions.
- 6. MEDICAL PROBLEMS: None noted or stated
- 7. PSYCHOPHYSICAL TESTS: Romberg Balance: Subject swayed approximately 3" in a circular motion and estimated 45 seconds as 30 seconds. Walk and Turn: Subject started walking immediately, lost balance during the instructions, stepped off the line, stopped walking, repeatedly used his arms for balance, and missed heel to toe One Leg Stand: Subject unable to complete the test using either foot. Finger to Nose: Subject missed tip of his nose on each attempt and his arm movements were very rigid.
- 8. CLINICAL INDICATORS: Subject exhibited immediate onset of HGN, vertical nystagmus, and lack of convergence. Blood pressure, pulse and body temperature were above the normal range
- 9. SIGNS of INGESTION: There was a strong chemical odor on the subject's breath.
- 10. STATEMENTS: Subject stated that he did not use any drugs.
- 11. OPINION of EVALUATOR: In my opinion Robert H. Ross is under the influence of Phencyclidine, or an analog, and unable to operate a vehicle safely
- 12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a blood sample.
- 13. MISCELLANEOUS: Three (3) discolored filtered cigarettes in a "Kool" box were found in the subject's right shirt pocket, and were sent to the laboratory for analysis.

Drug Influence Evaluation Rolling Log Y- OZZZ □ Patal □ Property Mark beorge □ Injury estee's Name (Last, First, Arresting Officer (Name, ID No.) Mark George 8-10-1964 Late Examined/Time/Location Breath Results: 0.04 Refused Central Testing
No What have you eaten today? 5-2-2000 2300 1234 Miranda Warning Given: XYes - No When? Have you been drinking? How much? Time of last drink? Pizza 5 pm Beer 5 PM When did you last sleep? Are you sick or injured? Time now? How long? Are you diabetic or epileptic? Do you have any physical defects? 

Yes 

No Are you under the care of a doctor or deptist? Yes No No response Attitude Withdrawn, non responsive passive. Are you taking any medication or drugs? 

Yes *No ath Chemical Odor Speech slow, slurred, attimes did not respond □ Reddened Conjunctiva Blindness: ■ Equal © Unequal Normal □ Bloodshot □ Watery None □ L Eye □ R Eye Corrective Lens: Pupil Size; FEqual Able to follow stimulus: ₽ Yes □ No Eyelids: □ Glasses □ Contacts, if so □ Hard □ Soft □ Unequal (explain) Normal □ Droopy Vertical Nystagmus? Yes O No Pulse & Time HGN Left Eye Right Eye 1 120,2310 4es Yes Lack of Smooth Pursuit Convergence GIVER Left Eye Right Eye 2326 4es Max. Deviation Immediate Immediate 2338 Angle of Onset Subject fell and ten Walk and Turn Test Cannot Keep Balance WWW WW W W W W Starts Too Soon 1" Nine 2nd Nine □ Sways While Balancing Stops Walking Misses Heel-Toe Uses Arms to Balance ms locked. Steps Off Line □ Hopping Raises Arms Constant vi □ Puts Foot Down Actual Steps Taken Describe Turn Swive led abruptly Cannot Do Test (Explain)
Staggered to the 1eft. Type of Footwear Internal Clock 40 Estimated At 30 Sec. Loafers Room Light Direct Pupil Size ○ Right △ Left Left Eye 4.0 3.5 Draw lines from spot touched Right Eye Reaction To Light Rebound Dilation Hippus □ Yes □ Yes -No Normal Visible Mot 104 Temp 100.5 Blood Pressure_ Muscle Tone: □ Normal □ Flaccid ■ Rigid Attach Photos Of Fresh Puncture Marks Comments Arms & weck very rigid t Medicine or Drug Have You Been Using? Where Were The Drugs Used? (Location) How Much? Time of Use? No response No response lo response Time DRE Notified Determine of Arrest
May 2, 2000 2.
Member Signature (Include Rapk) ID No. 1176 Reviewed By 1 Rule Out **≯** PCP □ I**h**halant Opinion of Evaluator: □ Alcohol □ Stimulant □ Medical □ Cannabis □ Depressant □ Hallucinogen □ Narcotic Analgesic

#### DRUG INFLUENCE EVALUATION

Page 2 of 2

LOG NO.

DRE: Officer John Blea

ARRESTEE: D. Schroeder

- LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR.
- 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS
- 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.
- 1. LOCATION: Examination of D. Schroeder took place in the DRE room, Denver PD Headquarters
- 2. WITNESS: Arresting Officer Officer Mark George
- 3. BREATH TEST: Officer George administered a breath test to Schroeder at 2300 hours, the result was

0.04%. At this time subject admitted she had consumed some beer.

- 4. NOTIFICATION / INTERVIEW of ARRESTING OFFICER: Writer was contacted by radio at 2245 hrs. and advised to return to Headquarters to conduct a DRE evaluation. Officer George informed me that he had observed the subject fail to obey a stop sign. At the time of the stop Schroeder was smoking a cigarette which gave off a strong chemical odor. Additional examination of the cigarette indicated the possibility of some form of string in the middle.
- 5. INITIAL OBSERVATIONS: Writer observed subject sitting quietly in the DRE room, staring at the floor, and taking no notice of the activity around her. It was necessary to instruct the subject twice to raise her head before she complied.
- 6. MEDICAL PROBLEMS: None noted or stated
- 7. PSYCHOPHYSICAL TESTS: Ms. Schroeder was very slow in responding to all instructions during this portion of the examination. Romberg Balance: Subject swayed approximately 3" in a circular motion and estimated 40 seconds as 30 seconds. Walk and Turn: Subject lost balance during the instructions, took the wrong number of steps, turned abruptly, stepped off the line, and repeatedly used her arms for balance. On the return she never touched heel to toe and simply took 12 "normal" steps. Her legs seemed very stiff and rigid. One Leg Stand: Subject fell after only three (3) seconds. Finger to Nose: Subject missed tip of her nose on each attempt and on one attempt missed her nose entirely.
- 8. CLINICAL INDICATORS: Subject exhibited immediate onset of HGN, vertical nystagmus, and lack of convergence. Blood pressure, pulse and body temperature were above the normal range.
- 9. SIGNS of INGESTION: There was a strong chemical odor on the subject's breath.
- 10. STATEMENTS: Subject stated that she had drank "one (1) beer" She did not respond to the questions regarding drug use or questions concerning the cigarette.
- 11. OPINION of EVALUATOR: In my opinion D. Schroeder is under the influence of Alcohol and Phencyclidine, or an analog, and unable to operate a vehicle safely.
- 12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a blood sample.
- MISCELLANEOUS: The confiscated cigarette was sent to the laboratory for analysis.

Three Hours

# SESSION XVII NARCOTIC ANALGESICS

# SESSION XVII NARCOTIC ANALGESICS

Upon successfully completing this session, the participant will be able to:

- o Explain a brief history of the Narcotic Analgesic category of drugs.
- o Identify common drug names and terms associated with this category.
- o Identify common methods of administration for this category.
- o Explain the symptoms, observable signs and other effects associated with this category.
- o Explain the typical time parameters, i.e., onset and duration of effects, associated with this category.
- o State the clues that are likely to emerge when the DRE Evaluation is conducted.
- Explain the procedures for examining and determining the ages of injection sites.
- o Correctly answer the "topics for study" questions at the end of this section.

Content Segments		Learning Activities	
A.	Overview of the Category	0	Instructor Led Presentations
В.	Possible Effects	0	Review of Drug Evaluation and Classification Exemplars
C.	On-Set and Duration of Effects	0	Reading Assignments
D.	Overdose Signs and Symptoms	O	Video Presentations
E.	Expected Results of the Evaluation	o	Slide Presentations
F.	Injection Site Examination		
G.	Expected Location of Injection Marks		
H.	Conclusion		





Analgesics)

NARCOTIC ANALGESICS

Total Lesson Time: Approximately 180 Minutes

Session title on wallchart.

Briefly review the objectives, content and activities of this session.

Point out that this category sometimes is called "The Opioids"; the drugs it contains either are found in Opium, or derive chemically from Opium, or produce effects similar to those of the Opium Derivatives.

The term "Opioid," however, most correctly refers to the synthetic subcategory of Narcotic Analgesics.



25 Minutes



XVII-1 ("Narcotic Analgesics Defined")

A. Overview of the Category

- 1. Narcotic Analgesic defined
  - a. A medical term, not a legal or police term.
  - b. An "Analgesic" is a drug that relieves pain. It differs from an anesthetic, in that it lowers one's perception of pain, rather than stopping nerve transmission.
  - c. Non-Narcotic Analgesics, such as Aspirin, Tylenol, and Motrin, relieve pain, but do <u>NOT</u> produce narcosis, which means numbness or sedation.

Clarification: Non-Narcotic Analgesics relieve pain, but do not alter mood. Therefore, they, in small amounts, are not psychoactive, and are not abused for their mind or mood altering actions.



XVII-2 ("Types of Narcotic Analgesics")

- d. A Narcotic is a drug derived from Opium, or produced synthetically that relieves pain, but also induces euphoria, alters mood, and produces sedation.
- 2. There are two subcategories of Narcotic Analgesics.
  - a. The Opiates: drugs that either contain or are derived from Opium.
    - (1) the natural alkaloids of Opium

(2) Opium derivatives.

Point out that a "natural alkaloid" is a substance that is found in another substance, and that can be isolated from it. Morphine, for example, is a natural alkaloid of Opium. Codeine is another example of a natural alkaloid.

The term "main ingredient" can be used as a synonym for "alkaloid."

Opium derivatives are obtained by chemically treating the Opium alkaloid. Opium Derivatives are therefore derived from Opium.

An analogy to help students understand the difference between an alkaloid and a derivative would be to compare opium to wheat. The "alkaloid" of the wheat would be whole wheat flour--a derivative of the wheat would be white flour (wheat flour which has been chemically treated)

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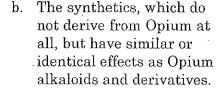
(Character-

istics common

to all Narcotic Analgesics)

#### Lesson Plan

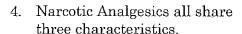
#### Instructor Notes



Point out that the synthetic Narcotic Analgesics are produced from a variety of nonopiate substances. Again, these are sometimes called "Opioids".

3. The natural alkaloids and the Opium Derivatives all come from Opium, which is sap from the seed pods of a particular type of poppy.

NOTE: The Opium poppy, or papaver somniferum (somniferum, Latin for the "carrier of sleep").



a. They will relieve pain.

Clarification: They produce analgesia.

- b. They will produce withdrawal signs and symptoms when the user is physically dependent, and drug use is stopped.
- Clarification: Physical dependence results from "chronic administration." This means that the drug has been taken at fairly regular intervals for a period of time.
- c. They will suppress the withdrawal signs and symptoms of chronic morphine administration.

Morphine is typically used as the standard for comparison with other Narcotic Analgesics.

<u>Clarification</u>: This means that the various Narcotic Analgesics can be substituted for each other to relieve withdrawal symptoms.



XVII-4 ("Commonly Abused Opiates")

- 5. Some commonly abused <u>Opiates</u>.
  - a. Powdered Opium (also known as smoking Opium)
    - o a simple refinement of raw Opium.

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#### **Instructor Notes**

- o used medically to treat diarrhea (administered orally)
- o remains popular as a drug of abuse (smoked) among some Asian-American communities.
- b. Morphine, the principal natural alkaloid of Opium.
  - o Morphine was first isolated from Opium in 1805.
  - o used medically to suppress severe pain (e.g., with terminal cancer patients).
  - o highly addictive
  - o at one time, Morphine was the most commonly abused Narcotic Analgesic.
- c. <u>Codeine</u> is another natural alkaloid of Opium.
  - o first isolated in 1832.
  - o Codeine's pain killing ability is much weaker than Morphine's.
  - used medically to suppress coughing or minor pain.
  - o Codeine is definitely an addictive drug.

The development of more effective opiates and synthetics has virtually eliminated its use medically. In recent years, there have been little street use of Opium. It is important to realize, however, that drug use trends can and do change.

Instructor, FYI: Named after Morpheus, the Greek God of dreams.

Morphine was widely used during the Civil War. Morphine addiction was termed "Soldier's disease."

Its technical name is Methylmorphine.

<u>Clarification</u>: Narcotic Analgesic addicts often turn to Codeine when they cannot get more popular drugs.

#### **Instructor Notes**



- d. <u>Heroin</u> is the most commonly abused illicit Narcotic Analgesic.
  - o derived from Morphine in 1874.
  - o Heroin was first thought to be a non-addictive substitute for Morphine.
  - o it was approved for general use by the American Medical Association in 1906.
  - o by the 1920's it was evident that Heroin was much more addictive than Morphine.
  - o importation and manufacture of Heroin have been illegal in this country since 1925.
- e. <u>Dilaudid</u> is another derivative of Morphine
  - o first produced in 1923.
  - o sometimes called "drug store Heroin", since it is commercially available from medical and pharmaceutical sources.
  - Dilaudid has the same addictive liabilities as does Heroin or Morphine.

Point out that the generic, or technical name for heroin is "Diacetyl Morphine".

Write "Diacetyl Morphine" on the chalkboard or flipchart.

Heroin is a Schedule I drug, which means it has no legitimate medical uses in the United States.

Technical Name: Hydromorphone Hydrochloride.

Aides	Aides Lesson Plan		Instructor Notes	
		o used medically for short term relief of moderate to severe pain, and to suppress severe, persistent coughs.		
		o can be ingested via injection, orally or in suppositories.		
	f.	<u>Hycodan</u> is a derivative of Codeine.	Technical Name: Hydrocodone.	
		<ul><li>o used medically to treat coughs.</li><li>o sometimes abused by addicts who are unable</li></ul>	Note: Vicodin is a commonly prescribed pain reliever containing Hydrocodone and Acetaminophen.	
		to obtain Morphine or Heroin.		
	g.	Percodan is another derivative of Codeine.	Technical Name: Oxycodone.	
		o Percodan is one of the most commonly prescribed Narcotic Analgesics.	It is also produced the under the brand name of "Percocet which is Percodan combined with Acetaminophen, such as Tylenol.	
		o it is somewhat less addictive than Morphine, but more than Codeine.		
		o another medicine, <u>Percobarb</u> , is a  combination of Percodan  and Barbiturate.	i.e., <u>Percobarb</u> combines a CNS Depressant with a Narcotic Analgesic.	
	h.	Metopon derives from Thebaine, another alkaloid of Opium.		
4		o Metopon is chemically similar to Morphine.		

#### **Instructor Notes**



XVII-5 ("Common Synthetic Opiates")

- o used to relieve chronic pain such as terminal cancer.
- 7. Some common <u>Synthetic</u> Opiates.
  - a. <u>Demerol</u> was first produced in 1939.
    - o Demerol is one of the most widely used Synthetic Opiates for relief of pain and for sedation.
    - o it is also the Narcotic Analgesic that is most frequently abused by medical personnel who become addicts.
    - o Demerol is widely used as an analgesic in childbirth.
    - o one medical advantage of Demerol is that it produces less respiratory depression than do other Narcotic Analgesics; thus, a fatal overdose is less likely with Demerol.
    - o Medical literature sometimes indicates that Demerol does not cause pupillary constriction. Enforcement experience indicates to the contrary.

Technical Name: Meperidine.

Point out that pupillary constriction ordinarily is one of the most reliable indicators of a Narcotic Analgesic.

- b. Methadone was developed in Germany during World War II and first marketed in America in 1947.
  - o Methadone's effects are similar to Morphine's, although they develop more slowly and last longer than do Morphine's effects.
  - o Methadone's withdrawal symptoms are slower and milder than are Morphine's.
  - o used extensively in
    "maintenance programs"
    as a substitute for
    Heroin for addicts
    undergoing therapy and
    treatment.
  - o In theory, the daily dose of Methadone given to an Heroin addict allows the addict to function normally with no physical need for up to 24 hours.
  - Methadone is also used medically to relieve moderate to severe pain, and to suppress coughing.
- c. Numorphan is a powerful analgesic with the same addictive properties as Morphine.
  - o used medically for the relief of chronic pain.

Methadone was developed in Germany because of wartime shortages of Morphine. Some experts have stated that the brand name for Methadone, "Dolophine," was derived from Adolph Hitler.

Ask students: "What is one of the most common medical uses of Methadone in this country?"

Remind students that one characteristic shared by all Narcotic Analgesics is that they suppress withdrawal symptoms of chronic Morphine administration.

Methadone's primary advantages are: it cannot be injected, and it has a much longer duration of effects than Heroin.

Aides	Lesson Plan	Instructor Notes	
	o sold in ampules (injection) and in suppositories.		
	o previously (pre-1972) it was sold in tablets, and was a favorite substitute for Heroin among addicts; addicts now generally prefer Dilaudid as an Heroin substitute.		
	d. The <u>Fentanyls</u> include several hundred "designer drug" analogs of Morphine.	"Sublimaze" is a brand name for Fentanyl. It is a Schedule II drug. It is frequently found in overdose situations. For	
	o first developed in 1965 as an intravenous anesthetic.	example, "Tango and Cash" and "Goodfellas", which contained Fentanyl, were sold	
	o legally produced as a pain killer.	in New York City in 1990 as Heroin. Many fatal overdoses occurred as a result.	
	o principal abused analog is "Three-Methyl Fentanyl".		
	e. Three-Methyl Fentanyl is  very powerful: about 100  times as powerful as  Morphine.		
	o can be fatal in very small amounts.		
	o can be absorbed through the skin: <u>very</u> dangerous to handle.	Remind about officer safety when handling any containers found on the suspect.	
	f. <u>MPPP</u> is an illegally manufactured analog of Demerol.		

MPPP is a powerfully addictive synthetic Narcotic Analgesic.

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#### **Instructor Notes**

- o in the course of producing MPPP, it often becomes contaminated with MPTP, a chemical producing paralysis similar to Parkinson's Disease.
- g. <u>Darvon</u> is a synthetic Narcotic of relatively low analgesic potency and relatively low addiction liability.
- 7. Methods of administration of Narcotic Analgesics vary from one drug to another.
  - a. Some are commonly taken orally.
  - b. Some are often administered in suppositories.
  - c. Some are smoked.
  - d. Some are snorted. (taken intranasally)
  - e. Heroin, and some others, usually are taken by injection.
  - f. Medically, some Narcotic Analgesics may be administered transdermally or through the skin.
- B. Possible Effects

Instructor, FYI: Parkinson's disease is a progressive neurological disorder characterized by resting tremors, shuffling gait, and muscle weakness.

Technical name: Propoxyphene.

Users have stated that the fear of contracting diseases, such as AIDS, from shared needles, has prompted them to either snort or smoke Heroin.

<u>If available</u>, show 35 mm slides of Heroin injection paraphernalia.

Solicit students' comments and questions concerning this overview of Narcotic Analgesics.



5 Minutes

#### **Instructor Notes**



XVII-6 ("Concept of Tolerance")

- As with nearly all the drugs of abuse, the effects produced by heroin or other Narcotic Analgesics depend on the tolerance that the user has developed for the drug.
  - a. People develop tolerance for Narcotic Analgesics fairly rapidly.
  - b. "Tolerance" means that the same dose of the drug will produce diminishing effects, or conversely that a steadily larger dose is needed to produce the same effects.

- c. A Narcotic Analgesic user who has developed tolerance and who is using his or her "normal" dose of the drug may exhibit little or no evidence of intellectual or physical impairment.
- d. Impairment is more evident with new users, and with tolerant users who exceed their "normal" doses.
- 2. Observable effects of Heroin and other Narcotic Analgesics.
  - a. Sedation "On the Nod"
    - o the condition known as "on the nod" is an apparently semiconscious state, resembling sleep.

Emphasize: Habitual users of drugs may develop tolerance to the drug. As a result, they may exhibit relatively little evidence of impairment on the psychophysical tests. Even tolerant drug users, when impaired, usually exhibit clinical evidence. (i.e. in the vital signs and eye signs - such as HGN)

Clarification: the tolerant addict who has injected his or her "normal" dose of Heroin may appear to be much less impaired than an inexperienced user who had taken the same dose.

<u>Point out</u> that "on the nod" occurs most often with new users or with users exceeding normal doses.



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#### **Instructor Notes**

- o the user's eyelids become very droopy.
- o their head will slump forward until the chin rests on the chest.
- o in this condition, the user usually can be aroused easily and will be sufficiently alert to respond to questions.
- b. Other effects.
  - o slowed reflexes
  - o slow and raspy speech
  - o slow, deliberate movements
  - o inability to concentrate
  - o slowed breathing
  - o skin cool to the touch
  - o possible vomiting
  - o itching of the face, arms or body
- C. Onset and Duration of Effects
  - 1. The psychological effects of Heroin begin virtually immediately after the addict has injected.

Remind students that the technical term for "droopy eyelids" is <u>Ptosis</u>.

<u>NOTE</u>: These effects may be dose-related, and most often occur with non-tolerant users.

Instructor, FYI: Technical terms are Hypopnea or Bradypnea.

Solicit students' comments and questions concerning possible effects of Narcotic Analgesics.



XVII-8A ("On-set and Duration of Effects of

Heroin")

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Aides
XVII-8B
XVII-8C

#### **Instructor Notes**

- A feeling of pleasure or euphoria.
- b. Relief from the symptoms of withdrawal.
- c. Relief from pain.
- 2. The observable signs will usually become evident within 5-30 minutes after the user has injected.
  - a. "On the nod"
  - b. Poor motor coordination
  - c. Depressed reflexes
  - d. Slowed breathing
- 3. The effects will usually be observable for up to 4-6 hours.
- 4. As the drug wears off, withdrawal signs and symptoms start to develop until the user injects again.

- a. Withdrawal <u>symptoms</u> usually begin to be felt by the addict within 4-6 hours following injection.
  - o chills
  - o aches of the muscles & joints
  - o nausea
  - o insomnia

Point out that the intensity of the euphoria will depend on a number of factors, one of which is the addict's tolerance. A heavily addicted user who is beginning withdrawal symptoms may experience only mild euphoria.

Remind students that the physical effects may not be observed at all, if the addict is tolerant and has injected a "normal" or "maintenance" dose.

<u>Point out</u> that the development of withdrawal symptoms implies that the Heroin has worn off, so that the addict is no longer under the influence.

As with nearly all drugs, the withdrawal signs and symptoms are essentially the opposite of the "high" or intoxicated state.

The early stages of withdrawal constitute the "downside effects of Narcotic Analgesics. DRE's should never state that an individual is "under the influence of the downside" of a drug.





- b. Withdrawal <u>signs</u> start to become observable 8-12 hours following injection.
  - o sweating
  - o goose bumps (Piloerection) on the skin
  - o yawning
  - o tearing
  - o runny nose
  - o vomiting
- 5. Withdrawal signs and symptoms closely resemble those of Influenza or the common cold.
  - a. These symptoms begin to intensify from 14-24 hours after injection, and may be accompanied by goose bumps (piloerection), slight tremors, loss of appetite and dilation of the pupils.
  - b. Approximately 24-36 hours after injection, the addict experiences insomnia, vomiting, diarrhea, weakness, depression and hot and cold flashes.
  - c. Withdrawal symptoms and signs generally reach their peak 2-3 days after injection:

Point out that "sweating" usually is the first withdrawal sign to appear.

"Piloerection" means "hair standing up".

Point out that yawning, tearing, runny nose and vomiting usually appear only after marked withdrawal of many hours.

Point out that "withdrawal" signs of Narcotic Analgesics are essentially the opposite of their "under the influence" signs.



XVII-9C



XVII-9D



Aides	Lesson Plan	Instructor Notes
	o muscular and abdominal cramps	
	o severe tremors and twitching	Point out that the involuntary tremors and twitching of the legs give rise to the expression
	o elevated temperature	"kicking the habit".
	d. The user at this point is nauseated, gags, vomits and may lose 10-15 pounds within 24 hours.	d
	e. The withdrawal syndrome continues to decrease in intensity over time, and is usually greatly reduced by the fifth day, disappearing in one week to 10 days.	
	f. A common misconception regarding withdrawal from Narcotic Analgesics is that they may be fatal. In reality, however, although Narcotic withdrawal is extremely uncomfortable, it rarely, if ever, proves fatal.	Solicit students' comments and questions concerning onset and duration of the effects of Narcotic Analgesics.
	D. Overdose Signs and Symptoms	
5 Minutes	Narcotic Analgesics depress respiration.	Point out that this is an effect that Narcotic Analgesics have in common with CNS
	a. In overdoses, the user's breathing will become slow and shallow.	Depressants.
	b. Death can occur from severe respiratory depression.	
	c. The danger of death is heightened by the fact that the addict may not know the strength of the drug he or she is taking.	Clarification: the percentage of pure Heroin in the sample the addict uses may be much higher than what the addict expects and is used to.

- 2. Other signs and symptoms of an overdose of a Narcotic Analgesic include clammy skin, convulsions and coma, blue lips and pale or blue body, extremely constricted pupils (unless there is brain damage, in which pupils may be dilated), recent needle marks, or perhaps a needle still in the user's arm.
- 3. Narcotic Analgesic overdoses are sometimes treated by the administration of a Narcotic antagonist such as Narcan. A Narcotic antagonist works at neuron receptor sites, blocking or counteracting the effects of Narcotic Analgesics. In effect, these substances precipitate withdrawal. The short duration of effects produced by Narcotic antagonists, however, require continued medical monitoring of the user.
- E. Expected Results of the Evaluation
  - Observable evidence of impairment.
    - a. Standardized Field Sobriety Tests.
      - o Neither Horizontal Gaze Nystagmus nor Vertical Gaze Nystagmus will be present.

E.g., "Tango and Cash" and "Goodfellas" were sold on the street as high grade Heroin. Rather, these contained the much more potent Fentanyl, resulting in many fatalities.

Point out that a person suffering from Narcotic Analgesic overdose may appear to be in shock.

Solicit students' comments and questions concerning signs and symptoms of an overdose of Narcotic Analgesics.

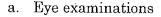


But remind students that Nystagmus could be present if the user has taken Heroin and PCP, or alcohol or some other CNS Depressant, or an Inhalant.

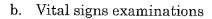
Aides	Lesson Plan	Instructor Notes
		Point out that, if the user has injected enough Narcotic Analgesic to exceed his or her level of tolerance, his or her performance of the Standardized Field Sobriety Tests will be uncoordinated and "rubberlegged", similar to that caused by CNS Depressants.
	o Performance on Walk and Turn and One Leg Stand will be impaired, and will reflect the slow and deliberate movements caused by this category of drugs.	
	o Performance on Romberg and Finger to Nose will also be impaired. Generally, the user will appear drowsy, possibly "on the nod," and exhibit slow and deliberate movements.	
XVII-10B ("General Indicators")	b. General indicators  o "Track marks" o "On the nod" o Droopy eyelids o Slowed reflexes o Slow, low, raspy speech o Facial itching o Dry mouth o Euphoria o Nausea o Pupils visibly and obviously constricted	If available, show 35 mm slides of typical addicts' "track" marks.  Caused by the release of Histamines.
	2. Evidence associated with the physiologic examinations.	



XVII-10C (Eye Examinations)



- Eyes will not exhibit a Lack of Convergence.
- Pupil size generally will be constricted (below 3.0 mm in diameter)
- Pupils reaction to light will be little or none visible.
- If the effects of the Narcotic Analgesic are wearing off, hippus may be evident.



- Blood pressure will be down.
- Pulse will be down.
- Body temperature will be down.
- Muscle tone will be normal or flaccid.

Summary

Point out that constricted pupils are one of the most reliable indicators of a Narcotic Analgesic. The technical term for "constricted pupils" is "Miosis."

NOTE: "Hippus" means pulsating pupils, i.e., alternately expanding and contracting in diameter.

Remind students that these cardiovascular indicators may not be present if the suspect is a tolerant user who has taken

a "normal" dose of the drug.



XVII-10D

Examinations")

("Vital Signs

XVII-11 ("Narcotic Analgesics Symptomatology Chart")

### Demonstrations

Video tape demonstrations.

Show video tape of suspect(s) under the influence of Narcotic Analgesics.



Aides	Lesson Plan	Instructor Notes
	b. Drug Evaluation and Classification exemplars demonstrations.	Relate behavior/ observations to the symptomatology chart.  Refer students to the exemplars found at the end of Section XVII of their student manuals.  Solicit students' comments or
		questions concerning Expected Results of the Evaluation.
	F. Injection Site Examination	
30 Minutes	<ol> <li>Examination of suspect's injection sites can give many clues to their drug habits.</li> </ol>	The slang term for an injection site is a "mark".
	a. Many drugs can be injected.	The presence of injection sites doesn't ensure the suspect is
	<ul> <li>Injection sites are a sign of drug use which may or may not be recent.</li> </ul>	under the influence of drugs.  Examination of ingestion sites
,	c. May be evidence of habitual use.	is just one of the twelve steps in the evaluation.
	2. The trauma to the skin, muscles and the blood is the basic concept of injection sites.	
	3. Drugs and medication are injected into the body in three ways.	
•	a. Legal injections are usually	Abbreviated as I/M.
	Intramuscular.	Medicine or medical procedures, such as tuberculin skin test, injection of insulin is injected Intramuscular.
}	b. Subcutaneous, means just under the skin.	Commonly referred to as "skin popping".

- c. For medically drawing of blood or emergency medical procedures, the injection is made into a blood vessel. Veins are usually used. Arteries are deep, thus not lending themselves to injection.
- 4. The primary instrument for injection is the hypodermic syringe.
  - a. It consists of a hollow needle, a tube and a plunger.
  - b. Needles vary in size, with the primary variance being the inside diameter of the needle or the gauge.
  - c. Most medical procedures utilize a small gauge needle, which means a large needle inner diameter. (usually 16 22 gauge).
  - d. Illegal injections are usually made with a large gauge needle, which is a small inner diameter. (usually about 26 gauge).
- 5. The user's equipment is commonly referred to as a "hype kit" or "works".
  - a. The kit contains a "cooker" which is any device such as a bottle cap, a metal spoon or etc., that is used to heat the drug with water to form an injectable solution.

Insulin is never injected into a blood vessel, because the person would go into a comma.

Abbreviated as I/V.

IMPORTANT RULE: the larger the gauge, the smaller the inside diameter of the needle.

A 26 gauge needle is used by a diabetic.

The hypodermic marks are smaller and are therefore, less noticeable making it more difficult for the DRE to see them.

Aides	Lesson Plan	Instructor Notes
	b. A handle to hold the "cooker" over the flame.	
	c. Matches, lighters (primarily disposable, adjustable flame types) used to heat the substance in the "cooker".	
	d. A tourniquet, which can be a rubber tubing, a tie, belt, etc. It is tied around the arm, above the injection site, to cause the vein to bulge or rise, thus making it easier to inject.	
	e. "Cottons" are the cotton balls or cigarette filters used to "purify" the drug. The user places the "cottons" into their cooker and draws the drug up through the cottons.	The cottons are saved for later use since they contain some of the drug.
·	6. As an expert, you may be asked in court to describe the difference between a legal and an illegal injection site.	There may be multiple injections, if the technician is unable to find a vein during the first try.
	a. The legal mark is usually intramuscular. Some exceptions would be in an emergency, blood donation or lab tests.	
-	b. Usually there will be only one mark and it will be larger than the typical illegal injection.	
İ	c. Legal injections are made with new, sterile needles.	
	d. The illegal mark is usually	Abbreviated as O/V.

over a vein.

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### **Instructor Notes**

- e. There will usually be multiple marks in various stages of healing. It takes approximately two weeks for a "mark" to totally heal.
- f. Users frequently use the same needle over and over again. Thus making it become dull or barbed.
- g. Since the used needles make it more difficult to pierce the skin and vein, the injections sites may be jagged.
- Use of old, dirty and shared needles cause the spread of infections and diseases such as AIDS.
- 7. Users may frequently use the same spot to inject, as an attempt to reduce their likelihood of detection.
  - a. The veins may become hard and thick from continuous injections and makes them difficult to find.
  - b. After about 10 to 20 injections, a large sore forms causing the site to enlarge and bruise. Upon close examination, the site reveals there are numerous puncture wounds in the same area, overlapping each other.

For example, the Heroin addict will inject approximately four to six times each day (every four to six hours). Therefore, they will inject approximately 2,000 times in one year.

Frequently the needles are carried in pockets or socks and the rubbing against clothing causes them to be dull or barbed.

A barbed needle may tear the skin on the way in and on the way out.

ALWAYS WEAR RUBBER GLOVES PRIOR TO CONDUCTING THE EXAMINATION

The technical term is "Thrombosed".

Write Thrombosed on the chalk board or flip chart.

This is referred to as "tunnel" or "corn".

Write tunnel and corn on a flip chart.

The healing is greatly retarded.



- 8. Basic principles of puncture healing.
  - a. Any needle that punctures the skin leaves a scab. A scab is simply a crust formed by the drying of the discharge from the puncture.
  - b. These dried remains fill the gap caused by the puncture of the skin. As the fluids dry, they harden (clot and gel).
  - c. There are no exact timetables for wounds to heal, but there are some general guidelines.
  - d. Scabs develop within about 18 24 hours after a puncture.
  - e. After about 14 days a scab usually starts to peel or flake and then falls off. The skin under the scab is shrivelled and is lighter in color than the surrounding tissue.
- 9. There is no exact science to classifying the age of puncture wound. Some general guidelines are:
  - a. Fresh puncture wounds are defined as under 12 hours after injection and will be a red dot and have an oozing appearance.

Scab is the dried remains of blood, plasma (a cellular, colorless fluid part of the blood), lymph fluid (a thin fluid that bathes all the tissues of the body) and puss (a thick yellowish/greenish fluid that forms at an injection site).

Chronic disease, poor nutrition and etc. retard the puncture healing process.

A general rule: when the scab first forms, it is bright red. With age, the color gets darker and darker.

Users sometimes inject under a scab to hide multiple puncture wounds. This is referred to as "trap dooring".



Wounds)

- b. <u>Early</u> puncture wound is 12
  96 hours after injection. It will have a light scab, light bruise, reddened border and a crater appearance.
- c. Late puncture wound is 5 14 days and will have a dark scab, dark bruise and the crater will flatten.
- d. Healing puncture wound is over 14 days. The scab will be flaking and falling off with shriveled light colored skin underneath.
- 10. Other indicators of injection sites:
  - a. In an attempt to hide puncture wounds, users may inject into tattoos.
  - b. Tattooing also refers to dark carbon deposits that result from using a flame to "sterilize" a needle. Carbon deposits on the needle are then injected into the skin, causing a tattoo effect.
  - c. A "track" is a hardened part of a vein where numerous injections have been administered. The entire vein becomes scarred and hardened and with time may no longer be able to inject into. The area becomes silvery-blue in color and raised. This is referred to as "silver streaks".

Tattoos that are designed to hide puncture wounds are frequently colored and found on the inner arms.

AS A GENERAL RULE: one inch of tracks indicates that approximately 50 - 100 separate injections have been administered in this area.



#### 20 Minutes

- G. Expected Location of Injection Marks
  - 1. Prior to conducting the injection site examination, always remember to wear gloves.
  - 2. Injection sites may be located anywhere on the suspects' body.
    - a. The arms are most frequently used because the veins here are large and easily accessible.
    - b. The ankles are frequently used because the marks can be easily covered with socks.
    - c. The user may even use their neck because the marks can be hidden by hair or makeup.
    - d. They will basically use any part of their body where there is a vein.
  - 3. Conduct a thorough, slow, methodical examination of the suspect's arms beginning with the left.
    - Using a magnifying light, examine the inner arm as it is extended with the palm facing you.
    - b. Beginning at the bicep slowly examine the arm.Document the findings of your examine.

An ideal light is a 10 power light.

- c. Ask the suspect to contract the arm, so that he or she grasps their shoulder.

  Beginning at the wrist, slowly examine the arm to the elbow and document the results.
- d. Next examine the outer arm as it is extended palm facing downward. Start the examination at the shoulder moving to the wrist.
- e. Subject should extend and spread their fingers as the examination is conducted on the hands. Examine both sides of the hands, with particular attention to the areas between the fingers, under watch bands and rings.
- 4. Conduct the entire procedure for the right side.
- 5. Ankles are the next most common injection area.
  - a. Subject should be instructed to remove their shoes and socks to allow the DRE to examine them for puncture wounds.
  - b. The most common area is on the back of the foot.
- On a case by case basis, the DRE may need to examine other parts of the body for marks.

This forces the individual's veins to protrude.

Suspects sometimes hide hypodermic needles in their socks, shoes and the heel compartments of their shoes.

Aides		Lesson Plan	Instructor Notes			
		a. ALWAYS follow your Agencies rules, policies and procedures and laws regarding invasive type searches.				
	Н.	Conclusion				
15 Minutes	1.	The injection site examination may reveal evidence of recent use.				
	2.	The presence of marks however, doesn't mean drug influence or impairment at the time of the evaluation.				
	3.	A slow methodical examination, using a magnifying light, is required to obtain evidence.				
	4.	Conducting an injection mark examination is a skill. As with all skills, such as taking blood pressure, competency improves with practice.	Solicit students' comments and questions concerning the injection site examination.			

### **Session XVII**

### **Narcotic Analgesics**



### **Narcotic Analgesics**

Upon successfully completing this session, the participant will be able to:

- Explain a brief history of the Narcotic Analgesic category of drugs
- Identify common drug names and terms associated with this category
- Identify common methods of administration for this category
- Explain the symptoms, observable signs and other effects associated with this category

Drug Evaluation & Classification Training

XVII-OA

### Narcotic Analgesics

(continued)

- Explain the typical time parameters, i.e., onset and duration of effects, associated with this category
- State the clues that are likely to emerge when the DRE process is conducted
- Explain the procedures for examining and determining the ages of injection sites
- Correctly answer the "topics for study" questions at the end of this section

Drug Evaluation & Classification Training

XVII-Q8

### Narcotic Analgesic:

An "Analgesic" is a drug that relieves pain. It differs from an anesthetic, in that it lowers one's perception of pain, rather than stopping nerve transmission.

Drug Evaluation & Classification Training

XVII-1

### **Types of Narcotic Analgesics**



The opiates





· The synthetics



Drug Evaluation & Classification Training

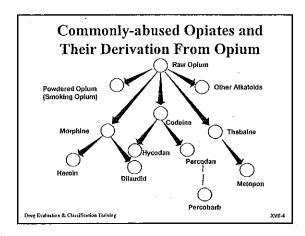
CVII-2

## Three Characteristics Common to All Narcotic Analgesics

- 1. Relieve pain
- 2. Produce withdrawal signs and symptoms
- 3. Suppress the signs and symptoms of chronic morphine withdrawal

Drug Evaluation & Classification Training

XVII-3



### **Common Synthetic Opiates**

- Demerol
- Methadone
- Numorphan
- Fentanyl
- · Three-Meth Fentanyl
- MPPP
- Darvon

Drug Evaluation & Classification Training



XVI]-5

## The Concept of Tolerance for a Drug

- 1. The same dose of the drug will produce diminishing effects
- 2. A steadily larger dose is needed to produce the same effect

Drug Evaluation & Classification Training

XVII-6

### "On the Nod"

- · Semi-conscious
- · Droopy eyelids (ptosis)
- Head slumped forward, chin on chest
- · Easily awakened
- · Alert to questions



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Drug Evaluation & Classification Training

## On-set and Duration of Heroin's Effects

- Immediate
  - Pleasure or euphoria
  - Relief from withdrawal
  - Relief from pain





Drug Evaluation & Classification Training

XVII-8A

### On-set and Duration of Heroin's Effects (continued)

- 5-30 minutes: Onset of physical effects
  - "On the nod"
  - Poor motor coordination
  - Depressed reflexes
  - Slowed breathing



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Drug Evaluation & Classification Training

## On-set and Duration of Heroin's Effects

(continued)

 Physical effects usually are observable for up to 4-6 hours







Drug Evaluation & Classification Training

XVII-80

## Signs and Symptoms of Withdrawal From Heroin

Symptoms begin: 4-6 hours following injection

- Chills
- Aches
- Nausea
- · Insomnia

Drug Evaluation & Classification Training

XVII-9

## Signs and Symptoms of Withdrawal From Heroin

(continued)

Signs appear: 8-12 hours following injection

- Sweating
- · Goose bumps
- · Yawning
- · Tearing
- · Runny nose
- Vomiting

Drug Evaluation & Classification Training

XVII-9E

## Signs and Symptoms of Withdrawal From Heroin

(continued)

Signs and symptoms intensify: 14-24 hours after injection

- · Similar to influenza or the common cold
- Goosebumps
- Slight tremors
- · Loss of appetite
- Dilation of pupils
   Final pupils
   Classification Training

vial.ar

## Signs and Symptoms of Withdrawal From Heroin

(continued)

Situation worsening: 24-36 hours after injection

- Insomnia
- Vomiting
- Diarrhea
- Weakness
- Depression
- Hot and cold flashes

Drug Evaluation & Classification Training

XVII-9D

# Signs and Symptoms of Withdrawal From Heroin

(continued)

Reaching the peak: 2-3 days after injection

- · Muscular and abdominal cramps
- · Severe tremors and twitching
- · Elevated temperature
- · Sharp loss of weight

Drug Evaluation & Classification Trainin

XVII-9E

## **Evaluation of Suspects Under the Influence of Narcotic Analgesics**

#### SFST Evidence:

- · HGN or Vertical Gaze Nystagmus none
- Performance on Walk and Turn and One Leg Stand will be impaired and will reflect the slow and deliberate movements
- Performance on Romberg and Finger to Nose will also be impaired
- Generally, the user will appear drowsy, possibly "on the nod"

Drug Evaluation & Classification Training

XVII-10A

## **Evaluation of Suspects Under the Influence of Narcotic Analgesics**

#### General Indicators:

- "Track marks"
- · Facial itching
- · On the nod
- · Dry mouth
- Droopy eyelids
- Euphoria
- · Slowed reflexes
- Nausea
- Slow, low, raspy speech
- Pupils visibly and obviously constricted (miosis)

Drug Evaluation & Classification Training

YVIL+AB

## **Evaluation of Suspects Under the Influence of Narcotic Analgesics**

#### Eye Examinations:

- · Lack of convergence none
- · Constricted pupils (miosis)
- · Reaction to light little or none visible
- As the effects of the drug wear off, hippus (pulsating pupils) may be evident

Drug Evaluation & Classification Training

XVII-10C

## **Evaluation of Suspects Under the Influence of Narcotic Analgesics**

#### Vital Signs:

- · Blood pressure down
- · Pulse down
- · Body temperature down
- · Muscle tone normal or flaccid

Drug Evaluation & Classification Training

XVII-10D

### Narcotic Analgesic Symptomatology Chart

HGN	None
VGN	None
Lack of Convergence	None
Pupil Size	Constricted
Reaction to Light	Little or None Visible
Pulse Rate	Down
Blood Pressure	Down
Temperature	Down
Muscle Tone	Normal or Flaccid

Drug Evaluation & Classification Trainin

XVII-11

### Classifying the Age of Puncture Wounds

- Fresh Under 12 hours after injection; will be a red dot and have an oozing appearance
- Early 12-96 hours after injection; will have a light scab, light bruise, reddened border and a crater appearance
- Late 5-14 days after injection; will have a dark scab, dark bruise and the crater will flatten
- Healing Over 14 days after injection; scab will be flaking and falling off with shriveled light-colored skin underneath

Drug Evaluation & Classification Training

XVII-12



000579	Drug Influenc	e Evaluation		
Evaluator Dan Webb	DRE No 0323	Rolling Log No. 946		
Recorder/Witness Stan O'Dell	Crash: K None □ Fatal □ Injury			
estee's Name (Last, First, MI)	DOB Sex	Race Arresting Officer (N	······································	
laugha, Jerry T.	5-18-1980 M		1 47650 I	
Date Examined/Time/Location 8-15-2000 2120 3 - 2	Precinct PD 1	reath Results 0.00 Refused strument #/234	Chemical Test  Refused	VUrine ABlood
	have you eaten today? Whe	n? Have you been drinking?	How much?	Time of last drink?
	othing M/	4 Nothing	<u> </u>	N/A
MIGHT Today 3 hours			Are you diabetic or e □ Yes MNo	pileptic?
Do you take insulin?	Do you have any physical de	fects? □ Yes No	Are you under the ca a doctor or dentist?	re of □ Yes <b>™</b> No
Are you taking any medication or drugs? UYes **Ne	Attitud On One Catur	e but sleepy.	Coordination, Ver	4 100se
	Breath Normal	e but steep y.	Stumbling Face 1/	
Speech	Eyes:   Reddened Conju	nctiva Blindness:	Norma	racking:
how and raspy.	Normal □ Bloodshot □			Equal □ Unequal
	Size: #Equal equal (explain)	Able to follow s	timulus: #Yes□No I	Eyelids: /ev/ Droopy
Pulse & Time HGN	Left Eye Right Eye Ve	ertical Nystagmus? 🗆 Yes 🚜 No	One Leg Stand	u reachediols
1. 60 / 7/25 Lack of Smooth Pursuit	NO NO Co	onvergence	17 3050	conds
2. 64 / 2140 Max. Deviation	No No	Right Eye Left Eye	(1)	9
3. 60 / 2153 Angle of Onset	NONE NONE		_ \ <b>\</b>	(1)
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dellocar	a all all	Steps Off Line		
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Draw lines from spot touched	Right Eye		Oral Cavity	Hear
N S S	77		-5	/ · · / ·
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		GHT ARM		+issue
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(4) X = 1	<u>\</u>		ale :	
	6	/skac		
(5)	~	Stare		
Blood Pressure //O / 64 Temp	98.0.	Puncture wound	, —	
Muscle Tone: □ Normal		with red dot Attach Photos Of Free	sh Puncture Marks	
: Medicine or Drug Have You Been Using?	How Much? Time of Use?			
1 -	ly question	s No a	rswer	
Date/Time of Arrest dug 15 2000/2100	DRE Notified 2/10	Eval Start Time 2/20	Time Completed	2210
Member Signature (Mank) Well 5		Reviewed By:	Floegel	
	I Alcohol □ Stimulan I Depressant □ Hallucino		□ Inhelant □ Cannabis	

		DRUG INFLUENCE EVALUATIO	ON Page <u>2</u> of <u>2</u>	
L	OG NO.	DRE: Sergeant Dan Webb	ARRESTEE: Jerry T. Vaughn	·· ··
5.	INITIAL OBSERVATI	ESS 3. BREATH TEST 4. NOTIFICATION ONS 6. MEDICAL PROBLEMS 7. PSYCH N 10. SUSPECTS STATEMENTS 11. OPIN	N / INTERVIEW ARRESTING OFCR. HOPHYSICAL 8. CLINICAL INDICATORS NION 12. TOXICOLOGY SAMPLE 13. MISC.	(
1.	LOCATION: Exami	ination of Jerry T. Vaughn, took place in	the DRE room, 3rd Pct.	
2.	WITNESS: Arresting	g Officer - Trooper Stanely R. O'Dell		
3.	BREATH TEST: To	rooper O'Dell administer a breath test to	Vaughn at 2100 hours, the result was 0.00%.	
4.	NOTIFICATION /	INTERVIEW of ARRESTING OFFIC	ER: Writer was contacted by radio and	
	advised to return to	the precinct to conduct a DRE evaluation	. Tpr O'Dell informed me that he had observed	
	the subject's vehicle	weaving through the traffic lanes. Subje	ect exhibited poor performance on the SFSTs, but th	ere
	no odor of an alcoho	olic beverage.		
5.	INITIAL OBSERVA	ATIONS: Writer observed subject sitting	g quietly in the DRE room. He appeared to be asleep	);
	eyes were closed, he	ad nodded forward, breathing was slow.	Subject responded to questions and became more a	lert
	as time passed. His	voice was low and raspy. He licked his li	ips repeatedly.	
6.	MEDICAL PROBL	EMS: None noted or stated		
7.	PSYCHOPHYSICA	L TESTS: Romberg Balance: Subject	swayed approximately 3" side to side and	
	estimated 50 seconds	s as 30 seconds. Walk and Turn: Subject	lost balance during the instructions, missed heel to	toe,
	stepped off the line,	and used his arms for balance. One Leg S	Stand: Subject put his foot down	
	swayed and used his	arms for balance. Finger to Nose: Subject	ect missed tip of his nose on each attempt.	
8.	CLINICAL INDICA	ATORS: Subject's blood pressure was be	elow the normal range. The pupils were constricted a	and
	showed little or no v	isible reaction to light. Subjects eyelids	were droopy.	
9.	SIGNS of INGESTI	ION: Subject had "track" type scars on bo	oth the left and right forearms, and a fresh oozing	
	puncture wound on	the back of the right hand.		
10.	STATEMENTS: Su	ubject denied using any medicine or drug	s and refused to answer any questions regarding the	e
	puncture wound on	the back of his right hand.		
11.	OPINION of EVAL	LUATOR: In my opinion Jerry T. Vaug	ghn is under the influence of a Narcotic Analgesic	
	and unable to operat	e a vehicle safely		
12.	TOXICOLOGICA	L SAMPLE: Subject agreed to provide b	poth a urine and a blood sample.	
13.	MISCELLANEOU	S:		
			,	
	• • • • • • • • • • • • • • • • • • • •			
			· · · · · · · · · · · · · · · · · · ·	

	000581		ıfluence E				
	Evaluator Toland, Steve	DRE No O.Z.	56 Ro	lling Log No	- 0100		
	Recorder/Witness Bradley. T.	Crash: YCS □ Fatal	□ None □ Injury	<b>∦</b> Pro	pperty	, , , , , , , , , , , , , , , , , , , ,	
(	restee's Name (Last, First, MI)	DOB 4-29-196	io Sex Ra		resting Officer (Na	me, ID No.)	a MPO
•	Date Examined/Time/Location Meso 11-1-2000 1615 Centra	POLL	Breath Instru	Results O.O		Chemical Test	□ Urine
	Miranda Warning Given: XYes □ No What	have you eaten tod	lay? When?	Have you	a been drinking?	How much?	Time of last drink?
		, , , , ,		□ Yes Æ	othing No	Are you diabetic o	r epileptic?
	Time now? When did you last sleep? How long?  Last A. a Far hour  Do you take insulin? Drys MNo	Do you have an	ny physical defects:	□ Yes 😿	No -	□ Yes ♣ No Are you under the	care of
	Are you taking any medication or drugs?   Yes  No					a doctor or dentist	? □Yes ♥No 2
	Re you taking any medication of drugs: 11 165 11 100	C	operational	-100		Sloppy-	Stumbling
	Speech		dened Conjunctive		Blindness:	"ace Norma	A ( Tracking:
	Slow and deliberate	<b>≵</b> Normal □ B	lloodshot □ Wate		X None □ L Ey		¥ Equal □ Unequal
		Size: ÆEqual equal (explain)			Able to follow st	imulus: #Yes □ No	Eyelids;  □ Normal *Droopy
	Pulse & Time HGN	<del></del>		Nystagmus?	□ Yes ≰No	One Leg Sta	
	Lack of Shiooth I disuit	<del> </del>	Converg	ence ht Eye	Left Eye		
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/	Walked	very slow	<i>''9</i> ')	Misses Hee	1 1		Jses Arms to Balance
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	Draw lines from spot touched	Lung		7.5		Oral Cavity	clear
	1,010,		Hippus □ Yes		d Dilation Read	ction To Light Z.7	He to
		''	Photo a		Puncture inds, 3 redd	none Visi	ouncture would with scabs
		71	AIGHT AA	XXX		O'TSEFT ARM W	nth scabs
		<u>\</u>				M. T.	
	3	26	(		ar tissue	_	
	Blood Pressure 100 / 60 Temp	97.0 .		-			
	Muscle Tone:   Normal A Flaccid Rigid Comments:   One of the sandneck very in the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of the comments of	relaxed.		Attach 1	Photos Of Fres	h Puncture Marl	<b>c</b> s
(	t Medicine or Drug Have You Been Using?		Time of Use?	Where V	Vere The Drugs Us		
`	1 None Refused 10 Unswer						
	Date/Time of Arrest //-/-Zooo/600  Member Signature (Int.) dde Rank)	DRE Notified	ID No.	Eval Sta	1615	Time Complet	ed 1710
	steve loland		ID No. 3529	Reviewe	u 59: M.	ryor	
- 1		Alcohol Depressant	□ Stimulant □ Hallucinogen	□ PCP	<b>ℓ</b> tic Analgesic	□ Inhalant □ Cannabis	

82		
	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>
LOG NO.	DRE: Sgt. Steve Toland	ARRESTEE: D. Bursten
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / ONS 6. MEDICAL PROBLEMS 7. PSYCHO 1 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS
1. LOCATION: Exami	nation of D. Bursten was conducted at the M	1esa PD holding facility
2. WITNESS: Arresting	g Officer - Officer T. Bradley #4779 MPD	
3. BREATH TEST: W	riter observed Officer Bradley administer a	breath test to Bursten, the result was 0.00%.
4. NOTIFICATION /	INTERVIEW of ARRESTING OFFICER	t: Writer was contacted by radio and advised
to return to the hold	ing facility to conduct a DRE evaluation. O	fficer Bradley informed me that the subject had been
involved in a car cra	sh at the intersection of Dobson and Main S	t. Subject exhibited poor performance on the
SFSTs, but there wa	as no odor of an alcoholic beverage.	
5. INITIAL OBSERV	ATIONS: Writer observed subject sitting qu	netly in the DRE room. He was scratching his
face and neck. His	eyelids were droopy and his voice was raspy	
6. MEDICAL PROBL	EMS: None noted or stated	
7. PSYCHOPHYSICA	AL TESTS: Romberg Balance; Subject sw	ayed approximately 3" in a circular motion and
estimated 50 second	s as 30 seconds. Walk and Turn: Subject sto	epped out of position during the instructions, stopped
walking and used hi	s arms for balance. One Leg Stand: Subject	put his foot down swayed and used his arms for
balance. Finger to N	Nose: Subject missed tip of his nose four tin	nes.
8. CLINICAL INDICA	ATORS: Subject's blood pressure, body ten	nperature and one pulse were all below the normal
range. The pupils w	ere constricted and showed litle or no visibl	e reaction to light. Subjects eyelids were droopy
9. SIGNS of INGEST	ION: Subject had three puncture wounds or	the right forearm and four puncture wounds with
scars on the left fore	arm	
10. STATEMENTS: S	ubject invoked his Miranda Rights.	
11. OPINION of EVA	LUATOR: In my opinion D. Bursten is un	der the influence of a Narcotic Analgesic
and unable to opera	te a vehicle safely	
12. TOXICOLOGICA	L SAMPLE: Subject agreed to provide a b	lood sample.
13. MISCELLANEOU	JS:	
		•

	Drug Influence Evaluation						
	Evaluator /ETZLAFF, 6. A.	126 DRE No XX	<b>ZZ-3</b> Rol	ling Log No.	-164		
	Recorder/Witness Parker, S.	Crash:	<b>D</b> None □ Injury	□ Prop			
	restee's Name (Last, First, MI) Shechan	DOB 05-16-19	366 M Rac		esting Officer (Na	me, ID No.) Znieski	×5484
		Parker Joil		Results: O. O.		Chemical Test  □ Refus	♥Urine □ Blood ed
	Miranda Warning Given:	What have you eaten t	oday? When?		been drinking?	How much?	Time of last drink?
	By: TETZCAFE, G. Time now?/ When did you last sleep? How By This morning 4  Do you take insulin? Thes	w long? Are you sick		□Yes 🕦	lo.	Are you diabetic o	or epileptic?
	Do you take insulin? Yes ** Idon't fake anything		any physical defects?			Are you under the a doctor or dentis	
	Are you taking any medication or drugs? □ Yes	Salca Salca	estic and	Sulle	7	Coordination Poor Staggering	
		Breath /	Cormal			Face Pale	100
	Speech Low numbled Raspy		eddened Conjunctiva Bloodshot   Water	у	Blindness: ■None □ L Ey		Tracking: Equal © Unequal
	Corrective Lens: □ None  □ Glasses □ Contacts, if so □ Hard □ Soft  Lemoved	Pupil Size: Æ Equal  Unequal (explain)			Able to follow st	imulus: 🏕 Yes □ No	Eyelids: Very  Normal Droopy
	Pulse & Time HGN	<del>  ,                                   </del>		Nystagmus? (	⊐ Yes □ No	One Leg St	and (22) (24)
	1. 60 / 2210 Lack of Smooth I	Pursuit Yes No	No Converg	ence it Eye	Left Eye		
	3. 58 / ZZ 30 Angle of Onset		NONE	<b>2</b> ) }	n Balanca	<b>Q</b>	R
	Approx. Approx.	teretereter S	2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	Cannot Kee Starts Too S	Soon I" Nine 2	L R	/ Sways While Balancing
	stopp of ter	red Counting	out tous	Stops Walkin Misses Heel-		W W	Uses Arms to Balance
				Steps Off Lin Raises Arms		11/1/	Hopping
	Internal Clock Describe 3	ournAs instru	Cann	Actual Steps	7	9 Type of Foo	Puts Foot Down
	SS_Estimated At 30 Sec. 60+	very slowly	/•	NA			gtips
	○ Right △ Left	,	Left Eve	-	<u> </u>	rect Nasal Area	Clear
	Draw lines from spot tou	ched	Right Eye /		2.0 /. 2.0 /·	S Oral Cavity	Clear
	N PIE	>, K)	Hippus □ Yes P No	Rebound		tion To Light He to non	e visible
	2		AIGHT AAI			LEFT ARM	R
	4)	$\times$		·	,	-	3 PUNCTURE RED WOUNDS RED
	55	6	·  (			PHOTO AREA	XX DOTS DOZING
Blood Pressure //O / 70 Temp 97.9 °						7,404	
}	Blood Pressure // / / / Temp // / 9 °  Muscle Tone: □ Normal □ Flaccid □ Rigid Comments:  Attach Photos Of Fresh Puncture Marks						
,	at Medicine or Drug Have You Been Using? 100 thing, do I look like	I do dope"	Time of Use?	Where W	ere The Drugs Us	sed? (Location)	lack"
ŀ	Date/Time of Arrest 2/30, 0	(et) 15 (5 (1 ) ( ) ( ) ( )	2140	Eval Star	rt Time	Time Comple	eted
}	Member Signature (Include Rank)	Well & I	ID No. 726	Reviewed	1By: -	Alexander -	-
l	Opinion of Evaluator: □ Rule Out □ Medical	# Alcohol  Depressant	□ Stimulant □ Hallucinogen	☐ PCP	ic Analgesic	□ Inhalant □ Cannabis	
1	□ Mcdrcal•	- Debieseane	~ randomoken	r∢αrcor	~ THIMPEOIL	- Cannanis	

D <del>584</del>		
	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>
LOG NO.	DRE: Sgt. Gary Tetzlaff	ARRESTEE: Thomas Sheehan
5. INITIAL OBSERV.	ITNESS 3. BREATH TEST 4. NOTIFICATION ATIONS 6. MEDICAL PROBLEMS 7. PSYCHOTION 10. SUSPECTS STATEMENTS 11. OPIN	OPHYSICAL 8. CLINICAL INDICATORS
1. LOCATION: Ex	amination of Roger J. Kurrus, took place in th	e DRE room, Jail Division, Parker Center
2. WITNESS: Arres	sting Officer - Officer D. Kvznieski	
3. BREATH TEST	: Writer observed Off. Kvznieski administer b	reath test to Sheehan, the result was 0.00%.
4. NOTIFICATION	N / INTERVIEW of ARRESTING OFFICE	R: At 2140 writer was contacted by Off. Kvznieski
who requested a	DRE evaluation. Sgt. Page informed me that k	e had observed subject driving westbound at 15 mph
on Longlook Lar	ne and the then failed to obey the stop sign at t	he intersection with Thunderhill Rd. Subject reacted
slowly and stopp	ed in the traffic lane approximately 800' past	the point where the emergency lights had been
activated. Subjec	ct appeared to be asleep and had his eyes close	d and his chin on his chest.
5. INITIAL OBSE	RVATIONS: Writer observed subject at 2200	hrs. He was wearing a three piece business suit with
no neck tie. Subjec	ct walked slowly, staggered and stumbled. He	swayed constantly while standing still, and his head
nodded forward re	peatedly. Subject spoke slowly in a low raspy	voice.
6. MEDICAL PRO	BLEMS: None noted or stated.	
7. РЅУСНОРНУЅІ	CAL TESTS: Romberg Balance: Subject sw	rayed approximately 2" front to back and estimated
55 seconds as 30	seconds. Walk and Turn: Subject lost his bala	nce during the instructions stepped off the line,
missed heel to too	e, and used his arms for balance. One Leg Sta	nd: Subject swayed, raised his arms, and put his foot
down. Finger to N	Nose: Subject missed tip of his nose on each at	tempt and used the wrong hand on the 3rd trial.
8. CLINICAL INDI	ICATORS: Subject's pupils were constricted,	systolic blood pressure was below the normal range.
His pulse was be	low the normal range on two (2) occasions. H	is eyelids were droopy.
9. SIGNS of INGES	STION: Subject's left arm had three (3) recen	puncture wounds and a one inch "track mark" scar.
10. STATEMENTS	: Subject stated that he did not used any drugs	Stated "Do I look like I do dope?" When asked
	ouncture wounds, subject said "Go have a hear	
11. OPINION of EV	ALUATOR: In my opinion Thomas Sheeha	n is under the influence of a Narcotic Analgesic,

12. TOXICOLOGICAL SAMPLE: Subject agreed to provide both a urine sample and a blood sample.

and unable to operate a vehicle safely.

13. MISCELLANEOUS: It appears that the subject is right handed.

Two Hours and Thirty Minutes

MID-COURSE REVIEW

### MID-COURSE REVIEW

This is an after-normal-class-hours session that students are free to attend or not, but are encouraged to attend. Its principal purpose is to help solidify the knowledge and skills they have begun to acquire, from the PRE-School and from the first four days of this school.

This session <u>must</u> be conducted in a highly interactive fashion. Don't simply present information or conduct demonstrations. Make the <u>students</u> do it. Ask questions, and call upon students to conduct the demonstrations that are required. Try to involve everybody, and convey your gratitude for the fact that they have attended the session.

### Content Segments

- A. Drugs, Drug Categories and the Drug Influence Evaluation
- B. Eyes and Vital Signs
- C. Physiology
- D. Questions and Answers

### **Learning Activities**

- o Instructor/Student Dialogues
- o Student-Led Demonstrations

#### **Instructor Notes**



MCR-1 (Review of Drugs, ...)

### MID-COURSE REVIEW

Drugs, Drug Categories and the A. **Drug Influence Evaluation** 

1. Define the word "drug".

2. Name the seven categories.

a. Name the six sub-categories of Depressants.

b. Name three sub-categories of CNS Stimulants.

Name two sub-categories of Narcotic Analgesic.

3. Identify the category for each of the listed drugs.

Desoxyn

Total Session Time: Approximately 150 Minutes

Any substance, which when taken into the human body can impair the ability to operate a vehicle safely.

CNS Depressants, CNS Stimulants, Hallucinogens, Phencyclidine, Narcotic Analgesics, Inhalants & Cannabis.

Barbiturates, Non-Barbiturates, Anti-Anxiety Tranquilizers, Anti-Depressants, Anti-Psychotic Tranquilizers, & Combinations of the first five.

Cocaine, the Amphetamines, and "Others".

Opiates and Synthetics.



CNS Depressant

CNS Stimulant



Aides	Lesson Plan	Instructor Notes
	c. Secobarbital	CNS Depressant
	d. Dilaudid	Narcotic Analgesic
	e. Alprazolam	CNS Depressant
	f. Phenyl Cyclohexyl Peperidine	Phencyclidine
	g. "Ecstasy"	Hallucinogen
	h. ETOH	CNS Depressant
	i. Numorphan	Narcotic Analgesic
	j. Psilocybin	Hallucinogen
	4. List the twelve components of the Drug Influence Evaluation in the proper sequence.	Breath Alcohol test; Interview of Arresting Officer; Preliminary Examination; Eye Examinations; Divided Attention Tests; Vital Signs Examinations; Darkroom Examinations; Check for Muscle Tone; Injection Sites Inspection; Statements of Suspect; Evaluator's Opinion; Toxicological Examination.
	a. Demonstrate the Preliminary Examination.	Allow student-demonstrations to refer to the standard Drug Influence Evaluation Form.
	b. Demonstrate the Eye Examinations.	Be sure to provide appropriate positive feedback and constructive criticism of the demonstrators' performances.
	c. Demonstrate the Administration of the Divided Attention Tests.	
	d. Demonstrate the Vital Signs Examinations.	·

Aides	Lesson Plan		Instructor Notes
		e. Demonstrate the Darkroom Examinations.	
		f. Demonstrate the Check for Muscle Tone and the inspection for Injection Sites.	
MCR-3 (Name the)	5.	Identify the category for each of the listed drugs.	
	,	a. Demerol	Narcotic Analgesic
		b. Cylert	CNS Stimulant
		c. Chlordiazepoxide	CNS Depressant
		d. Ketamine	Phencyclidine
		e. Percodan	Narcotic Analgesic
		f. Ritalin	CNS Stimulant
		g. Isopropanol	CNS Depressant
		h. Bufotenine	Hallucinogen .
		i. Thebaine	Narcotic Analgesic
		j. Methaqualone	CNS Depressant
	В.	Eyes and Vital Signs	
50 Minutes  MCR-4 (Eyes			
and Vital)			
	1.	Name the three clues of Horizontal Gaze Nystagmus.	Lack of smooth pursuit; distinct nystagmus at maximum deviation; angle of onset.

Aides		Lesson Plan	Instructor Notes	
		a. Demonstrate the check for "Lack of smooth pursuit".		
		b. Demonstrate the check for "Distinct nystagmus at maximum deviation".	Ask the student-demonstrator: How long should the eye be held at maximum deviation? (About four seconds)	
		c. Demonstrate the check for "Angle of Onset".	Ask the student-demonstrator: What is the formula that expresses the approximate relationships between BAC and Angle of Onset? (BAC = 50 - Angle)	
	2.	Name the categories of drugs that will cause Horizontal Gaze Nystagmus.	CNS Depressants, Phencyclidine, Inhalants.	
7.7		a. Name the categories that will cause <b>Vertical</b> Gaze Nystagmus.	Same as above.	
		b. Demonstrate the check for Vertical Gaze Nystagmus.	Ask the student-demonstrator: How long should the eye be held at maximum elevation? (About four seconds)	
	3.	Name the test that is always administered immediately after Vertical Gaze Nystagmus.	Lack of Convergence.	
		a. Demonstrate the test for Lack of Convergence.		
		b. Name the categories of drugs that usually will cause Lack of Convergence.	CNS Depressants; Phencyclidine; Inhalants; Cannabis.	
	4.	Name the lighting conditions under which we make estimations of pupil size.	Room light; near-total darkness; direct light.	
}		a. Demonstrate the room light pupil size estimation procedure.		

Aides		Lesson Plan	Instructor Notes
	b	o. Demonstrate the near-total darkness procedure.	
	с	. Demonstrate the direct light procedure.	Ask the student-demonstrator: How large should the circle of light appear on the suspect's face for the direct-light check? (Approximately the same as the eye socket)
			Ask the student-demonstrator: How long should the light be shined directly into the suspect's eye? (Fifteen seconds)
	d	. Name the other things a DRE looks for while shining the light directly into the suspect's eye.	Pupil reaction to light; hippus; rebound dilation.
	e	. How quickly must the pupil start to constrict if it is considered to exhibit normal reaction to light?	Within one second.
	f.	Define Hippus.	A rhythmic pulsating of the pupils of the eyes, as they dilate and constrict within fixed limits.
	g	. Define Rebound Dilation.	Rebound dilation is a period of constriction followed by dilation with a change equal to or greater than 2 mm.
		tate the normal range of pupil	3.0 mm to 6.5 mm (in diameter).



these...)

terms.

Define each of the listed

Aides	Lesson Plan	Instructor Notes
	o Miosis	Abnormally constricted pupils
	o Mydriasis	Abnormally dilated pupils
	o Ptosis	Droopy eyelids
	b. What kinds of drugs will cause dilation of the pupils?	CNS Stimulants; Hallucinogens; Cannabis (although sometimes only slight dilation, if any).
	c. What kinds of drugs will cause constriction?	Narcotic Analgesics.
MCR-6 (More	6. Identify the category for each of the listed drugs.	
drugs)	a. Fentanyl	Narcotic Analgesic
	b. Halcion	CNS Depressant
	c. Librium	CNS Depressant
·	d. Peyote	Hallucinogen
	e. Darvon	Narcotic Analgesic
	f. Preludin	CNS Stimulant
	g. Diazepam	CNS Depressant
	h. Biphetamine	CNS Stimulant
	i. Hycodan	Narcotic Analgesic
	j. Percobarb	Combination of CNS Depressant and Narcotic Analgesic
	7. Define "Pulse".	The expansion and relaxation of an artery, generated by the pumping action of the heart.
		(Also acceptable: The expansion and relaxation of an artery, caused by the surging flow of blood.)

Aides		Lesson Plan	Instructor Notes
	a.	Define "Pulse Rate".	The number of pulsations in an artery per minute.
	b.	Define "Artery".	A strong, elastic blood vessel that carries blood from the heart to the body tissues.
	c.	Define "Vein".	A blood vessel that carries blood back to the heart from the body tissues.
MCR-7A&B	d.	Identify the location of each listed pulse point.	
(Where are)		o Radial	In the wrist, at the base of the thumb.
		o Brachial	In the crook of the arm.
		o Carotid	In the neck, on either side of the Adam's Apple
	e.	Demonstrate a pulse measurement, using the left Radial pulse point.	
	f.	State the normal range of adult human pulse rate.	60 to 90 beats per minute.
	g.	Name the drug categories that usually cause elevated pulse rate.	CNS Stimulants; Hallucinogens; Phencyclidine; Inhalants; Cannabis.
	h.	Name the drug categories that usually cause lowered pulse rate.	CNS Depressants; Narcotic Analgesics.
;	8. De	fine "Blood Pressure".	The force exerted by blood on the walls of the arteries.
	a.	How often does a person's blood pressure change?	It is <b>always</b> changing, from instant to instant.
i vi	b.	When does the blood pressure reach its highest value?	When the heart is fully contracted, and blood is sent rushing into the arteries.

Aides	_	Lesson Plan	Instructor Notes
	C.	When does the blood pressure reach its lowest value?	When the heart is fully expanded, just before it starts to contract for the next "pumping" action.
22 3	d.	Name the two medical instruments that are used to measure blood pressure.	Select a student to come to the chalkboard or flipchart and print "SPHYGMOMANOMETER" and "STETHOSCOPE".
	e.	Name the sounds that we hear through the stethoscope when we make a blood pressure measurement.	Select a student to come to the chalkboard or flipchart and print "KOROTKOFF SOUNDS".
	f.	What does this "Hg" mean?	Instructor: Print "Hg" on the chalkboard or flipchart.  Chemical symbol for the
96 944			element Mercury; abbreviation for the Latin word Hydrargyrum, meaning "Mercury".
T I I	g.	In what units is blood pressure measured?	Millimeters of Mercury. Instructor: Print "mm" on the chalkboard or flipchart, right in front of the "Hg".
	h.	Suppose that, at some particular instant, a person has a blood pressure of 120 mmHg. What does that "120 mmHg" mean?	It means the pressure would be strong enough to push a column of liquid Mercury up a glass tube to a height of 120 millimeters.
			INSTRUCTOR, IF ONE IS AVAILABLE, DISPLAY A SPHYGMOMANOMETER THAT HAS A LIQUID MERCURY PRESSURE GAUGE.

Aides	Lesson Plan	Instructor Notes
	i. Name the types of dr that usually cause a blood pressure.	· · · · · · · · · · · · · · · · · · ·
	j. Name the types of di that elevate blood pr	
MCR-8	k. State the meaning of the listed terms.	feach of
(Some technical)	o Systolic	The highest value of blood pressure.
	o Diastolic	The lowest value of blood pressure.
	o Bradycardia	Abnormally slow heart rate; pulse rate below the normal range.
	o Tachycardia	Abnormally rapid heart rate; pulse rate above the normal range.
	o Hypertension	Abnormally high blood pressure.
	o Hypotension	Abnormally low blood pressure.
	l. State the normal ran systolic blood pressu	
	m. State the normal ran diastolic blood pressu	
	n. Demonstrate the measurement of bloo pressure.	Tell the student-demonstrator to explain out loud everything he or she does to take blood pressure measurement.

Aides	Lesson Plan	Instructor Notes
	C. Physiology	
20 Minutes  MCR-9 (Physiology)		
	1. Define "Physiology".	The study of the functions of living organisms and their part.
	2. What is the expression we use to remember the names of the ten major body systems?	Select a student to come to the chalkboard or flipchart, and print "MURDERS INC" vertically.
	a. What is M for?	Muscular (Have a student print out each name).
	b. What is <b>U</b> for?	Urinary
	c. What is the first R for?	Respiratory (or, Reproductive)
	d. What is $D$ for?	Digestive
	e. What is E for?	Endocrine
·	f. What is the second $\mathbf R$ for?	Reproductive (or, Respiratory)
ļ	g. What is $S$ for?	Skeletal
	h. What is I for?	Integumentary
	i. What is $N$ for?	Nervous
	j. What is C for?	Circulatory
	3. State the word that means "dynamic balance involving levels of salts, water, sugars and other materials in the body's fluids".	Homeostasis.

#### Lesson Plan

#### **Instructor Notes**

- 4. Which artery carries blood from the heart to the lungs?
  - a. What is unique about the Pulmonary artery, compared to all other arteries?
  - b. What are the Pulmonary veins?
  - c. What is unique about the Pulmonary veins?
- Name the various types of nerves.
  - a. Sensory Nerves, carry messages to the brain.
  - b. Motor Nerves, carry messages from the brain.
  - c. Voluntary Nerves are motor nerves that carry messages to the muscles that we consciously control.
  - d. Autonomic Nerves are motor nerves that carry messages to the muscles and organs we do not consciously control.

Pulmonary.

- (1) it is the only artery that takes blood from the right side of the heart;
- (2) it is the only artery that carries **deoxygenated** blood (i.e., blood that is depleted of oxygen).

The veins that carry blood back to the heart from the <u>lungs</u>.

(1) they are the only veins that bring blood to the left side of the heart; (2) they are the only veins that carry **oxygenated** blood.

Ask students to "fill in" the missing names. Use an acetate overlay on Visual MCR-10 to "write in" the students' responses.

Also known as Afferent Nerves.

Also known as Efferent Nerves.



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#### Lesson Plan

#### **Instructor Notes**



- e. Sympathetic Nerves are autonomic nerves that carry messages commanding the body to react to fear, stress, excitement, etc.
- f. Parasympathetic Nerves are autonomic nerves that carry messages to produce relaxed and tranquil activities.
- 6. Define each of the listed terms.
  - a. Neuron
  - b. Synapse
  - c. Neurotransmitter
  - d. Axon
  - e. Dendrite
- D. Questions and Answers

Clarification: Sympathetic nerves carry the brain's "fire alarms" and "wake up calls".

Clarification: Parasympathetic nerves carry the brain's "all clear" and "at ease" messages.

A nerve cell; the basic "building block" of a nerve.

The gap or space between two nerve cells.

A chemical that flows across the synapse, to carry a message from one neuron to the next.

The end of a neuron that sends out the neurotransmitter.

The end of a neuron that receives the neurotransmitter.

Segment D: As long as necessary

Solicit and answer students' questions about anything covered thus far in their training.



#### **Mid-Course Review**

Review of Drugs, Drug Categories, and the Drug Influence Evaluation





#### What Kinds of Drugs are These?

- Xanax
- · "Ecstasy"
- Desoxyn
- ETOH
- · Secobarbital
- Numorphan
- Dilaudid
- Psilocybin
- Alprazolam
- Phenyl Cyclohexyl Piperidine

Drug Evaluation & Classification Training

MCR-2

#### Name the Categories:

- Demerol
- Ritalin
- Cylert
- Isopropanol
- · Chlordiazepoxide
- Bufotenine
- Ketamine
- Thebaine
- Percodan
- Methagualone

Drug Evaluation & Classification Training

MCR-3

#### **Eyes and Vital Signs Review**



Drug Evaluation & Classification Training

MCR-4

#### What Do These Words Mean?

- Miosis
- Mydriasis
- Ptosis

Drug Evaluation & Classification Training

LICEE

#### **More Drugs to Categorize**

- · Fentanyl
- Preludin
- Halcion
- Diazepam
- Librium
- Biphetamine
- Peyote
- Hycodan
- Darvon
- Percobarb

drug Evaluation & Classification Training

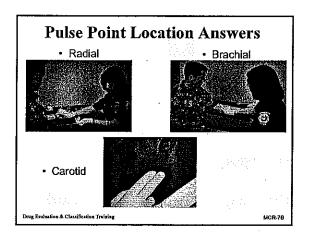
MCR-6

# Where Are These Pulse Points Located?

- Radial
- Brachial
- Carotid

Drug Evaluation & Classification Training

ICR-7A

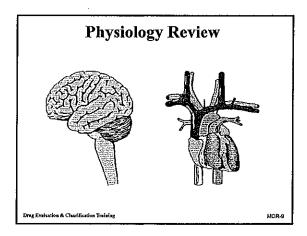


# Some Technical Terms to Define

- Systolic
- · Diastolic
- Bradycardia
- Tachycardia
- Hypertension
- Hypotension

Drug Evaluation & Classification Training

MCR-8



# Classification of Nerves All Nerves Motor Nerves (AKA "Afferent" Nerves) Autonomic (not conscious control) Sympathetic Fear, stress, etc. Parasympathetic relaxation, tranquillity, etc. Drug Evaluation & Chasification Training MGR-10

## Some More Technical Terms to Define

- Neuron
- Synapse
- Neurotransmitter
- Axon
- Dendrite

Drug Evaluation & Classification Training

MCR-11

Forty-Five Minutes

### SESSION XVIII

PRACTICE: TEST INTERPRETATION

#### SESSION XVIII PRACTICE: TEST INTERPRETATION

Upon successfully completing this session, the participants will be able to:

- o Analyze the results of a complete DRE examination and identify the category or categories of drugs affecting the individual examined.
- o Articulate the bases for the drug category identification.

#### Content Segments

- A. Interpretation Demonstrations
- B. Interpretation Practice

#### Learning Activities

- o Instructor Led Demonstrations
- o Small Group Practice
- o Participant Led Presentations

Aides	Lesson Plan	Instructor Notes
- N	PRACTICE: TEST INTERPRETATION	Total Lesson Time: Approximately 45 Minutes
		Point out the "Test Interpretation" wallchart.
XVIII-O (Objectives)		Briefly review the objective content and activities of this session.
	A. Interpretation Demonstrations.	,
20 Minutes		
	1. Case #1 "Subject Martinez"	Direct students to turn to the "Subject Martinez" exemplar in Section XVIII of their manual.
	a. Preliminary Examination.	Review the results of the Preliminary Examination of Subject Martinez.
		Ask students: "What category or categories of drugs would produce preliminary examination results consistent with this exemplar?" Probe to draw out the basis for students' responses.
	b. Eye Examinations.	Review the results of the eye examination of Subject Martinez.
		Ask students to discuss the category or categories of drugs that would cause these eye examination results.
	c. Psychophysical Tests.	Review the results of the psychophysical tests of Subject Martinez.

Aides	Lesson Plan	Instructor Notes
		Ask students to discuss the category or categories of drugs that would produce these psychophysical tests results.
	d. Vital Signs Examinations.	Review the results of the Vital Signs Examinations of Subject Martinez.
		Ask students to discuss the category or categories of drugs that would cause these results.
	e. Dark Room Examinations.	Review the results of the Dark Room Examinations of Subject Martinez.
		Ask students to discuss the category or categories of drugs that would produce these results.
	f. Other evidence.	Review the results of the examinations for injection sites and muscle rigidity, and of the final interview of Subject Martinez.
·		Ask students to comment on the category or categories of drugs that would be consistent with all of the evidence on this exemplar.
	g. Opinions of Evaluator.	<u>Point out</u> that the evidence indicates that Subject Martinez is under the influence of PCP.
		Solicit students' questions concerning this demonstration.
	2. Case #2: "Subject Groves".	Direct students to review the "Subject Groves" exemplar.
		·

Aides	Lesson Plan	Instructor Notes
	a. Preliminary Examination.	Review the results of the Preliminary Examination of Subject Groves.
		Ask students: "What category or categories of drugs would produce preliminary examination results consistent with this exemplar?" Probe to draw out the basis for students' response.
	b. Eye Examinations.	Review the results of the eye examinations of Subject Groves.
		Ask students to discuss the category or categories of drugs that would cause these eye examination results.
	c. Psychophysical Tests.	Review the results of the psychophysical tests of Subject Groves.
		Ask students to discuss the category or categories of drugs that would produce these psychophysical test results.
	d. Vital Signs Examinations	Review the results of the Vital Signs Examinations of Subject Groves.
		Ask students to discuss the category or categories of drugs that would produce these results.
	e. Dark Room Examinations.	Review the results of the Dark Room Examinations of Subject Groves.

Aides	Lesson Plan	Instructor Notes
		Ask students to discuss the category or categories of drugs that would produce these results.
	f. Other evidence.	Review the results of the examinations for injection sites and muscle rigidity, and of the final interview of Subject Groves.
		Ask students to comment on the category or categories of drugs that would be consistent with all of the evidence on this exemplar.
	g. Opinions of Evaluator.	Point out that the evidence indicates that Subject Groves is under the influence of a Narcotic Analgesic.
	B. Interpretation Practice	Solicit students' questions concerning this demonstration.
25 Minutes		
·	1. Team practice	Assign students to work in teams of three or four members.
		Tell teams that they are to review four exemplars (Subjects Hatos, Schafer, Johnson and Sholly). Team members are to discuss the evidence among themselves and reach a conclusion concerning the category or categories of drugs; if any.
		Teams will present their conclusions to the entire class.

Aides	Lesson Plan	Instructor Notes
	a. Review and discussion of exemplars by teams.	Allow teams approximately 1 minutes to review the three exemplars and reach their conclusions.
	<ul> <li>b. Feedback of results.</li> <li>o Subject Martinez</li> <li>o Subject Groves</li> <li>o Subject Hatos</li> <li>o Subject Schaffer</li> <li>o Subject Johnson</li> <li>o Subject Sholly</li> </ul>	Poll the teams to determine their conclusions concerning the category or categories of drugs present in each subject Offer appropriate comments concerning the teams' performance.
`	2. Session Wrap up.	Solicit students' comments and questions concerning this practice session.

#### DRUG CATEGORIES FOR INTERPRETATION PRACTICE

**SUBJECT** 

CATEGORY(TES)

Martinez

PCP

Groves

Narcotic Analgesic

Hatos

CNS Stimulant and ETOH

Schaffer

PCP and CNS Depressant

Johnson

PCP and Narcotic Analgesic

Sholly

Medical rule out

## **Session XVIII**

Practice: Test Interpretation



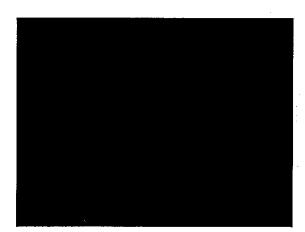
## **Practice: Test Interpretation**

Upon successfully completing this session, the participants will be able to:

- Analyze the results of a complete Drug Evaluation and Classification examination and identify the category or categories of drugs affecting the individual examined
- Articulate the bases for the drug category identification

Drug Evaluation & Clearification Training

XVIII-0



**Drug Influence Evaluation** 0112 Recorder/Witness None □ Property □ Fatal Injury S. Richardson estee's Name (Last, First, MI) Arresting Officer (Name, ID No.) Sex Chuck Crum W Breath Results: 0.03 7 Refused Chemical Test **U**rine □ Blood Instrument# 1234 □ Refused 122/2000 2330 Time of last drink? Miranda Warning Given: XYes □ No What have you eaten today? Have you been drinking? How much? Nothing Nothing By: Wayne Warner When did you last sleep? How long? Are you sick or injured? □ No Are you diabetic or epileptic? -Yes -No Not SIC/s t's late Kwer have any physical defects? 

Yes 

No Are you under the care of □ No □ Yes Not SICK No + SICK No Answer
Coordination Hoor Unsteady
Staggering Are you taking any medication or drugs? 

Yes 

No NON Responsive - Passive Not Sick Chemicalodor Tracking: ⊭ Equal □ Unequal □ Reddened Conjunctiva Blindness: □ Normal #Bloodshot □ Watery KNone □LEye □REye Able to follow stimulus: FYes □ No Eyelids: Pupil Size: #Equal Normal □ Droopy □ Glasses □ Contacts, if so □ Hard □ Soft □ Unequal (explain) HGN Left Eye Right Eye Vertical Nystagmus? ¥Yes Pulse & Time 2340 4es Lack of Smooth Pursuit Convergence Right Eye Left Eye 2356 Max. Deviation 0010 Angle of Onset Walk and Turn Test "Moonwalking Legs & arms rigid. Romberg Balance Cannot Keep Balance Approx. Starts Too Soon 1st Nine 2nd Nine Sways While Balancing Stops Walking Misses Heel-Toe □ Uses Arms to Balance Steps Off Line □ Hopping Raises Arms  $\sqrt{\sqrt{}}$  $\sqrt{\nu}$ Actual Steps Taken Describe Turn Turned Dackward Cannot Do Test (Explain) Type of Footwear Internal Clock Loafers 30_Estimated At 30 Sec. stopped for losec. after tun Pupil Size Room Light Darkness Direct ○ Right △ Left Left Eye 6.0 Draw lines from spot touched Oral Cavity Right Eye Rebound Dilation Reaction To Light Hippus O Yes Norma **#**No □ Yes XNo 2 Temp_ Blood Pressure Muscle Tone: □ Normal Attach Photos Of Fresh Puncture Marks Arms and legs. Time of Use? Where Were The Drugs Used? (Location) How Much? Haswel Time Completed Time DRE Notified 23 No. 2379 PCP □ Inhalant □ Rule Out □ Alcohol □ Stimulant Opinion of Evaluator: ☐ Narcotic Analgesic □ Cannabis □ Hallucinogen □ Medical □ Depressant

	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>	
LOG NO.	DRE: Trooper Wayne Warner	ARRESTEE: J.M. Martinez	
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / I ONS 6. MEDICAL PROBLEMS 7. PSYCHOR N 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS	
1. LOCATION: Exami	nation of J.M. Martinez took place in the DF	RE room, SP Albany, Troop T.	
2. WITNESS: S. Richar	rdson (NHTSA) and Chuck Crum (IACP)		
3. BREATH TEST: W	riter administered breath test to Martinez, the	he result was 0.00%.	
4. NOTIFICATION / I	INTERVIEW of ARRESTING OFFICER	: Writer was the arresting officer.	
5. INITIAL OBSERVA	ATIONS: Writer observed subject seated in	the drivers position of a blue, 1996 Oldsmobile, NY	
registration "277 BR	X". Vehicle was stationary in the Northbour	nd lane of Hannover Ave., at the intersection with	
Hugenot St. The traf	fic light was green and the other vehicles ha	d to pull out and around subject's vehicle.	
6. MEDICAL PROBL	EMS: None noted or stated		
7. PSYCHOPHYSICA	L TESTS: Romberg Balance: Subject swa	yed approximately 3" side to side.	
Walk and Turn: Subj	ect lost his balance during the instructions, s	topped walking, turned backwards.	
He paused for approximately ten (10) seconds after turning and exhibited muscle rigidity in his arms and legs			
throughout the test. One Leg Stand: Subject raised his arms, put his foot down, staggered and nearly fell			
at this point the test was stopped. Finger to Nose: Subject missed tip of his nose four times.			
8. CLINICAL INDICATORS: Subject had HGN, Vertical Nystagmus and Lack of Convergence. His pulse was			
above the normal.			
9. SIGNS of INGESTI	ION: Subject's breath had a strong chemical	l odor.	
10. STATEMENTS: Subject was very passive throughout the evaluation and was very slow at responding to questions.			
He repeatedly answered "not sick" to questions concerning the use of medication. He also failed to respond to a			
couple of the quest	ions		
11. OPINION of EVALUATOR: In my opinion J.M. Martinez is under the influence of			
and unable to opera	te a vehicle safely.		
12. TOXICOLOGICA	L SAMPLE: Subject agreed to provide a ur	ine sample.	
13. MISCELLANEOU	S:		
	494500		

Drug Influence Evaluation
Rolling Log No. 76-19 Evaluator 000612 CLARK, KEN Recorder/Witness □ Fatal □ Injury □ Property DELVECHIO, J.J. stee's Name (Last, First, MI) DOB Race Arresting Officer (Name, ID No.) Sex DELVECHIO OUES ROBERT Date Examined/Time/Location 3RD PCT Breath Results: , and Refused Chemical Test ✓ Urine Instrument# □ Refused AUG 15 1996 UBPD 1234 Miranda Warning Given: ♥Yes □ No What have you eaten today? When? Have you been drinking? How much? Time of last drink? NOTHING CLARK FRIED CHECKEN Time now? When did you last sleep? How long? Are you sick or injured? Are you diabetic or epileptic? □ Yes 🧇 No MIDNIGHT LAST NIGHT Do you take insulin? Do you have any physical defects? 

Yes No Are you under the care of Coordination STUMBLING Are you taking any medication or drugs? □ Yes □ No (のりまた) E FOR Attitude Breath NORMAL ONOR
BREATHING SLOW & SHALLOW BACKPAIN AS NEEDED NORMAL Tracking: Blindness: Eyes: 

Reddened Conjunctiva <page-header> Equal 🗆 Unequal Normal □ Bloodshot □ Watery **#**None □LEye □REye SLOW KNYUMBLES Corrective Lens: **№**None Able to follow stimulus: ♣Yes □ No Eyelids: Pupil Size: 😼 Equal □ Glasses □ Contacts, if so □ Hard □ Soft □ Unequal (explain) □ Normal 👿 Droopy Pulse & Time HGN Left Eye Right Eye 1.60,010 Lack of Smooth Pursuit NO NO Convergence Right Eye Left Eye Max. Deviation NO Angle of Onset NONE Romberg Balance Walk and Turn Test Cannot Keep Balance 🗸 🗸 Starts Too Soon Approx. 1st Nine 2nd Nine Sways While Balancing Stops Walking M Uses Arms to Balance Misses Heel-Toe Steps Off Line □ Æløpping Raises Arms ✓ Puts Foot Down Actual Steps Taken Describe Turn LOST BALANCE Type of Footwear Internal Clock Cannot Do Test (Explain) Estimated At 30 Sec. STAGEFRED TO RIGHT Pupil Size Room Light Darkness Direct ○ Right △ Left Left Eye Draw lines from spot touched Oral Cavity Right Eye Hippus Rebound Dilation Reaction To Light LIME □ Yes 🞜 No □Yes 🏚 No 100 Temp <u>77.</u> 8 Blood Pressure 106 Muscle Tone: □ Normal Flaccid □ Rigid Attach Photos Of Fresh Puncture Marks Comments: ARMS & NECK RUBBELY t Medicine or Drug Have You Been Using? How Much? Time of Use Where Were The Drugs Used? (Location) AROUND OUPLE OF PELLS FOR MY BACK RESTAURANT D EUNER Eval Start Time Time DRE Notified Time Completed 0025 □ Inhalant Opinion of Evaluator: □ Rule Out □ Alcohol □ Stimulant  $\square$  Depressant  $\square$  Hallucinogen □ Cannabis □ Medical □ Narcotic Analgesic

	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>		
LOG NO.	DRE: Sgt. Ken Clark	ARRESTEE: Robert G. Groves		
1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.				
1. LOCATION: Exami	nation of Robert G. Groves, took place in th	e DRE room, 3rd Pct. Virginia Beach PD		
2. WITNESS: Arresting	, Officer - Trooper J.J. Delavecchio			
3. BREATH TEST: W	riter observed Trooper J.J. Delavecchio adm	ninister a breath test to Groves, the result was 0.00%		
4. NOTIFICATION /	INTERVIEW of ARRESTING OFFICER	: Writer was contacted by radio and advised to		
return to the precinct	to conduct a DRE evaluation. Tpr Delave	cchio informed me that he had observed		
the subject's vehicle	drifting across the center line and driving 1	5 mph in a 45 mph zone. Tpr Delavecchio further		
stated that the subjec	t admitted to taking "a few" pain pills.			
5. INITIAL OBSERVA	ATIONS: Writer observed the subject seated	d int he breath testing room VBPD. Subject appeared		
sleepy with his eyes	closed and head nodded forward. He was co	poperative throughout the examination.		
6. MEDICAL PROBL	EMS: Subject stated that he had taken codie	ene pills to alleviate back pain, and that he'd had an		
apointment with his	doctor earlier that day. He further stated tha	the was not experiencing any pain at this time.		
7. PSYCHOPHYSICAL TESTS: Romberg Balance: Subject swayed side to side and front to back, and estimated 53				
seconds as 30 seconds. Walk and Turn: Subject lost his balance during the instructions, missed heel to toe, and lost				
his balance while turn	his balance while turning. One Leg Stand: Subject raised his arms, put his foot down, and swayed. Finger to Nose:			
Subject missed tip of	nose on each attempt.			
8. CLINICAL INDIC	8. CLINICAL INDICATORS: Subject's blood pressure was below the normal range and his pupils were constricted.			
9. SIGNS of INGESTION: None were evident				
10. STATEMENTS: S	ubject stated he had taken "a couple of pills	for my back". He also staed that the pills contained		
Codiene.				
11. OPINION of EVALUATOR: In my opinion Robert G. Groves is under the influence of a				
and unable to operate a vehicle safely.				
12. TOXICOLOGICA	L SAMPLE: Subject agreed to provide a un	ine sample.		
13. MISCELLANEOU	S:			
		·		

**Drug Influence Evaluation** Rolling Log No. Evaluator 2838 49 XVIII-3 KLIMA Recorder/Witness Crash: □ None □ Fatal □ Injury □ Property estee's Name (Last, First, MI) DOB Sex Race Arresting Officer (Name, ID No.) 3  $\omega$ STEPHEN UNSWORTH J. #1811 PHOEDIX PD COTA Date Examined/Time/Location Breath Results: □ Refused Chemical Test XUrine □ Blood MARICOPA COUNTY Instrument# □ Refused 2300 0.04 11.25.2000 358C| Miranda Warning Given: p∀Yes □ No What have you eaten today? When? Have you been drinking? How much? Time of last drink? RUAST BEEF 2 Hes AGO JUNG TO ZZALO A KLIMA 2 HRS 460 Time now? When did you last sleep? How long? Are you sick or injured? □ Yes KNo Are you diabetic or epileptic? □Yes X No 8HRS LAST NIGHT Do you take insulin? Do you have any physical defects? Yes No Are you under the care of a doctor or dentist? □ Yes 😿 No Are you taking any medication or drugs? 

Yes XNo Attitude Coordination COOPERATIVE & NERWOUS STUMBLING POOR JERKY Breath Face DOOR OF ALCOHOLIC Eyes: Reddened Conjunctiva BEVERAGE NORMAI Tracking: Speech NORMAL BUT VERY Blindness: None □LEye □REye Normal □ Bloodshot □ Watery M Equal □ Unequal TALKATHE Corrective Lens: Pupil Size: Equal Able to follow stimulus: ¥Yes □ No Evelids: □ Glasses □ Contacts, if so □ Hard □ Soft □ Unequal (explain) Mormal □ Droopy Pulse & Time HGN Left Eye Right Eye Vertical Nystagmus? 

☐ Yes 

✓ No One Leg Stand 1, 100 / 2340 Lack of Smooth Pursuit YES 455 Convergence Right Eye Left Eye 2, 104 / 2349 Max. Deviation No  $\mathcal{O}_{\mathcal{O}}$ 8. 108 / 3358 Angle of Onset NONIE このじゃ Romberg Balance Walk and Turn Test Cannot Keep Balance Starts Too Soon Approx. Approx. 1st Nine 2nd Nine Sways While Balancing Stops Walking Misses Heel-Toe √ Uses Arms to Balance Steps Off Line □ Hopping Raises Arms ✓ Puts Foot Down Actual Steps Taken Internal Clock Describe Turn Cannot Do Test (Explain) Type of Footwear Estimated At 30 Sec. N/A LOAFERS AS INSTRUCTED Nasal Area REDNESS ま Pupil Size Room Light Darkness Direct ○ Right △ Left ULCERATIONS Left Eye Draw lines from spot touched  $( \omega, \mathcal{O} )$ 8.5 5.5 **Oral Cavity** Right Eye ( . O E.8 5.5 CLEAR Rebound Dilation Hippus □ Yes Reaction To Light ¢#No □ Yes 💆 No SLOW NO VISIBLE MARKS 146 99.2 Blood Pressure Temp Muscle Tone: A Normal □ Flaccid □ Rigid Attach Photos Of Fresh Puncture Marks Comments: t Medicine or Drug Have You Been Using? How Much? Time of Use? Where Were The Drugs Used? (Location) "I DIDIO TECHS FORID I" I DIDAT DO ADO NE NA Time Completed Date/Time of Arrest Time DRE Notified 33 <u>20</u> Eval Start Time 11-25-200 2310 0100 11-26-2000 Member Signature (Include Rank) ID No. Reviewed By: D. LAMM 10084 Opinion of Evaluator: □ Rule Out □ Stimulant □ Alcohol □ Inhalant □ Medical □ Hallucinogen □ Depressant □ Narcotic Analgesic □ Cannabis

000615				
	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>		
LOG NO.	DRE: Lt. Joseph Klima	ARRESTEE: Stephen H. Hatos		
1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.				
1. LOCATION: Exami	nation of Stephen H. Hatos, took place in the	e DRE room, Maricopa County Jail		
2. WITNESS: Arresting	g Officer - J. Unsworth #1811			
3. BREATH TEST: Of	fficer Unsworth administer a breath test to H	atos, the result was 0.04%		
4. NOTIFICATION /	INTERVIEW of ARRESTING OFFICER	: Writer was contacted by radio and advised to		
return to the jail to co	onduct a DRE evaluation. Officer Unsworth	informed me that he had observed the subject		
driving at excessive	speed and he failed to stop at a red traffic lig	tht. Officer Unsworth further stated that the subject		
appeared nervous an	d performed poorly on the SFSTs.			
5. INITIAL OBSERVA	ATIONS: Writer observed the subject seated	d in the breath testing room. Subject was very		
talkative, repeatedly	shifted his weight from foot to foot, and exl	nibited nervous abrupt movements with his hands.		
When not speaking he appeared to grind his teeth. There was also an odor of alcoholic beverage on the subjects				
breath.				
6. MEDICAL PROBL	EMS: None noted or stated			
7. PSYCHOPHYSICAL TESTS: Subject performed all of the tests in a stumbling jerky fashion. Romberg Balance:				
Subject swayed approximately 3" side, and estimated 20 seconds as 30 seconds. Walk and Turn: Subject				
lost his balance durir	lost his balance during the instructions, and stopped walking and used his arms for balance. One Leg Stand: Subject			
raised his arms, put l	nis foot down, and swayed. Finger to Nose:	Subject missed tip of nose on each attempt.		
8. CLINICAL INDICATORS: Subject's blood pressure and pulse were above the normal range.				
9. SIGNS of INGESTION: None were evident				
10. STATEMENTS: Subject stated, "I didn't snort anything"				
11. OPINION of EVALUATOR: In my opinion Stephen H. Hatos is under the influence of a				
and unable to opera	te a vehicle safely.			
12. TOXICOLOGICA	L SAMPLE: Subject agreed to provide a ur	ine sample.		
13. MISCELLANEOU	S:	-		

**Drug Influence Evaluation** Rolling Log No. Evaluator 000616
POFF MEL
Recorder/Witness 422 XVIII -4 Crash: X None O Fatal □ Injury □ Property JOHN MCKAY (TEXAS DECP)

Pastee's Name (Last, First, MI) DÓB Arresting Officer (Name, ID No.) Sex Race LHAFER, BRIND 3:15.66 POFF, M. # 314 ω HPD Date Examined/Time/Location Chemical Test Breath Results: □ Refused Ŭrine □ Blood CENTRAL Instrument # □ Refused 00EC 1005, 71 CAR 0.00 WITOY. CO12838 Miranda Warning Given: XYes □ No What have you eaten today? Have you been drinking? How much? Time of last drink? By: M. POFF CHINESE FOOD JUST WATER A/CA الساندال Time now? When did you last sleep? How long? Are you sick or injured? □ Yes >KNo Are you diabetic or epileptic? ARGUT □ Yes #No 2 HOURS LAST NIGHT 8 PM □ Yes No Do you take insulin? Do you have any physical defects? Yes No Are you under the care of a doctor or dentist? 

XYes □ No FOR STRESS Coordination Attitude COOPERATIVE POOR STAGGERING DETARHED Breath Face NORMAL COLOR BLANK STARE VALIUM 10mg TWICE A DAY Speech THICK, SLURRED, SLOW CHEMICAL ODOS Eyes: Reddened Conjunctiva Blindness: Tracking: □ Normal □ Bloodshot X Watery None OLEye OREye  $\mathbf{X}$ Equal  $\square$  Unequal でのののファ ≭None Corrective Lens: Pupil Size: KEqual Able to follow stimulus: X Yes □ No Eyelids: □ Glasses □ Contacts, if so □ Hard □ Soft □ Unequal (explain) Normal Droopy Pulse & Time HGN One Leg Stand Left Eye Right Eye Vertical Nystagmus? XYes □ No dg 13310 Lack of Smooth Pursuit Convergence YE5 Right Eye Left Eve 13225 Max. Deviation YES 94 12235 30° Angle of Onset Walk and Turn Test PEPEAT Romberg Balance Cannot Keep Balance 🗸 🗸 Starts Too Soon_ Approx. Approx. 1st Nine 2nd Nine か Sways While Balancing Stops Walking Misses Heel-Toe Uses Arms to Balance Steps Off Line C Hopping Raises Arms IJ Actual Steps Taken Puts Foot Down 0 10 Internal Clock Describe Turn Cannot Do Test (Explain) Type of Footwear حل Estimated At 30 Sec. TURNOD BACKWADS 4/4 PUNINING SHOES Pupil Size Room Light Darkness Direct Nasal Area VERY RIGID ARM MOJEMENTS Right A Left CLEAR Left Eye Draw lines from spot touched 4.0 4.0 6.0 Oral Cavity PED SUDJUCT US DUTAGOS Right Eye 4.0 6.0 0.4 Rebound Dilation Hippus □ Yes Reaction To Light **K**No □ Yes XNo SLOW Blood Pressure___ Temp 99.2. 100 Muscle Tone: ☐ Normal ☐ Flaccid Kigid Comments: RIGIO ARMS & NECK

* Medicine or Drug Have You Been Using? How Attach Photos Of Fresh Puncture Marks Where Were The Drugs Used? (Location) Time of Use? or Thaid I YESTERDAY DUYTHING ELSE. Date/Time of Arrest Time DRE Notified Eval Start Time Time Completed 2120 01-17-01 1D No. 314 Member Signature (Include Rank) Reviewed By: 12355 Opinion of Evaluator: □ Rule Out □ PCP □ Alcohol □ Stimulant □ Inhalant □ Medical □ Depressant □ Hallucinogen □ Narcotic Analgesic □ Cannabis

000617	:			
	DRUG INFLUENCE EVALUATION	Page_2_of_2		
LOG NO.	DRE: Officer Mel Poff	ARRESTEE: Brian Schafer		
1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.				
1. LOCATION: Examination of Brian Schafer took place in the DRE room, HPD				
2. WITNESS: Mr. John McKay (Texas DECP Cordinator)				
3. BREATH TEST: Writer administered breath test to Schafer the result was 0.00%.				
4. NOTIFICATION / INTERVIEW of ARRESTING OFFICER: Writer was the arresting officer.				
5. INITIAL OBSERVATIONS: Writer observed subject seated in the drivers position of a blue, 1990 Oldsmobile, NJ				
registration "297 BXX". Vehicle was stationary in the driving lane of Easton Ave., at the intersection with				
West St. The traffic light was green and the other vehicles had to pull out and around subject's vehicle.				
6. MEDICAL PROBL	ÆMS: None noted or stated			
7. PSYCHOPHYSICA	L TESTS: Romberg Balance: Subject swa	ayed approximately 2" in a circular motion and		
estimated 46 seconds as 30 seconds. Walk and Turn: Subject lost his balance during the instructions, missed heel to				
toe, stopped walking,	toe, stopped walking, stepped off the line, turned backwards, and returned taking ten (10) steps.			
One Leg Stand: Subj	ject raised his arms,put his foot down, and s	wayed, Finger to Nose: Subject missed tip of his		
nose, and had very ri	igid arm movements.			
8. CLINICAL INDICA	ATORS: Subject had HGN, Vertical Nystag	mus and Lack of Convergence. His pulse, and blood		
pressure were above	the normal range.			
9. SIGNS of INGEST	ION: Subject's breath had a strong chemical	l odor and a red coating on the tongue.		
10. STATEMENTS: Subject stated he regularly takes Valium for stress. He further stated "I don't do anything else."				
11. OPINION of EVALUATOR: In my opinion Brian Schafer is under the influence of				
and unable to opera	ite a vehicle safely.			
12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a urine sample.				
13. MISCELLANEOUS:				

**Drug Influence Evaluation** Evaluator 4000618 and, R.C. Crash: **≥**None Kochubka, D.

**stee's Name (Last, First, MI) □ Property □ Fatal □ Injury DOB Sex Arresting Officer (Name, ID No.) Johnson, Steve 4-15-1965 M Breath Results 0.00 Refused Chem 732 MPOC Chemical Test □ Urine → Blood 3/18/2000 2030
Miranda Warning Given: Yes No USCP
What have you eaten today? Instrument# □ Refused Have you been drinking? When? How much? Time of last drink? Some toast, this morning Coffee wlong? Are you sick or injured? Tyes ANO By: Kochubka. D. N/A When did you last sleep? How long? Are you diabetic or epileptic? 8 hours Do you have any physical defects? I Yes A No Are you under the care of Are you taking any medication or drugs? 

Yes You Attitude Passive - Cooperative Breath Holitosis Jushed blank □ Reddened Conjunctiva Tracking: Speech Blindness: Slow, lowand raspy
Corrective Lens: Prone Normal □ Bloodshot □ Watery None □LEye □REye 🗷 Equal 🗅 Unequal Corrective Lens: Pupil Size: 🚜 Equal Able to follow stimulus: FYes □ No Evelids: □ Ûnequal (explain) □ Glasses □ Contacts, if so □ Hard □ Soft □ Normal *Droopy Pulse & Time Left Eye HGN Right Eye One Leg Stand , Zo38 Yes 405 Lack of Smooth Pursuit Convergence Right Eye Left Eye Yes Yes Max. Deviation ,2103 92 Angle of Onset Romberg Balance Walk and Turn Test Cannot Keep Balance Starts Too Soon Approx. R Sways While Balancing √2nd Nine Stops Walking □ Uses Arms to Balance Misses Heel-Toe Steps Off Line O Hopping Raises Arms Puts Foot Down Actual Steps Taken Describe Turn abrupt swivet Followed by staggering Internal Clock Cannot Do Test (Explain) Type of Footwear 50 Estimated At 30 Sec. Room Light Pupil Size Darkness Direct ○ Right / Left Left Eye Z. 0 2.0 Draw lines from spot touched Oral Cavity, Right Eye Reaction To Light Little or none VISIBLE. Rebound Dilation Hippus □ Yes **y**≰No Numerous puncture wounds NUMBER ARM WITH SCALS Scartissur Temp Muscle Tone: KNormal ☐ Flaccid ☐ Rigid Attach Photos Of Fresh Puncture Marks Comments: Medicine or Drug Have You Been Using? How Much? Time of Use? Where Were The Drugs Used? (Location) I didn't use nothing I didnituse it. I don't do that anymore. Mar Signature (Include Rank) Eval Start Time Z035 Time DRE Notified 2020 2010 ID No. 102 Reviewed By: Opinion of Evaluator: □ Stimulant а РСР □ Rule Out □ Alcohol □ Inhalant □ Medical □ Depressant □ Hallucinogen □ Narcotic Analgesic □ Cannabis

000619		· · · · · · · · · · · · · · · · · · ·	
· -	DRUG INFLUENCE EVALUATION	Page_2_of_2	
LOG NO.	DRE: Sgt. Richard Studdard	ARRESTEE: Steve Johnson	
1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.			
1. LOCATION: Exami	nation of Steve Johnson took place in the Dl	RE room, US Capitol Police HDQT.	
2. WITNESS: Arresting	Officer D. Kochubka, MPDC and Officer	G. Bird USCP	
3. BREATH TEST: O	officer D. Kochubka administered breath te	st to Johnson the result was 0.00%.	
4. NOTIFICATION /	INTERVIEW of ARRESTING OFFICER	: Writer was on duty at USCP HDQTs administering	
the DRE knowledge	examination when notified that Officer Kocl	nubka was in route with a "drugee". Officer	
Kochubka stated he l	and observed the subject walking eastbound	on East Capitol St., staggering and stumbling. He	
appeared dazed conf	used and mumbling softly. He further stated	l that the subject was wearing only shorts, a tee shirt,	
and w as barefoot. Th	ne temperature at the time was approximatel	y 34' F. No odor of alcoholic beverage was detected.	
5. INITIAL OBSERVA	ATIONS: Writer observed subject as he was	being brought into the building. He repeatedly	
staggered, stumbled, exhibited a blank stare and appeared to be unaware of his surroundings.			
6. MEDICAL PROBL	EMS: None noted or stated		
7. PSYCHOPHYSICA	L TESTS: Romberg Balance: Subject swa	yed approximately 3" side to side and estimated 50	
seconds as 30 second	s. Walk and Turn: Subject lost his balance d	uring the instructions, stepped off the line, stopped	
walking, repeatedly r	nissed heel to toe, and raised his arms for ba	lance. One Leg Stand: Subject raised his arms, put	
his foot down, swaye	ed, and raised his arms for balance. Finger	to Nose: Subject had to be reminded several times	
to keep his eyes closed, and consistently missed the tip of the nose.			
8. CLINICAL INDICA	ATORS: Subject had HGN, Vertical Nystag	mus and Lack of Convergence. His pulse was above	
the normal range, an	d his blood pressure and temperature were v	vithin the normal range. Pupils were constricted.	
9. SIGNS of INGESTION: Subject's had numerous scars resembling track marks on both arms, and had a fresh			
oozing puncture wound on the right arm.			
10. STATEMENTS: Subject stated, "No I didn't use anything", "I didn't use it " and "I don't do that anymore"			
11. OPINION of EVAL	LUATOR: In my opinion Steve Johnson is	under the influence of	
and unable to opera	te a vehicle safely.		
12. TOXICOLOGICA	L SAMPLE: Subject agreed to provide a bl	ood sample.	
13. MISCELLANEOUS:			
· ·			

**Drug Influence Evaluation** Rolling Log No. 4213 Evaluator 000620 VAUGHA GATES 8092 Evaluator 445 XVIII-6 Recorder/Witness Crash: □ None ズ Fatal □ Injury □ Property estee's Name (Last, First, MI) DOB Sex Race Arresting Officer (Name, ID No.) CHP OFFICER SEAN NAUL. LOLLY 10-3-75 M CAMEDON H W 12890 Date Examined/Time/Location Breath Results: □ Refused Chemical Test Urine □ Blood □ Refused Instrument# 6-10-2000 1245 CHP SACRAMENTO - OO Miranda Warning Given: □ Yes □ No What have you eaten today? When? Have you been drinking? How much? Time of last drink? NOTHING AT TOOK I NOTHING YET 1/A When did you last sleep? How long? Time now? Are you sick or injured? □ Yes ⋉No Are you diabetic or epileptic? □ Yes ⋉No ABOUT 2 DAY?" VEAH 10 SEC PAUSE 00 ... HU Do you take insulin? Do you have any physical defects? UYes XNo Are you under the care of UM...NOT YET I DON'T GO TO THE DOC. Are you taking any medication or drugs? Yes Do Attitude Coordination COOPELATIVE BUT SULL TO RESPON SUCIO ANDO SHAKEY I TOOK TYLEWOL THIS Breath mores, is a NOTHING UNUSUAL DOMING UNUSUAL Eyes: Reddened Conjunctiva Blindness: Speech Low Voice, Show Tracking: K Equal □ Unequal □ Normal □ Bloodshot XWatery None □LEye □REye AUG SOMETIMES SCURPED CLOUDY Pupil Size: DEqual RT PUPIL 2 mm

# Unequal (explain) LACKER IN ALL X None Corrective Lens: Able to follow stimulus: X Yes □ No Eyelids:  $\square$  Glasses  $\square$  Contacts, if so  $\square$  Hard  $\square$  Soft Normal □ Droopy LIGHTING COND. Pulse & Time HGN Left Eye Right Eye Vertical Nystagmus? □ Yes Vo One Leg Stand 1. 120 / 1245 Lack of Smooth Pursuit Convergence 20 NO Right Eye Left Eye 2. 120 / 1305 Max. Deviation 20 NO 8. 120 / 1345 Angle of Onset こひりに HUNE Walk and Turn Test WOULD NOT PLACE Romberg Balance Cannot Keep Balance Starts Too Soon Approx. Approx R/ Sways While Balancing 1" Nine 2nd Nine Stops Walking STATED, " THIS IS IMPUSSIBLE" STEPPED Misses Heel-Toe □ Uses Arms to Balance OFF UNE AND WOULD Steps Off Line □ Hopping Raises Arms こうしていしつ Puts Foot Down Actual Steps Taken 20 Internal Clock Describe Turn Cannot Do Test (Explain) Type of Footwear _Estimated At 30 Sec. と REPUSED TO COMPLETE GOOLK 2TOS Pupil Size Room Light Darkness Direct Nasal Area ○ Right △ Left CLEAS Left Eye 7.5 Draw lines from spot touched 5.5 5.0 Oral Cavity Right Eye 3.5 5.5 3.0 CLEAR Hippus ☐ Yes Rebound Dilation Reaction To Light **Σ**(Nο □ Yes X No NORMAL NONE OBSERVED 900° 180 Blood Pressure (CO Temp Muscle Tone: Normal □ Flaccid □ Rigid Attach Photos Of Fresh Puncture Marks Comments: Medicine or Drug Have You Been Using? How Much? Time of Use? Where Were The Drugs Used? (Location) ST TWO TYLENOL NO AUSCUED THISMORNICE NO ANSWER Date/Time of Arrest Time DRE Notified **Eval Start Time** Time Completed (6 - 10 - 2000) Member Signature (Include Rank) 1205 1345 ID No. Reviewed By: OFFICES KNOWIES Opinion of Evaluator: □ Rule Out □ Alcohol □ Stimulant п РСР □ Inhalant Medical □ Depressant □ Hallucinogen □ Narcotic Analgesic □ Cannabis

#### DRUG INFLUENCE EVALUATION

Page 2 of 2

LOG NO.

DRE: Officer Vaughn Gates

ARRESTEE: Cameron H. Sholly

- 1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR.
- 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS
- 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.
- 1. LOCATION: Examination of Cameron Sholly took place in the DRE examination room Sacramento CHP
- 2. WITNESS: Arresting Officer Officer Sean Nava, #12890
- 3. BREATH TEST: Writer administered breath test to Sholly the result was 0.00%
- 4. NOTIFICATION / INTERVIEW of ARRESTING OFFICER: I was advised via dispatch to respond to Southwest

  Division to conduct an evaluation at the request of Officer Nava. Officer Nava stated that the subject had been
  a driver of a vehicle involved in a fatal crash.
- 5. INITIAL OBSERVATIONS: Writer first observed the defendant standing next to the breath testing instrument at the rear door of Sacramento Station. He was standing upright on his own without assistance and was not swaying.
- 6. MEDICAL PROBLEMS: The defendant did state that high blood pressure runs in his family and defendant sometimes stutters uncontrollably.
- 7. PSYCHOPHYSICAL TESTS: During the instruction portions of all the divided attention tests. Defendant appeared to be confused. When asked if he understood the instructions of the test, Sholly would say "yes" or "yeah" but would still appear to be confused. I had to continually show the defendant how to perform the test, and after the defendant would perform the test, he would still appear to not have understood what he had just done. The defendant would not complete or even attempt to complete the walk and turn test. He just stated "This is impossible" and stand there staring at the line on which he had been standing. I had to physically move the defendant's right foot in front of his left foot on the line during the instruction phase, even after repeated demonstrations he didn't seem to understand. Romberg Balance: Subject estimated 15 seconds as 30 seconds. Williams exhibited non-bilateral impairment on certain divided attention tasks: for example during the finger to nose test, he correctly touched his nose with his right index finger, but missed on all three occasions with his left hand.
- 8. CLINICAL INDICATORS: Subjects pulse and systolic blood pressure were above the normal range. He was sweating heavily around the neck and chest area. Pupils were unequal (2 millimeters) in all light levels.
- 9. SIGNS of INGESTION: None were evident
- 10. STATEMENTS: Defendant stated "I do not use (stutter pause) drugs at all, I only took two Tylenol this morning.

  (long pause 10 -seconds) I don't drink that much anymore, either."
- 11. OPINION of EVALUATOR: In my opinion, Cameron Sholly is not exhibiting any symptoms of drug intoxication but was possibly exhibiting signs of mental impairment.
- 12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a urine sample

One Hour and Thirty-Five Minutes

SESSION XIX

INHALANTS

#### SESSION XIX INHALANTS

Upon successfully completing this session, the participant will be able to:

- o Explain a brief history of the Inhalant category of drugs.
- o Identify common drug names and terms associated with this category.
- o Identify common methods of administration for this category.
- o Explain the symptoms, observable signs and other effects associated with this category.
- o Explain the typical time parameters, i.e., onset and duration of effects, associated with this category.
- o State the clues that are likely to emerge when the drug evaluation and classification process is conducted for a person under the influence of this category of drugs.
- o Correctly answer the "topics for study" questions at the end of this Section.

Content Segments		<u>Lear</u>	Learning Activities	
A.	Overview of the Category	0	Instructor Led Presentations	
В.	Possible Effects	o	Review of Drug Evaluation and Classification Exemplars	
C.	Onset and Duration of Effects	0	Reading Assignments	
D.	Overdose Signs and Symptoms	o	Video Presentations (If Available)	
E.	Expected Results of the Evaluation	o	Slide Presentations	





XIX-OA&B (Objectives)



15 Minutes



XIX-1 ("Major Types of Inhalants")

#### **INHALANTS**

- A. Overview of the Category
  - 1. Inhalants are breathable chemicals that produce mind altering results.
    - a. Inhalants vary widely in terms of the chemicals involved and the specific effects produced.
    - b. Depending on the nature of the particular Inhalant, the effects produced may be similar to those of CNS Stimulants, Depressants or Hallucinogens.
  - 2. There are three major subcategories of Inhalants.
    - a. Volatile Solvents
    - b. Aerosols
    - c. Anesthetic gases
  - 3. The <u>Volatile Solvents</u> include a large number of readily available substances, none of which are intended by their manufacturers to be used as drugs.

Total Lesson Time: Approximately 95 Minutes

Session title on wall chart.

Briefly review the objectives, content and activities of this session.

#### INSTRUCTOR NOTES:

Inhalants are sometimes called "Deliriants," in that they may produce delirium. Delirium is usually a brief state characterized by incoherent excitement, confused speech, restlessness and possible hallucinations.

"Volatile" means that they evaporate easily to produce fumes.

#### Aides

#### Lesson Plan

#### **Instructor Notes**



XIX-2 ("Volatile Solvents")

- a. One widely abused Volatile Solvent is plastic cement, or "model airplane glue".
- b. Plastic cement includes the following volatile chemicals.
  - o Toluene
  - o Acetone
  - o Naphtha
  - o Aliphatic Acetates (straight-chained hydrocarbons)
  - o Hexane
  - o Cyclohexane
  - o Benzene
- c. Other frequently abused Volatile Solvents include:
  - o Gasoline
  - o Kerosene
  - o lighter fluid
  - o household cements and glues
  - o fingernail polish remover
  - o paint thinners
  - o engine degreasers
  - o typewriter correction fluid (liquid paper)
  - o paints (particularly oil or solvent based)
  - o dry cleaning fluids
  - o spray paints
- 4. Aerosols are chemicals discharged from a pressurized container by the propellant force of a compressed gas.

Ask students to name a Volatile Solvent that often is abused as a drug.

Contains Naphtha Rubber Cements contain Benzene

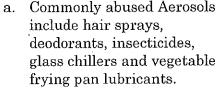
Contains Acetone

Older stocks contain Trichlorethylene.



#### Lesson Plan

#### **Instructor Notes**



- b. All of these abused Aerosols contain various hydrocarbon gases that produce drug effects.
- 5. The overwhelming majority of abusers of Volatile Solvents and Aerosols are children.
  - a. Male Inhalant abusers outnumber females
- 6. The third subcategory, Anesthetic gases.
  - a. Anesthetic gases are drugs that abolish pain.
  - b. They are used medically during surgical procedures such as childbirth, dental surgery, etc.
  - c. Anesthetic gases that sometimes are abused as Inhalants:

Nitrous Oxide

- o Ether
- o Chloroform

These substances have a long history of medical use and illicit use, e.g., Ether abuse dates to the 1790's in England, where it was taken orally. Chloroform was used in 1849 in England as a childbirth anesthetic.

Nitrous oxide has been used since 1845. It is still used in certain dental procedures.



XIX-4 ("Typical Abusers")



XIX-5 ("Anesthetic Gases") E.g., Freon, which is now available primarily in many medical Aerosols.

If available, display 35 mm slides of typically abused Aerosols.

Some reasons: These substances appear in nearly every household. They are inexpensive and readily accessible.

Adults may be more frequent users of the anesthetic gases subcategory than of the Aerosols or Volatile Solvents.

- d. Other common Inhalants in this subcategory that do not relieve pain are:
  - o Amyl Nitrite
  - o Butyl Nitrite
  - o Isobutyl nitrite and Butyl nitrite have essentially identical effects to Amyl nitrite.
- 7. Inhalants obviously are ingested by breathing, or inhaling, their fumes.
  - a. Some are ingested directly from the source.

 Some are soaked into rags, handkerchiefs or tissue papers for repeated inhalation. Nitrous Oxide is a propellent for whipped cream. Drug paraphernalia stores often sell Nitrous Oxide in cartridges that are identical to carbon dioxide containers. They are termed by users "whippets", and are allegedly sold to purchasers as devices to propel whipped cream.

Nitrites are vasodilating substances used medically to relieve angina pectoris (heart-related chest pains) and for treatment of cyanide poisoning. In angina, the nitrites work by dilating blood vessels near the heart so that more blood can reach the heart. Nitroglycerin, ordinarily not abused as an intoxicant, is also used for this purpose.

Common slang and brand names for the nitrites are: "Rush" and "Locker Room".

Examples: Amyl Nitrite and Butyl Nitrite are sold in small glass bottles or bulbs. The user simply opens the bottle and breathes in the fumes. They have been marketed in drug paraphernalia stores as room deodorizers.

Aides	Lesson Plan			Instructor Notes	
		c.	Some are placed in paper or plastic bags which the user places over the face or head. These may be placed in twist lock beverage containers.		
		d.	Some are used by breathing the fumes or vapors from balloons.	Solicit students' comments or questions concerning this overview of Inhalants.	
		e.	Some common street names that Inhalant users use are: huffing, hacking, ballooning, and glading.		
	B.	Po	ssible Effects		
10 Minutes	The effects of Inhalants vary somewhat from one substance to another.		newhat from one substance		
	2.		mmon effects of Inhalants clude:		
XIX-6 ("Effects of Inhalants")		a.	Inebriation similar to alcohol intoxication.	In fact, many of the Inhalants are classified as Depressants in medical texts. Their effects,	
		b.	Bizarre thoughts.	consequently, often mirror	
		c.	Dizziness and numbness.	Alcohol intoxication.	
		d.	Euphoria and grandiosity.		
		e.	Floating sensations.		
		f.	Distorted perceptions of space and time.		
		g.	Possible hallucinations.		
		<b>h.</b>	Nausea and excessive salivation.		
		i.	Drowsiness and weakness.		

Aides	Lesson Plan	Instructor Notes
5 Minutes	j. Light headedness.  k. Altered shapes and colors.  l. Antagonistic behavior.  m. Intense headaches.  3. Persons under the influence of Inhalants generally will appear confused and disoriented, and their speech will be slurred.  C. On-Set and Duration of Effects  1. Inhalants' effects are felt virtually immediately.  2. Duration very much depends on the particular substance.  a. Amyl Nitrite, Isobutyl Nitrite produce effects that last a few seconds up to 20 minutes.	Solicit students' questions and comments concerning possible effects of Inhalants.  Point out that the route of passage of the drugs from lungs to brain can be traveled very quickly.  Inhalation of these produces a distinct "rush" similar to that of the related substance, Nitrous Oxide.  Users claim these Nitrites enhance sexual excitement. This may occur from dilation of genital arteries (vasodilation) and relaxation of other smooth
	<ul> <li>b. The effects of nitrous oxide last 5 minutes or less.</li> <li>c. Glue, paint, gasoline and other commonly abused Inhalants produce effects that last several or more hours. (Generally 6-8 hours for most volatile solvents depending on exposure.)</li> </ul>	Point out that residue of these substances may be deposited inside the nostrils, causing the user to breathe the fumes constantly.  Solicit students' comments and questions concerning the time parameters of Inhalants.



#### 5 Minutes

- D. Overdose Signs and Symptoms
  - 1. There is a risk of death due to overdose of Inhalants.
    - a. Some Inhalants will depress the Central Nervous System to the point where respiration ceases.
    - b. Others can produce instant death from heart failure.
    - c. Overdoses of Inhalants frequently induce severe nausea and vomiting: If the user vomits while he or she is unconscious, death can result from aspiration of the vomitus.
  - 2. Death can also result indirectly, if a person places a plastic bag over the head, loses consciousness and suffocates.
  - 3. Long term abuse of Inhalants can cause permanent damage to the Central Nervous System, and greatly reduced mental and physical abilities.
  - 4. Evidence also exists of liver, kidney, bone and bone marrow damage resulting from long term Inhalant abuse.
  - 5. There is no well defined withdrawal syndrome for these substances. Physical dependence has not been documented, although habituation is common.

All solvents make the heart more sensitive to adrenaline. This sometimes causes a dangerous cardiac arrhythmia. The term "sudden sniffing death" (SSD) has been used to describe death resulting from physical exertion and the breathing of Inhalants in an enclosed, poorly ventilated space.

Solicit students' questions and comments concerning overdose signs and symptoms.

#### Aides

#### Lesson Plan

#### **Instructor Notes**



60 Minutes



XIX-7A ("SFST Evidence")

- E. Expected Results of the Evaluation.
  - 1. Observable evidence of impairment.
    - a. Standardized Field Sobriety Tests.
      - o Horizontal Gaze Nystagmus will generally be present.
      - o Vertical Gaze Nystagmus may be present.
      - o Performance on the Walk and Turn and One Leg Stand tests will be impaired.
      - o Performance on the Romberg and Finger to Nose tests will be impaired.
    - b. General indicators
      - o odor of the inhaled substance
      - o possible traces of the substance around the face and nose and on the hands or clothing
      - o bloodshot, watery eyes
      - o confused, disoriented appearance

Emphasize that, with Inhalants, there is significant variation in effects from one substance to another.

<u>Point out</u> that immediate onset of Nystagmus may be observed.

Point out that high doses (for that individual) of Inhalants may cause Vertical Gaze Nystagmus.

<u>Point out</u> that subjects will tend to take slow, deliberate steps on the Walk and Turn, and will tend to stagger.

<u>Point out</u> that subjects will tend to sway when performing this test.



XIX-7B ("General Indicators")

Aides	Lesson Plan	Instructor Notes
XIX-7C ("Eye Exam-	o muscle tone varies - flaccid or normal  o flushed face, possibly sweating  o slow, thick, slurred speech  o non-communicative  2. Evidence associated with the physiologic examinations.  a. Eye examinations  o Lack of Convergence will be present.	Speech usually clears up quickly when substance is no longer being inhaled.
inations")	o Pupil size will be normal but may be dilated.	Anesthetic gases may produce some dilation, although usually not to the extent seen with CNS Stimulants or Hallucinogens. No Inhalants produce pupillary constriction.
XIX-7D ("Vital Signs Examina- tions")	<ul> <li>o Reaction to light will be slowed.</li> <li>b. Vital signs examinations</li> <li>o blood pressure will be up or down</li> </ul>	NOTE: The Anesthetic Gases generally lower blood pressure while elevating pulse rate. The Volatile Solvents and the Aerosols usually elevate both blood pressure and pulse rate.  The lowering of blood pressure by Anesthetic Gases is due to their vasodilation effect. The heart compensates for this vasodilation by increasing its heart rate.

Aides Lesson Plan Instructor No	tes
o pulse will be up Pulse increase is due to factors, including oxygen displacement. The heat faster in order to body tissues with a surguply of oxygen.	en art may supply
o effect on body temperature may be up, down or normal.	
XIX-8 ("Inhalants Symptomatol ogy Chart")	
4. Demonstrations  a. Video tape demonstrations (if available)  b. Drug Evaluation and Classification exemplar demonstrations  Before students to the exemplars found at the Section XIX of their st manuals.  Relate the items noted exemplars to the Symptomatology chart Solicit students' comm questions concerning eresults of the evaluation.	avior/ t. e end of udent on the . ents and xpected on of
subjects under the infl Inhalants.	uciice oi

## **Session XIX**

#### Inhalants



#### Inhalants

Upon successfully completing this session, the participant will be able to:

- Explain a brief history of the Inhalant category of drugs
- Identify common drug names and terms associated with this category
- Identify common methods of administration for this category
- Explain the symptoms, observable signs, and other effects associated with this category

Drug Evaluation & Classification Training

XIX-OA

# Inhalants (continued)

- Explain the typical time parameters, (i.e., on-set and duration of effects, associated with this category)
- State the clues that are likely to emerge when the drug evaluation and classification process is conducted for a person under the influence of this category of drugs
- Correctly answer the "topics for study" questions at the end of this section

Drug Evaluation & Classification Fraining

XIX-0B

#### **Major Types of Inhalants**

- · Volatile solvents
- · Aerosols
- · Anesthetic gases

Drug Evaluation & Classification Training

XIX.

#### **Volatile Solvents**

- · Plastic cement ("model airplane glue")
- · Petroleum products
  - Gasoline
  - Kerosene
- · Lighter fluid
- Household cements and glue
- Fingernail polish remover
- · Paint thinners
- · Typewriter correction fluid
- · Paints (particularly oil or solvent based)
- · Dry cleaning fluids
- · Spray paints

Drug Evaluation & Classification Training



XIX-2

#### Aerosols

- · Hair sprays
- Deodorants
- Insecticides
- · Glass chillers
- · Frying pan lubricants



Drug Evaluation & Classification Training

XIX-3

#### **Typical Abusers of Inhalants**

- Children
- · Males outnumber females
- · Poor children are significantly overrepresented

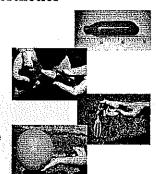
Drug Evaluation & Classification Trainin

YIY.

#### Anesthetics

- Ether
- Chloroform
- · Nitrous Oxide
- · Amyl Nitrite
- · Butyl Nitrite
- Isobutyl Nitrite

Drug Evaluation & Classification Training



#### **Effects of Inhalants**

- Inebriation similar to alcohol intoxication
- Bizarre thoughts
- Dizziness and numbness
- Euphoria and grandiosity
- Floating sensations
- Distorted perceptions of space and time
- · Possible hallucinations

Drug Evaluation & Classification Training

- Nausea and excessive salivation
- Drowsiness and weakness
- · Lightheadedness
- Altered shapes and colors
- Antagonistic behavior
- · Intense headaches

XIX-6

# **Evaluation of Suspects Under the Influence of Inhalants**

#### SFST Evidence:

- · Horizontal Gaze Nystagmus will be present
- Vertical Gaze Nystagmus present (high dose for that individual person)
- Impaired performance will be evident on Walk and Turn and One Leg Stand
- Impaired performance will be evident on Romberg and Finger to Nose

Drug Evaluation & Classification Training

XIX-7A

#### Evaluation of Suspects Under the Influence of Inhalants

#### **General Indicators:**

- · Odor of the inhaled substance
- · Possible traces of the substance around the face and nose
- · Bloodshot, watery eyes
- · Confused, disoriented appearance
- · Lack of muscle control
- · Flushed face, possibly sweating
- · Slow, thick, slurred speech
- Non-communicative

Drug Evaluation & Classification Training

XIX-7B

# **Evaluation of Suspects Under the Influence of Inhalants**

Eye Examinations:

- · Lack of Convergence will be present
- Pupil size normal*
- · Pupil reaction to light will be slow

*may be dilated

Drug Evaluation & Classification Training

XIX-7C

# **Evaluation of Suspects Under the Influence of Inhalants**

#### Vital Signs:

- Blood pressure may be up or down*
- · Pulse will be up
- Effect on body temperature will be up, down or normal

*up with volatile solvents or aerosols; down with anesthetic gases

Drug Evaluation & Classification Training

XIX-7

### Inhalants Symptomatology Chart

HGN	Present
VGN	Present (High dose for that individual)
Lack of Convergence	Present
Pupil Size	Normal*
Reaction to Light	Slow
Pulse Rate	Up
Blood Pressure	Up or down**
Temperature	Up, down, or normal
Muscle Tone	Normal or flaceid

*But may be dilated

**Up with volatile solvents or aerosols; down with anesthetic gases

Drug Evaluation & Classification Training

XIX-8



Drug Influence Evaluation				
Evaluator BUSTRUM, ROB	DRE No 4212	Rolling Log No. 131		
Recorder/Witness Bill MOPRISON	Crash: ≯None □ Fatal □ Injur			
estee's Name (Last, First, MI)	DOB Sex	Race Arresting Officer (		
Date Examined/Time/Location	<u>16-8-821 M</u>	Brooth Rocultors AND Refused	SON, BILL MONT CO.  Chemical Test • Urine Blood	
7-2-2000 3200	DPD TRAFFIC DIV	Instrument #1234	□ Refused	
		nen? Have you been drinking		
By: R. BUSTRUM H	amburger _	6 PM Just WATER		
Time now? When did you last sleep? How long	Are you sick or injured?	□ Yes 🌘 No	Are you diabetic or epileptic?	
Do you take insulin?	Do you have any physical	defects? □Yes ₽No	Are you under the care of a doctor or dentist? ☐ Yes • No	
Are you taking any medication or drugs?   Yes	No Attitude	ive but cazed	Coordination Very 200 R COULD BARELL STAND	
	Breath	- 1 d 0	Face PAINT SMEARS .	
Speech	Eyes:   Reddened Con	junctiva Blindness:	Tracking:	
Sturres/mumbles	□ Normal □ Bloodshot	·	Eye □ R Eye	
Corrective Lens:	il Size: Equal nequal (explain)	Able to follow	v stimulus: ▼Yes □ No Eyelids:  Normal □ Droopy	
Pulse & Time HGN	Left Eye Right Eye	Vertical Nystagmus? □ Yes 🎳 No	One Leg Stand 105 Tool of the leg Stand	
1. 104 / 3210 Lack of Smooth Pursui	yes yes	Convergence Right Eye Left Eye	W GOV	
2. 102 / 224 Max. Deviation	ves ves	( <del>2</del> )	0 1 40 60	
3. 104 / 2240 Angle of Onset	<u> </u>		Sur Sur Sur Sur Sur Sur Sur Sur Sur Sur	
Romberg Balance Walk and Turn Approx. Approx.	STOPPED 1	Cannot Keep Balance Starts Too Soon	L R	
		Stops Walking 1st Nin	e 2 nd Nine	
Tost STOPPED CO	IDANT STAND	Misses Heel-Toe	□ □ Uses Arms to Balance	
		Steps Off Line	□ Hopping	
Staggered Stag		Raises Arms Actual Steps Taken	D Puts Foot Down	
NEARLY FELL		-		
Internal Clock  NA Estimated At 30 Sec.  Describe Turn  NA		Cannot Do Test (Explain) UNABLE TO STAI HEEL- TOE	Type of Footwear	
	Pupil Siz	e Room Light Darkness	Direct Nasal Area DROP Paint	
1, 10(1) 1: 0"755171	Left Eye	4.0 6.5	3.5 Oral Cavity OctoR	
palmoniose	Right Eye		3.5 OF PAINT	
	Hippus	☐ Yes Rebound Dilation	Reaction To Light	
Palm OF Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched To Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touch Touched Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touch Touched Touch Touch Touched Touch Touch Touch Touched Touch Touch Touch Touched Touch Touch Touch Touched Touch Touch Touch Touched Touch Touch Touch Touched Touch Touch Touch Touched Touch Touch Touch Touched Touch Touch Touch Touched Touch Touch Touch Touched Touch Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touch Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched Touched T	N/	#No □Yes #No	NORMAL	
2		MRA THRIA	LEFT ARM PAINT SYMEARS	
	PA'	PAINT	smears det	
3 TEST Administe	rect 6 5m	Paint		
110 100 001				
Blood Pressure TO / DO Tel  Muscle Tone: Normal   Flaccid   Rigid  Comments:		Attach Photos Of I	Fresh Puncture Marks	
nat Medicine or Drug Have You Been Using?	How Much? Time of U		•	
	11A-1- 1   -	$\alpha$ $\Gamma$	Door	
IL SNIFFED a little Gold	Notmuch ABOU		PARK	
Date/Time of Arrest	me DRE Notified	Eval Start Time	Time Completed	
Date/Time of Arrest  1-2-2000  Member Signature (include Rank)	me DRE Notified  2145  Sept. ID No. 4	Eval Start Time	Time Completed 2245	

000638			
	DRUG INFLUENCE EVALUATION	Page 2_ of 2	
LOG NO.	DRE: Sgt. Rob Bustrum	ARRESTEE: Jay Graves	
5. INITIAL OBSERVATION	SS 3. BREATH TEST 4. NOTIFICATION / NS 6. MEDICAL PROBLEMS 7. PSYCHOL 10. SUSPECTS STATEMENTS 11. OPINIC		
1. LOCATION: Examina	ntion of Jay Graves took place in the DPD	Traffic Div.	
2. WITNESS: Arresting (	Officer Bill Morrison, Mont. Co.		
3. BREATH TEST: Arre	esting Officer Bill Morrison administered	breath test to Graves the result was 0.00%.	
4. NOTIFICATION / IN	TERVIEW of ARRESTING OFFICER	: Writer was contacted by radio and advised	
to return to the holding	g facility to conduct a DRE evaluation. Of	ficer Morrison stated he had arrested the subject for	
failing to obey a traffic	c control device, at Colfax and 6th Ave. S	ubject was cooperative, dazed, and unable	
to perform the SFSTs.	A can of Krylon Gold spray paint was for	und on the front seat of the subjects vehicle along	
paint soaked rags.			
5. INITIAL OBSERVAT	FIONS: Writer observed subject seated in	the DRE room, he appeared passive and dazed.	
Paint smears were visi	ble on his hands chin and upper lip.		
6. MEDICAL PROBLE	MS: None noted or stated		
7. PSYCHOPHYSICAL	TESTS: Romberg Balance: Subject una	ble to perform test, and it was terminated for his	
safety. Walk and Turn	: Subject unable to perform test, and it was	s terminated for his safety. One Leg Stand: Subject	
unable to perform test,	unable to perform test, and it was terminated for his safety. Finger to Nose: Subject was seated and used the palm		
of his hand to touch his	s nose on each attempt.		
8. CLINICAL INDICAT	ORS: Subject had HGN, and Lack of Co	nvergence. His pulse and blood pressure were	
above the normal rang	e.		
9. SIGNS of INGESTIO	ON: Subject's breath had a strong chemical	odor "like paint." There were paint smears on his	
face and hands.			
10. STATEMENTS: Sub	oject was asked "how much paint did you s	niff today?" He replied, "I sniffed a little gold - not	
to much - just a little	bit". When asked when and where he'd sr	niffed, he replied, "about 8 o'clock in the park".	
11. OPINION of EVALU	JATOR: In my opinion Jay Graves is und	ler the influence of an Inhalant and	

unable to operate a vehicle safely.

13. MISCELLANEOUS:

12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a urine sample.

	O00639 Drug Influence Evaluation						
	ZARRAGA JOSE	DRE No OG6	6 .	Rolling Log N	8-151		
	Recorder/Witness RICK LOMBARDI	Crash: □ Fatal	® Nor □ Inju		Property		
	stee's Name (Last, First, MI)	DOB O	Sex		Arresting Officer (Na	<u> </u>	N1. 10 or 05
ŀ	Date Examined/Time/Location	9-1-82	F F	Breath Results:	LOMBAY	Chemical Test	VewPort PD
	12-11-2000 2000	<tw kla<="" td=""><td>as PD</td><td>Instrument #</td><td>1,03^{0 nemoca} 1,34</td><td>□ Refuse</td><td></td></tw>	as PD	Instrument #	1,03 ^{0 nemoca} 1,34	□ Refuse	
f	Miranda Warning Given: ◆Yes □ No What	have you eaten to	1/ ~1		you been drinking?	How much?	Time of last drink?
		1e PIZZE		erschool Wit	1e Coolers		MPM
	Time now? When did you last sleep? How long?	Are you sick	· ·		□ No	Are you diabetic o □ Yes <b>\$</b> No	r epileptic?
ļ	Do you take insulin?	Do you have		ZZY ANG defects? □ Yes	d WOOLY	Are you under the	care of
						a doctor or dentist	? □Yes <b>©</b> tNo
Ì	Are you taking any medication or drugs? □ Yes PNo	Attitude COOPei	VITAS	e. Very Sin	ow to Respond	Poor ST	AGGERING
		Breath ,			SOLÎNG	Face Flus	3
ŀ	Speech		eddened Cor	njunctiva	Blindness:  None □ L E	1	Tracking:  © Equal □ Unequal
	Slow Sherred AND LOW		vidioousnot	D Watery			
		Size: Equal qual (explain)			Atte to follow s	timulus: 🏕 Yes □ No	Eyelids:
	Pulse & Time HGN	Left Eye I	Right Eye	Vertical Nystagm	us? □ Yes 💋 No	One Leg Sta	
	1. 100 / 2010 Lack of Smooth Pursuit	Yes	Yes	Convergence Right Eye	Left Eye	O J	
	2. 100 / 3024 Max. Deviation	Yes 1	YES	Right Eye	(Lett Eye		
	3. 100 / 2036 Angle of Onset	350	35°				eb 18
[	Romberg Balance Walk and Turn Te Approx. Approx.	st			Keep Balance Too Soon	L R	6
7	34 34 3		PERE	Stops W	1" Nine Valking		Sways While Balancing
\	Y Y Y TEST &	HOPPED	5T	Misses I	Heel-Toe	<b>1</b>	Jses Arms to Balance
	CIRCULATION OF THE	مشهشه	•	Stone O	ff Line	0 01	Hopping
	COAK POR	Litelan	GTA!	Raises Actual S	Arms Steps Taken	<b>6</b> 0 1	Puts Foot Down
	Internal Clock Describe Turn	<u> </u>	5	Cannot Do Tes	t (Explain) TPST	Type of Foo	twear
	Estimated At 30 Sec.			Stopped	When hear	ely Tenn	is Shoes
Ì	○ Right △ Left		Pupil Si	ze Room Light		Direct Nasal Area	RUNNY NOSE
	_		Left Eye	5.0	6,5 1	( II	of Gasoline
	Draw lines from spot touched	L	Right Ey		6.5		
	N (10)	$\mathcal{D}$	Hippus		ound Dilation Re	action To Light	ASOLIANE
	14	N)		No □ Ye	es <b>&amp;</b> No	NORMA	K
		J		RIGHT ARM		Visit All	
	2	<u> </u>			' n	A CONTRACTOR OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF TH	
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	\$ 6	<b>8 A</b>		95			$\rightarrow$
ļ	Blood Pressure 146 / 104 Temp	<u>98.8</u> •	-				
	Blood Pressure / / / Temp  Muscle Tone:   Normal   Flaccid   Rigid	<u>40.50</u> °	- F				,
	Comments:					esh Puncture Mar	· ks
(		How Much?	Time of	Use? Who	ere Were The Drugs U	1 7, 1	+10:01
	L dudn't Sniff ANYTH	DRE Notified	12 00	SPS L	didn't	Time Comple	Wight.
	Member Signature (Include Rays)	DRE Notified	ID No.	Pay	2000	10/10	40
	Member Signature (Include Renk)  Opinion of Evaluator:   Rule Out	Alcohol	O Stim	411	Lew	I Inhalant	
		Depressant			arcotic Analgesic	□ Cannabis	

## 000640 DRUG INFLUENCE EVALUATION Page 2 of 2 LOG NO. DRE: Jose Zarraga ARRESTEE: C. Mashburn 1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC. 1. LOCATION: Examination of C. Mashburn took place in the DRE room, Central Testing Unit, Stockton P.D. 2. WITNESS: Arnie Trotter, California Office of Traffic Safety 3. BREATH TEST: Writer administered breath test to Mashburn the result was 0.03%. 4. NOTIFICATION / INTERVIEW of ARRESTING OFFICER: Writer was the arresting officer. 5. INITIAL OBSERVATIONS: Writer observed subject walking northbound in the northbound lane of traffic on State St. Vehicular traffic was moderate to heavy, and oncoming vehicles were forced to swerve to avoid her. She was staggering, stumbling, and reeling as she walked. 6. MEDICAL PROBLEMS: None noted or stated 7. PSYCHOPHYSICAL TESTS: Romberg Balance: Subject swayed approximately 3" in a circular manner, nearly fell and estimated 19 seconds as 30 seconds. Walk and Turn: Subject lost his balance during the instructions, staggered and nearly fell. The test was terminated for subject's safety. One Leg Stand: Test was terminated for the subject's safety. Finger to Nose: Subject was seated, and missed the tip of her nose each time. On #5 and #6 subject used the wrong finger. 8. CLINICAL INDICATORS: Subject had HGN, and Lack of Convergence. Her pulse and blood pressure were above the normal range. 9. SIGNS of INGESTION: Subject's breath had a strong odor of gasoline. 10. STATEMENTS: Subject was asked "where did you sniff the gasoline?" She replied, "I didn't sniff anything, I don't do gas." Subject was then told that there was an odor of gasoline on her breath and asked "what time did you sniff the gas?" She replied, "I didn't do it tonight." 11. OPINION of EVALUATOR: In my opinion C. Mashburn is under the influence of Alcohol and an Inhalant and unable to operate a vehicle safely. 12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a urine sample. 13. MISCELLANEOUS:

Sixty Minutes

## SESSION XX

PRACTICE: VITAL SIGNS EXAMINATIONS

#### SESSION XX PRACTICE: VITAL SIGNS EXAMINATIONS

Upon successfully completing this session, the participants will be able to:

- o Conduct examinations of pulse and blood pressure.
- o Articulate the vital signs examination procedures.
- o Document the results of the vital signs examinations.

# Content Segments Learning Activities A. Procedures For This Session o Instructor Led Presentations B. Pulse Measurements o Students Hands On Practice C. Blood Pressure Measurements o Instructor Led Coaching D. Session Wrap Up o Student Led Coaching





XX-O (Objectives)



10 Minutes

PRACTICE: VITAL SIGNS EXAMINATIONS

- A. Procedures For This Session
  - 1. Participants will work in three or four member teams.
    - a. At any given time, one member of the team will be engaged in conducting and recording vital signs examinations of another member.
    - b. The remaining member(s) will help coach and critique the student who is conducting the examinations.
    - Students will take turns serving as test administra-tor, test subject and coach.
  - 2. Teams initially will practice taking one another's <u>pulse</u>.

Total Lesson Time: Approximately 60 Minutes

Point out "Practice Sessions" wallchart.

Briefly review the objectives, content and activities of this session.

REFER TO CHAPTER VII IF THERE ARE ANY QUES-TIONS ON VITAL SIGNS.

Make team assignments.

Emphasize that students can help each other learn by pointing out errors of omission or commission.

Point out that the student who is "coaching" should simultaneously take the subject's pulse along with the test administrator. Example: administrator can take pulse at subject's left wrist, coach can take it at subject's right wrist.

Aides	ides Lesson Plan Instr	
		Then, the administrator and coach can compare the measurements they obtain.  Demonstrate this, using a
		student subject and two instructors.
	3. Teams subsequently will practice taking one another's blood pressure.	NOTE: If specially designed training stethoscopes are available, the student coach can "listen in" on the blood pressure measurements being taken by the student administrator.
	4. Students will record their measurements, using the Vital Signs Examination Data Sheet.	Hand out copies of the Vital Signs Examination Data Sheet to each student.
		Solicit students' questions concerning procedures for this practice session.
20 Minutes	B. Pulse Measurements	Monitor teams and coach students as necessary and appropriate.
		Terminate this segment after 20 minutes, or after each student has administered a pulse measurement to each of their team members (whichever comes first).
25 Minutes	C. Blood Pressure Measurements	Monitor teams and coach students as necessary and appropriate.
		If a training Stethoscope is available, "listen in" on occasional blood pressure measurements to verify that the students are taking accurate measurements.

Aides	Lesson Plan	Instructor Notes
5 Minutes	D. Session Wrap up	Terminate this segment after 25 minutes, or after each student has measured the blood pressure of each member of their team (whichever comes first).  Offer appropriate comments and observations about the students' performance.  Solicit students' comments concerning the practice session.
	·	

## VITAL SIGNS EXAMINATIONS DATA SHEET

EXAMINER'S NAME	
DATE / /	
PULSE MEASUREMENTS	BLOOD PRESSURE MEASUREMENTS
SUBJECT'S NAME	SUBJECT'S NAME
TIME	TIME
PULSE POINT USED	SYSTOLIC
BEATS PER MINUTES	DIASTOLIC
SUBJECT'S NAME	SUBJECT'S NAME
TIME	TIME
PULSE POINT USED	SYSTOLIC
BEATS PER MINUTES	DIASTOLIC
SUBJECT'S NAME	SUBJECT'S NAME
TIME	TIME
PULSE POINT USED	SYSTOLIC
BEATS PER MINUTES	DIASTOLIC

## **Session XX**

Practice: Vital Signs Examinations



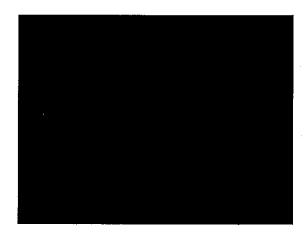
## Practice: Vital Signs Examinations

Upon successfully completing this session, the participants will be able to:

- Conduct examinations of pulse and blood pressure
- · Articulate the vital signs examination procedures
- Document the results of the vital signs examinations

Drug Evaluation & Classification Tenining

VV #



One Hour and Twenty-Five Minutes

SESSION XXI CANNABIS

#### SESSION XXI CANNABIS

Upon successfully completing this session, the participant will be able to:

- o Explain a brief history of Cannabis.
- o Identify common names and terms associated with Cannabis.
- o Identify common methods of administration for Cannabis.
- o Explain the symptoms, observable signs and other effects associated with Cannabis.
- o Explain the typical time parameters, i.e., onset and duration of effects, associated with Cannabis.
- O State the clues that are likely to emerge when the Drug Evaluation and Classification Process is conducted for a person under the influence of Cannabis.
- o Correctly answer the "topics for study" questions at the end of this Section.

Content Segments		Learning Activities	
A	Overview of the Category	0	Instructor Led Presentations
В.	Possible Effects	0	Review of Drug Evaluation and Classification Exemplars
C.	On-Set and Duration of Effects	0	Reading Assignments
D.	Overdose Signs and Symptoms	0	Video Presentations (If Available)
E.	Expected Results of the Evaluation	O	Slide Presentations

Aides	Lesson Plan	Instructor Notes
, ii .	CANNABIS	Total Lesson Time: Approximately 85 Minutes
Language and the second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second		Session title on wall chart.
XXI-OA&B		Briefly review the objectives, content and activities of this session.
(Objectives)		
	A. Overview of the Category	
10 Minutes	"Cannabis" is a category of drugs derived primarily from various species of Cannabis plants, such as Cannabis Sativa	If available, display 35 mm slides of Cannabis plants, leaves, flowers, etc.
	and Cannabis Indica.	INSTRUCTORS NOTE: Some jurisdictions as well as botan-
	a. Cannabis grows readily throughout the temperate zones of the world	ists don't recognize Cannabis Indica as a separate plant species.
, ²³ 3	. b. It has been cultivated for centuries.	Example: At the first permanent English settlement in America, Jamestown, VA, where it was grown to produce hemp.
	2. The primary psychoactive ingredient in Cannabis is Delta-9 Tetrahydrocannabinol.	Point out: "Δ- 9 THC" on wall chart.
	a. THC is found principally in the leaves and flowers of the plant rather than in the stem or branches.	18-20% in a Northern California Study
	b. Different varieties of the Cannabis have different concentrations of THC.	28-30% - Indoor Cultivated 35-37% Long Beach, CA Study in 1988-89



- c. One variety that has a relatively high concentration of THC is Sinsemilla, which is the unfertilized female Cannabis Sativa plant.
- 3. There are four principal forms of Cannabis.
  - a. Marijuana The dried leaves of the plant.
  - b. Hashish Basically a concentrated version of Marijuana, produced by crushing and boiling the leaves and allowing them to dry into a semi-solid mass.
  - c. <u>Hashish Oil</u> Also known as "Hash Oil", a liquid extracted from Hashish.
  - d. <u>Marinol</u> (or Dronabinol) A synthetic form of THC.
     This is a prescriptive drug used to inhibit vomiting. It is prescribed for certain cancer patients undergoing chemotherapy.

Nabilone - A synthetic form of THC and is used as an anti-vomiting agent.

- 4. Cannabis has some limited medical applications.
  - a. It lowers intraocular pressure, which can be helpful for Glaucoma patients.

Explanatory note: "Sinsemilla" is a Spanish derivative of the latin expression "sine semina" meaning "without seed".

Show Slides - of Special Types

"Dronabinol" is the generic, or chemical name for the synthetic THC. "Marinol" is the trade name used by the Roxane Laboratories, Inc.

"Intraocular": within the eyeball.

Cannabis lowers the intraocular pressure by dilating in size the blood vessels of the eyes (more size- less pressure).

Aides	Lesson Plan	Instructor Notes
		This causes reddening of the conjunctive.
	b. It suppresses nausea, and sometimes is recommended for cancer patients to relieve the nausea accompanying chemotherapy.	
	c. <u>Cannabidiol</u> , a non- psychoactive ingredient found in Cannabis, is used in treating Epilepsy; it helps to inhibit seizures.	
	d. Cannabis has also had some limited medical applications as:  o an appetite enhancer for victims of Anorexia Nervosa;	Marijuana has been legalized for medical treatment in some states including Arizona, Oregon, and California
	o a muscle relaxant;	
	o a tumor growth retardant.	
•	5. Marijuana usually is smoked.	
	6. Marijuana, Hashish and Hash oil also can be ingested orally, for example, baked in cookies or brownies and eaten.	
	7. In controlled studies, passive	·

inhalation of Marijuana smoke has resulted in behavioral effects as well as a measurable amount in toxicology samples.

Study does not address

quantitative amount of

physical impairment.

Solicit students' comments and questions concerning this overview of Cannabis.



#### 5 Minutes

B. Possible Effects

- 1. One major effect of Marijuana is that it appears to interfere with a person's ability to pay attention.
  - a. People under the influence of Marijuana simply seem not to pay attention, or to have very brief attention spans.
  - b. In particular, they do not divide their attention very successfully.
  - c. This can make them very unsafe drivers, since driving requires the ability to divide attention among many simultaneous tasks, i.e.
    - o steering
    - o operating the accelerator
    - o signaling
    - o observing other traffic
    - o recognizing traffic control devices
    - o shifting
  - d. People under the influence of Marijuana may attend to one or a few of these driving tasks, but simply ignore the other tasks.

<u>Clarification</u>: They have a difficult time dealing with more than one or two tasks at once.

<u>Ask</u> students: "What are some of the things that drivers have to do simultaneously?"

Loss of depth perception would be demonstrated by stopping improperly. Short attention span would be indicated by erratic speeds, failing to maintain a single lane and stopping for a red light then continuing on.

•	٠	-		
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#### Lesson Plan

#### **Instructor Notes**

- e. Because Marijuana impairs attention, Standardized Field Sobriety Tests like Walk and Turn and One Leg Stand are excellent tools for recognizing people under the influence of Marijuana.
- Remind students that WAT and OLS are <u>divided attention</u> Standardized Field Sobriety Tests.
- 2. Other effects of Marijuana.
  - a. Diminished inhibitions
  - b. Impaired perception of time and space
  - c. Disorientation
  - d. Body tremors
  - e. Eyelid tremors

- <u>Point out</u> that this may become evident when the suspect attempts to estimate the passage of 30 seconds when performing the Romberg test.
- Solicit students' comments or questions concerning possible effects of Marijuana.

5 Minutes



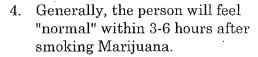
("On-set and Duration")

- C. Onset and Duration of Effects
  - Persons begin to feel and exhibit the effects within 8-9 seconds after smoking Marijuana.
  - 2. The effects reach their peak within 10-30 minutes.
  - 3. Depending on the amount smoked and on the concentration of THC in the Marijuana, the person will continue to feel and exhibit the effects for 2 3 hours.

NOTE: A 1985 Stanford University study shows pilots have difficulty in holding patterns and in lining up with runways for up to 24 hours after using Marijuana.

#### Lesson Plan

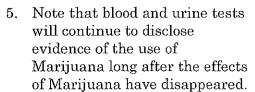
#### **Instructor Notes**



- The user may be impaired long after the euphoric feelings have ceased.
- Marijuana impaired performance at .25, 4, 8, 24 hours after smoking. While 7 of the 9 pilots showed some degree of impaired at 24 hours after smoking Cannabis, only one reported any awareness of the drugs effects.

In 1990 - a second Stanford

University Study shows:



Source Marijuana Alert, Peggy Mann (Bibliography)

NIDA Study, "Blood Brain Barrier"

a. Blood tests may disclose
 Marijuana use for at least 3
 days after smoking.

b. Urine tests may indicate the presence of metabolites of THC for a month or more.

Solicit students' comments and questions concerning onset and duration factors.

- c. There are two important metabolites, or chemical by-products of THC.
  - Hydroxy THC, which causes the user to feel euphoric.
  - o <u>Carboxy THC</u>, there is no evidence at this time that it is psychoactive.
- d. Hydroxy THC usually is eliminated from the blood plasma within six hours.
- e. Carboxy THC may be found in the blood plasma for several days following Marijuana use.





Write "Hydroxy THC: Causes Impairment and Euphoria" on

the chalkboard or flip chart.

Aides		Lesson Plan	Instructor Notes
	6.	Cannabis is a fat soluble (i.e., it dissolves easily into fatty tissue); therefore, it can remain for long periods in the brain tissue, which is about one-third fat.	
	7.	Cannabis principally is eliminated from the body in feces and urine.	
	D.	Overdose Signs and Symptoms	
5 Minutes	1.	Excessive or long term use of Marijuana can have very undesirable consequences.	
	2.	Marijuana has been observed to produce sharp personality changes, especially in adolescent users.	
	3.	It can create paranoia and possible psychosis.	
	4.	Long term effects include:	Ask students: "Is there danger of death from Cannabis overdose?"
		a. Lung damage	Answer: It is not likely that there is a <u>direct</u> risk of death
		b. Chronic Bronchitis	from an overdose. However, persons impaired by Cannabis
•		c. Lowering of Testosterone (male sex hormone)	may <u>behave</u> in foolishly dangerous ways, and become injured or killed as a result.
		d. Possible birth defects, still births and infant deaths	•
		e. Acute anxiety attacks	
		f. Chronic reduction of attention span	

Aides	Lesson Plan	Instructor Notes
60 Minutes	g. Increased sensitivity  h. Research indicates that life threatening overdoses rarely if ever occur.  i. Withdrawal - is similar to alcohol dependence withdrawal.  j. Physical dependence can occur with chronic use.  E. Expected Results of the Evaluation  1. Observable evidence of impairment	Solicit students' questions concerning signs and symptoms of Cannabis overdose.
XXI-3A ("SFST Evidence")	a. Standardized Field Sobriety Tests  o neither Horizontal nor Vertical Gaze Nystag- mus will be present.  o performance on the Walk and Turn and One Leg Stand tests will be impaired.	But remind students that Marijuana users often drink alcohol in conjunction with their smoking, and that others often lace their Marijuana with PCP. Either combination would cause Nystagmus.  Point out that, with suspects under the influence of Marijuana, poor performance on these tests usually will result principally from their inability to divide attention, and less so from impaired
	o performance on the Romberg and Finger to Nose tests will be	Remind students to be especially alert for evidence of the suspect's distorted

perception of time when performing the Romberg test.

impaired.

#### **Aides**

#### Lesson Plan

#### **Instructor Notes**



XXI-3B ("General Indicators")

- b. General indicators:
  - o odor of burnt Marijuana on suspect's breath, clothing, etc.
  - o marked reddening of the Conjunctiva (white part of the eyeball)

- c. Marijuana debris (leaves, seeds, etc.) in mouth or on clothing:
  - o body tremors
  - o disorientation
  - o relaxed inhibitions
  - o muscle tone is normal
- 2. Evidence associated with the physiologic examinations.
  - a. Eye examinations:
    - o Lack of Convergence generally will be evident.
    - o pupil size generally will be dilated or possibly normal.
    - o pupil reaction to light will be normal.

<u>NOTE</u>: Odor of Marijuana is similar to odor of burnt rope.

Properly called Conjunctival Injection.

This should not be confused with conjunctivitis which is a disease of the eye. The vasodilation is the primary cause of the reddening of the eyes not the Cannabis smoke.

Visine causes vaso-constriction in the eyes and is often used to reduce the reddening.



The content and potency could effect pupil size. The higher THC content will increase the likelihood of pupil dilation. However, Cannabis does not cause pupil constriction.

Α.	•	_1	_	_
-	1	a	ω	•

#### Lesson Plan

#### **Instructor Notes**

o DREs report a
phenomenon termed
"Rebound Dilation" in
suspects under the
influence of Marijuana.

Government grown Cannabis has a low THC levels. Studies using it tends to show a normal range of pupil size.

Clarification: "Rebound dilation" is a period of constriction followed by dilation with a change equal to or greater than 2 mm the final size determination being estimated at the end of a 15-second time period in which the light from the penlight is directed into the eye.

NOTE HOWEVER that this phenomenon has not been systematically investigated in controlled research.

Draw an eyeball on a balloon and squeeze it to demonstrate the difference between Hippus and Rebound.

NOTE: Remind students that the final size determination being estimated at the end of the 15-second time period in which the light from the penlight is directed into the eye. Caution should be used by the officer so as not to move the light beam or allow the bulb to change in light intensity.

Vasodilation - allows for greater blood flow but an increase in the amount of heat lost.



b. Vital signs examinations:

- o blood pressure generally will be up
- o pulse generally will be up
- o body temperature will be normal

Aides	Lesson Plan	Instructor Notes			
XXI-4 ("Cannabis Symptomatol ogy Chart")	3. Summary				
	4. Demonstrations				
	a. Video tape demonstrations (if available)	Show video tape of suspect(s) under the influence of Cannabis. Relate behavior/observations to the symptomatology chart.			
	b. Drug Evaluation and Classification exemplar demonstrations.	Refer students to the exemplars found at the end of Section XXI of their student manuals.			
		Solicit students' comments and questions concerning expected results of the evaluation.			
	• · · · · · · · · · · · · · · · · · · ·				

## **Session XXI**

#### Cannabis





#### **Cannabis**

Upon successfully completing this session, the participant will be able to:

- · Explain a brief history of Cannabis
- Identify common names and terms associated with Cannabis
- Identify common methods of administration for Cannabis
- Explain the symptoms, observable signs and other effects associated with Cannabis

Drug Evaluation & Clussification Training

XXI-0

# Cannabis (continued)

- Explain the typical time parameters, i.e., on-set and duration of effects, associated with Cannabis
- State the clues that are likely to emerge when the drug evaluation and classification process is conducted for a person under the influence of Cannabis
- Correctly answer the "topics for study" questions at the end of this section

Drug Evaluation & Classification Training

XXI-0E

#### Forms of Cannabis



Marijuana



Hashish oil

Drug Evaluation & Classification Training



Hashish



Marinol

XXII-

# On-set and Duration of Marijuana's Effects







- · 8-9 seconds User begins to feel and exhibit effects
- · 10-30 minutes Peak effects are reached
- · 2-3 hours User continues to feel and exhibit effects
- 3-6 hours User feels "normal"

Note: for days later, blood and urine tests will continue to disclose evidence of marijuana use

Drug Evaluation & Classification Training

XXI-2

#### **Metabolites of THC**

- Hydroxy THC
  - Causes impairment and euphoria
- · Carboxy THC

Causes impairment but no euphoria

Drug Evaluation & Classification Training

XXI-3

#### Evaluation of Suspects Under the Influence of Cannabis

#### SFST Evidence:

- · HGN or VGN none
- Impaired performance will be evident on Walk and Turn and One Leg Stand
- Impaired performance will be evident on Romberg and Finger to Nose

Drug Evaluation & Classification Training

XXI.3

# **Evaluation of Suspects Under the Influence of Cannabis**

#### General Indicators:

- · Odor of marijuana
- · Marked reddening of conjunctiva
- · Marijuana debris in mouth
- · Body tremors
- Disorientation
- · Relaxed inhibitions

Dong Evaluation & Classification Turking

xxi.a

# **Evaluation of Suspects Under the Influence of Cannabis**

#### Eye Examinations:

- · Lack of Convergence present
- · Pupil size will be dilated*
- · Pupil reaction to light will be normal

*possibly normal

Drug Evaluation & Classification Training

XXI-3C

#### Evaluation of Suspects Under the Influence of Cannabis

#### Vital Signs:

- · Blood pressure up
- · Pulse up
- · Body temperature normal

Drug Evaluation & Classification Training

XXI-30

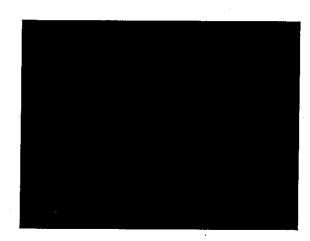
## Cannabis Symptomatology Chart

HGN	None
VGN	None
Lack of Convergence	Present
Pupil Size	Dilated*
Reaction to Light	Normal
Pulse Rate	Up
Blood Pressure	Up
Temperature	Normal
Muscle Tone	Normal

* Or possibly normal

Drug Evaluation & Classification Trainin

XXI.4



	O00663 Drug Influence Evaluation							
	Evaluator CONRAD MURRAY	DRE No //Z		Rolling Log No	5-025			
	Recorder/Witness Dave	Crash: □ Fatal	> None □ Injury	o Pr	coperty			
(	Stee's Name (Last, First, MI)	^{DOB} 5-23-46	) Sex		rresting Officer (N BURSTE)		)9 ISP	
	Date Examined/Time/Location	KNICTU C	Br In:	eath Results: 0.0 strument # 123	<u> </u>	Chemical Test	<b>p</b> Urine □ Blood d	
	Miranda Warning Given: Se Yes No What By: M (101) PQ	have you eaten tod	ay? When Arou	Have yo	ou been drinking?	How much?	Time of last drink?	
	Time now? When did you last sleep? How long?	Are you sick or	<u>5f</u>		ung AAN	Are you diabetic o	r epileptic?	
	Do you take insulin?	Do you have an	y physical def	reet Greets? 1 Yes	eat.	Are you under the	ARE YOU?	
	NO And I Con't take a lot of Si- Are you taking any medication or drugs? Yes BNO		Arnold	Δ.		a doctor or dentist		
	Drugs! Hell No!	Breath A.O.	<u>rous but</u>	•	<u>locperative</u>	Pace [	several lines	
	Speech	Eyes: □ Red	dened Conjun loodshot □ V		Blindness:	<u>    ⊢luishe/</u>  ve   □ R Eve	Tracking:  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Object  Obj	
	Corrective Lens: None Pupil	Size: 🕶 Equal		<b>y</b>		stimulus: <b>&amp;</b> Yes □ No	Eyelids:	
		qual (explain)	1. 73		D = W A N	One Low Ste	Mormal Droopy	
	Pulse & Time  1. 106 / 22 10  Lack of Smooth Pursuit			tical Nystagmus	er les prino	One Leg Sta	\$ 105 ped	
	2. 100 1227 Max. Deviation	NO	ND	Right Eye	Left Eye		100	
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	NA Estimated At 30 Sec.				minate o	Y Fell Type of Food	FERS	
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	Draw lines from spot touched	J	Left Eye Right Eye	5.5		)-2.5 Oral Cavity	1R	
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}	Member Signature (Medide Rank)	<u> </u>	^{ID No.} 466	Dames	200 ved By: 5	10 for 1	- <b>3</b>	
Ī	Opinion of Evaluator:   © Rule Out	Alcohol Depressant	□ Stimulant □ Hallucino	□ PCP	otic Analgesic	□ Inhalant \$Cannabis		

0664						
	DRUG INFLUENCE EVALUATION	Page 2 of 2				
LOG NO.	DRE: Murray Conrad	ARRESTEE: Ken Clark				
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / ONS 6. MEDICAL PROBLEMS 7. PSYCHOI N 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS				
1. LOCATION: Examin	ination of Ken Clark took place in the DRE r	room, Marion County Jail.				
2. WITNESS: Arresting	g Officer: Trooper David Bursten, Indiana St	tate Police				
3. BREATH TEST: Ar	rresting officer administered breath test to C	Clark the result was 0.00%.				
4. NOTIFICATION / J	INTERVIEW of ARRESTING OFFICEF	R: Writer was contacted by radio and advised				
to return to the holdi	ng facility to conduct a DRE evaluation. Tr	cooper Bursten stated he had observed the subject for				
operating a vehicle a	it a high rate of speed east bound on Purdue	Ave. and weaving around slower traffic. Subject				
seemed unconcerned	d about being stopped and readily admitted	driving fast. Subject stated, "I'm just out				
to enjoy myself toni	ght!"					
5. INITIAL OBSERVA	ATIONS: Writer observed subject seated in	the breathalyzer room and was laughing loudly and				
repeatedly saying "T	The machine says I'm not drunk." There was	s also reddening of the conjunctiva.				
6. MEDICAL PROBLI	EMS: None noted or stated					
7. PSYCHOPHYSICA	7. PSYCHOPHYSICAL TESTS: Romberg Balance: Subject unable to perform test, and it was terminated for his					
safety. Walk and Tur	n: Subject unable to perform test, and it was	s terminated for his safety. One Leg Stand: Subject				
unable to perform test	t, and it was terminated for his safety. Fing	ger to Nose: Subject was seated and missed the tip of				
his nose on each atten	mpt. Subject also exhibited eyelid tremors.	N.				
8. CLINICAL INDICA	TORS: Subject had lack of convergence, 1	pupils were dilated in near total darkness and				
rebound dilation was	s observed. Subject's pulse and blood press	sure were above the normal range.				
9. SIGNS of INGESTI	ION: Subject's breath had an odor of marijua	ana.				
10. STATEMENTS: Su	abject initially denied using any drugs. Whe	en told he looked and acted like someone who had				
smoked marijuana, l	he gigled and said, "come on, don't hassle n	me: this is bullshit." When asked how much pot he				
smoked, he replied,	"not much just a little." When asked where	e he smoked, subject paused and said, "No, I ain't				
saying no more."						
11. OPINION of EVAL	LUATOR: In my opinion Ken Clark is unde	er the influence of a Cannabis				
and unable to operate	te a vehicle safely.					
12. TOXICOLOGICAI	L SAMPLE: Subject agreed to provide a uri	ine sample.				
13. MISCELLANEOUS	S: Subject maintained a jovial and boisterous	s attitude throughout the entire evaluation				

Onness Drug Influence Evaluation							
Evaluator John, C.	lark	DRE No 1/Z	2	Rolling Log No	-0129		
Recorder/Witness Graham, G.	CHP	Crash: □ Fatal	a None □ Inju	3			
restee's Name (Last First MT		DOB	Sex		esting Officer (Na		
Peltier, Cha	rles E.	5-16-19				, G. M70	
Date Examined/Time/Location 9-11-2000 232	20 Par	ker Cen	ter	Breath Results 0.0 Instrument # /2	6 Refused 34	Chemical Test  • Refus	<b>≇</b> Urine □ Blood ed
Miranda Warning Given: *Yes	s □ No What	have you eaten t	oday? Wh	nen? Have you	been drinking?	How much?	Time of last drink?
•	t sleep? How long?	Are you sick	<u>ル ラ 内</u> or injured?	775 <i>G</i> 9€ Co U)	No No and	Are you diabetic o	2 hrs ago
Time now? When did you last I have Cast Night No icea I think Do you take insulin?	271131	Im n	of dru	nk either		don't 400	wlet mego
Do you take insulin?  I don't take a.	oyes ANo	Do you have	any physical (	ny physical defects?   Yes  No Are you under the care of a doctor or dentist?   Yes  No			care of
Are you taking any medication or	drugs? □ Yes WNo	Attitude	mpat	ient, anxi	005	Coordination L	en por
OFcoursenot		Breath 0	rative	AN ALCOR	HOLIC	Face Norm	ck, Stumbling
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Corrective Lens: MNone  Glasses Contacts, if so DE		Size: ÆEqual qual (explain)	•		Whis to ionom se	imulus: #Yes □ No	Eyelids:  ▶ Normal □ Droopy
Pulse & Time HO	in	Left Eye I	Right Eye	Vertical Nystagmus?	□ Yes □ No	One Leg St	and
	ck of Smooth Pursuit	425	4es	Convergence	Loft Pro		<b>3</b> )
	x. Deviation	4es	Yes	Right Eye	Left Eye		)   8
3. //0 /2353 An	gle of Onset	L	NONE				an Tanana
Romberg Balance Approx. Approx.	Walk and Turn Tes	st Leg	tremor	Cannot Kee Starts Too	Soon		eg Tremors
1 3" 4 3" 3" 3"				Stops Walki	1st Nine 2	od Nine	/ Sways While Balancing
	walked	very sl	owly	Misses Heel	l-Toe		Uses Arms to Balance
Steps Off Line   Hopping				Hopping			
EYELLO INECHORS		W	M	Raises Arms Actual Step	V V	$\frac{vv}{a}$	Puts Foot Down
Internal Clock	Describe Turn 5	taggereo	12	Cannot Do Test (Ex		Type of Foo	otwear
42 Estimated At 30 Sec.	Steps to	the rig	int	N	4	Runni	ng shoes
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Draw line	s from spot touched \	ſ				Brownish	
NO		+7				g on tongue	
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2	X /	$\frac{71}{}$					
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		<u></u>			isble.	110	
4 A Visible Mark							
Muscle Tone & Normal							
Comments:  Attach Photos Of Fresh Puncture Marks							
at Medicine or Drug Have You had a few glas	at Medicine or Drug Have You Been Using? . How Much? Time of Use? Smiks Where Were The Drugs Used? (Location) had a few glasses of wine that all. Cant Bs a Oh come on In not going to tell.						
Date/Time of Arrest 7-11-2000 2245			Eval Start Time 2350 Time Completed				
Member Signature (Include Rank		oh	ID No.//7	7 Reviewe		Kickande	
		Alcohol Depressant	□ Stimula □ Halluci		ic Analgesic	□ Inhalant • Cannabis	····

#### DRUG INFLUENCE EVALUATION

Page 2 of 2

LOG NO.

DRE: Sgt. Clark John

ARRESTEE: Charles E. Peltier

- 1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR.
- INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS
- SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.
- 1. LOCATION: Examination of Charles E. Peltier, took place in the DRE room, Parker Center, LAPD
- 2. WITNESS: Arresting Officer was Sgt. Gordon Graham, CHP
- 3. BREATH TEST: Sgt. Graham administered breath test to Peltier, the result was 0.06%.
- 4. NOTIFICATION / INTERVIEW of ARRESTING OFFICER: Writer was contacted by radio and advised to return to Parker Center to conduct a DRE evaluation. Sgt. Graham stated he had observed the subject traveling southbound on the San Diego Fwy. operating a vehicle with no head or tail lights. Upon stopping the vehicle, the subject stated, "hey I can see fine I don't need any f'ing lights cowboy!" Subject further stated "cute little bow tie --- you must be Little Bow Peep.
- 5. INITIAL OBSERVATIONS: Writer observed the subject seated in the breath testing room. Subject appeared anxious, impatient, and several times asked to be "let go". Generally he was polite and cooperative. His speech was slow and slurred, and he stumbled while walking,
- 6. MEDICAL PROBLEMS: None noted or stated
- 7. PSYCHOPHYSICAL TESTS: Romberg Balance: Subject swayed approximately 3" in a circular motion, and exhibited eyelid tremors, and estimated 42 seconds as 30 seconds. Walk and Turn: Subject lost his balance during the instructions, staggered while turning, raised arms and missed heel to toe.

One Leg Stand: Subject raised his arms, swayed, put his foot down, and exhibited leg tremors. Finger to Nose:

Subject missed tip of his nose five times and exhibited eyelid tremors.

- 8. CLINICAL INDICATORS: Subject's pulse and blood pressure were above the normal range. His pupils were dilated, there was lack of convergence, and HGN was present. There was also a reddening of the conjunctiva.
- 9. SIGNS of INGESTION: Subject had a brownish coloration on his tongue.
- 10. STATEMENTS: Subject admitted to drinking "a few glasses of wine" When subject was asked, "when did you smoke the marijuana?" He responded, "I guess I can't bullshit a bullshitter, can I?" "marijuana? who me?" and then laughed. When asked where he had used the marijuana, the subject replied, "oh, come on, I'm not going to tell you."
- 11. OPINION of EVALUATOR: In my opinion Charles E. Peltier is under the influence of Alcohol and Cannabis and unable to operate a vehicle safely.
- 12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a urine sample.
- 13. MISCELLANEOUS:

**Drug Influence Evaluation** Rolling Log No. 20%, Evaluator 5158 XX1-3 Crash: Recorder/Witness □ Property DOB estee's Name (Last, First, MI) Race Arresting Officer (Name, ID No.) Sex 10/5/53 JAIGHT JAMES B w FLAHAVAN, TOM CHP 13505 Date Examined/Time/Location Breath Results: □ Refused Chemical Test ▼Urine □ Blood Instrument# □ Refused 0.00 12-7-96 <u> 2300</u> AMADOR CHP OFFICE 012833A Miranda Warning Given: XYes 🗆 No What have you eaten today? Have you been drinking? How much? Time of last drink? When? 7 PH By: OFC. FLAHAUAN NOTHING I DOUT DEIUK COUPLE OF BURGERS When did you last sleep? How long? Are you sick or injured? □Yes ⊠No Are you diabetic or epileptic? □ Yes ⊠No IN JUST FINE 9 HRS MIUNIGHT □ Yes XNo Do you have any physical defects? 

Yes No Are you under the care of Do you take insulin? a doctor or dentist? □ Yes 🗖 No Coordination Are you taking any medication or drugs? 

Yes 

No Attitude RELAXED CARE FILZE, GENERALLY C ひらってんかべいし POOR STUMBLING OF COURSE NOT DURMAL Eyes: 

Reddened Conjunctiva Blindness: Speech Tracking: □ Normal | Bloodshot □ Watery None □LEye □REye Equal O Unequal Score & DELIBERATE ⊠None Pupil Size: Equal Able to follow stimulus: ¥ Yes □ No Corrective Lens: Normal Droopy □ Glasses □ Contacts, if so □ Hard □ Soft □ Unequal (explain) Pulse & Time HGN Left Eye Right Eye One Leg Stand Vertical Nystagmus? □ Yes X No 1 108 / 2307 Lack of Smooth Pursuit Convergence こり Right Eye Left Eye 2 110/2318 Max. Deviation NO 3. 108 /2325 Angle of Onset とうこうさ NONE Walk and Turn Test 105 PEPERTED Cannot Keep Balance Romberg Balance Starts Too Soon_ Approx. R Sways While Balancing 1st Nine 2nd Nine Stops Walking Uses Arms to Balance Misses Heel-Toe ALL Steps Off Line □ Hopping Raises Arms Puts Foot Down Actual Steps Taken Cannot Do Test (Explain) Internal Clock Describe Turn Type of Footwear ABRUPT SWINEL ABOUT FACE 5\ Estimated At 30 Sec. N/A LOAFERS Room Light Darkness Direct Nasal Area Pupil Size EYEUD TREMORS CHUCKLING ○ Right △ Left TUR 2064 00T CUERR-Left Eye 50-7.0 Oral Cavity BITS OF Draw lines from spot touched 7.5 GREEN LEAFY MATERIAL Right Eye 5.0 -7.0 (<u>.</u>, O 7.5 Rebound Dilation Reaction To Light Hippus □ Yes No. ≰(Yes □ No NOIZMAL LEFT ARM Action weims hery slikely 96 _ Temp98.8 Blood Pressure 140 Muscle Tone: XNormal OFlaccid ORigid Attach Photos Of Fresh Puncture Marks Comments: at Medicine or Drug Have You Been Using? Where Were The Drugs Used? (Location) How Much? Time of Use? TOOK I HO' SMOKE?" WHO ME?" HAH, HAH, HAH chit Remouber" المعتانات ا CHEC Date/Time of Arrest Eval Start Time Time DRE Notified Time Completed 2340 3300 Reviewed By: Member Signature (Include Rank) 7733 ている ミエ マスとい Opinion of Evaluator: □ Rule Out □ Alcohol □ Stimulant □ PCP □ Inhalant □ Medical □ Cannabis □ Depressant ☐ Hallucinogen □ Narcotic Analgesic

#### DRUG INFLUENCE EVALUATION Page 1 of 2 LOG NO. DRE: Sgt. Mike Pryor ARRESTEE: James B. Wright 1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC. 1. LOCATION: Examination of James B. Wright, took place in the DRE room, Colonie Police Department 2. WITNESS: Arresting Officer Tom Flahavan, CHP 3. BREATH TEST: Officer Flahavan administered breath test to Wright, the result was 0.00%. 4. NOTIFICATION / INTERVIEW of ARRESTING OFFICER: Writer was contacted by radio and advised to return to the Department to conduct a DRE evaluation. Trooper Kennedy stated he had observed the subject operating a vehicle at a very slow rate of speed (15/55) southbound on St. Rt 22. When the emergency lights were activated, subject's vehicle slowly drifted left, crossing the northbound lane, through a low hedge and finally coming to rest in a corn field. Subject climbed out of the vehicle laughing. 5. INITIAL OBSERVATIONS: Writer observed the subject seated int he breath testing room. Subject was humming softly. While interviewing Trooper Kennedy, the subject shouted, "Hey Brian, tell him about my wild ride tonight!" MEDICAL PROBLEMS: None noted or stated 7. PSYCHOPHYSICAL TESTS: Romberg Balance: Subject swayed approximately 2" in a circular motion, and exhibited eyelid tremors, and estimated 51 seconds as 30 seconds. Walk and Turn: Subject lost his balance during the instructions, started walking to soon, raised arms repeatedly, and never touched heel to toe. Subject twice requested that the instructions be repeated. One Leg Stand: Subject raised his arms, put his foot down, and swayed, Finger to Nose: Subject missed tip of his nose each time. 8. CLINICAL INDICATORS: Subject's pulse and blood pressure were above the normal range. His pupils were dilated they exhibited rebound dilation and there was lack of convergence. 9. SIGNS of INGESTION: Subject's breath had an odor of marijuana and there were bits of green vegetation on tongue and between the teeth. 10. STATEMENTS: Subject was asked, "when did you smoke the marijuana?" He responded, "what? smoke marijuana? who me?" and then laughed. When asked where he had used the marijuana, the subject replied, "oh, I don't know. Oh gee, seriously, I can't remember." 11. OPINION of EVALUATOR: In my opinion James B. Wright is under the influence of Cannabis and unable to operate a vehicle safely. 12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a urine sample.

13. MISCELLANEOUS: Subject exhibited eyelid tremors and chuckled throughout the evaluation.

Sixty Minutes

## SESSION XXII OVERVIEW OF SIGNS AND SYMPTOMS

#### SESSION XXII OVERVIEW OF SIGNS AND SYMPTOMS

Upon successfully completing this session, the participant will be able to:

- o Name the possible effects that may be observed in each major indicator of drug impairment.
- o Identify the effects that will most likely be observed with suspects under the influence of each drug category.

# Content Segments A. The Major Indicators and Their o Instructor Led Presentations Possible Effects B. Effects Associated With the Drug o Interactive Discussions Categories

Aides Lesson Plan **Instructor Notes** OVERVIEW OF SIGNS AND Total Lesson Time: **SYMPTOMS** Approximately 60 Minutes XXII-O (Objectives NOTE: PRIOR TO THE START OF THIS SESSION, DRAW THE FOLLOWING MATRIX ON THE CHALKBOARD OR FLIPCHART: Stimul Halluc PCP Inhal Possible Depress Narcot **Effects** HGN VGN Lack ConvPupil Size React Light Pulse Rate Blood PressTemp The Major Indicators and Their A.



#### 15 Minutes

- Possible Effects.
  - 1. The major indicators of drug impairment are:
    - a. Horizontal Gaze Nystagmus
    - b. Vertical Gaze Nystagmus
    - c. Lack of Convergence
    - d. Pupil Size

Point to the major indicators on the matrix.

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Aides	Lesson Plan	Instructor Notes
	e. The Reaction of the Pupils to Light.	Point out that the first five major indicators all concern the eyes.
	f. Pulse Rate	
	g. Blood Pressure	
	h. Body Temperature	Point out that the last three major indicators concern the vital signs.
		ANNOUNCE TO THE STUDENTS: WE WILL NOW REVIEW ALL OF THE POSSIBLE EFFECTS THAT WE MIGHT OBSERVE WITH EACH MAJOR INDICATOR.
	2. Possible effects that might be observed with <b>Nystagmus</b> .	
	<ul> <li>a. With Horizontal Gaze         Nystagmus, there are only         two possible effects that         might be observed.     </li> </ul>	
	<ul><li>o Either HGN will be present;</li><li>o or it will be none.</li></ul>	Under the "Possible Effects" column of the matrix, opposite "HGN", write: PRESENT OR
		NONE Point out that there is no drug that stops Horizontal Gaze Nystagmus. Some drugs cause HGN to be present, others do not; but there is no drug that "cures" HGN.
in opt		Now ask students: What are the possible effects we might observe with Vertical Gaze Nystagmus?

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#### Lesson Plan

#### **Instructor Notes**

- b. With Vertical Gaze
  Nystagmus, there are also
  only two possible effects.
  - o Either it will be **present**;
  - o or it will be none.
- 3. For Lack of Convergence, there are also only two possible effects.
  - a. Either Lack of Convergence will be **present**;
  - b. Or it will be none.
  - Just as with Nystagmus, there is no drug that "cures" Lack of Convergence.

- 4. For **Pupil Size**, there are three possible effects that might be seen.
  - a. The pupils might be **normal** of size;
  - b. or, the pupils might be dilated;
  - c. or, they might be constricted.

Opposite "VGN", write:
PRESENT
OR
NONE

Now ask students: What effects might we observe with Lack of Convergence?

Opposite "Lack Conv", write:
PRESENT
OR
NONE

Point out that, when we say that "Lack of Convergence is present", we mean that the eyes are **unable** to converge or cross properly.

Now ask students: What effects might we observe with Pupil Size?

Opposite "Pupil Size", write:

NORMAL
OR
DILATED
OR
CONSTRICTED

Now ask students: What effects might we observe with the pupils' reaction to light?

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#### Lesson Plan

#### **Instructor Notes**

- 5. There are a number of effects that might be observed in the pupils' Reaction to Light.
  - a. The pupils might react in a **normal** manner, i.e., by constricting somewhat in one second or less.
  - b. Or, the pupils might react slow, i.e., by constricting somewhat, but requiring more than one second to do so.
  - c. In some instances, you may observe very little, or no visible reaction to light.
  - d. If there is a visible reaction of the pupils, it is possible that two other effects might be seen.
    - o **Hippus**, i.e., pupils rhythmically pulsating in size.
    - o Rebound Dilation, i.e., a period of constriction followed by dilation with a change equal to or greater than 2 mm.
- 6. For each of the Vital Signs, there are three possible effects.
  - a. The pulse rate, or blood pressure, or body temperature could be normal.
  - b. Or, it could be UP.

Opposite "React Light", write:

NORMAL

OR
SLOW

OR

LITTLE TO NONE VISIBLE

Point out that we should <u>not</u> report that the "pupils did not react at all", but rather we should report "no visible reaction".

Opposite "Pulse Rate", write:

NORMAL

OR

UP

OR

DOWN

Aides	Lesson Plan	Instructor Notes
	c. Or, it could be <b>DOWN</b> .	Write exactly the same things opposite "Blood Press".
		Write exactly the same things opposite "Body Temp".
		Solicit students' comments and questions about the possible effects of the eight major indicators.
45 Minutes	B. Effects Associated with the Drug Categories.	Ask for a student to volunteer to state the major effects that usually will be seen in a suspect under the influence of a CNS Depressant. Correct the students' statements, as necessary, and write the correct effects on the matrix, under the "Depress." column.
	1. CNS Depressants.	
	a. HGN: present	· .
	b. VGN: present	i.e., at high doses for that individual.
	c. Lack Conv. present	individual.
	d. Pupil Size: <b>normal</b> , <u>except</u> with the specific depressant Methaqualone and Soma, which <b>dilates</b> pupils.	
	e. React Light: slow	
	f. Pulse Rate: down except Methaqualone and ETOH, which may elevate.	
	g. Blood Pressure: <b>down</b>	

Aides	Lesson Plan	Instructor Notes
	h. Body Temp: normal	Emphasize that these are the usual major effects that will be observed with CNS Depressants, but we cannot always be certain that all of these effects will be seen.  Thank the "volunteer" student for their help.  Pick another volunteer to state the usual major effects of CNS Stimulants. Correct the student's statements as necessary, and write the correct effects under the "Stimul" column.
	2. CNS Stimulants	
	a. HGN: none	
	b. VGN: none	
	c. Lack Conv: none	
	d. Pupil Size: dilated	
	e. React Light: slow  f. Pulse Rate: up  g. Blood Press: up  h. Body Temp: up	Emphasize that these are the effects usually seen with CNS Stimulants, but we can't guarantee that all of these effects will be observed in each and every case.  Thank the "volunteer" student for his or her help.  Select another volunteer to help with identifying the usual major effects of Hallucinogens.

Aides	Lesson Plan	Instructor Notes
	3. Hallucinogens	
	a. HGN: none	
	b. VGN: none	
	c. Lack Conv. none	
	d. Pupil Size: dilated	
	e. React Light: normal, certain Psychedelic Amphetamines cause slow reaction.	Point out that "Reaction to Light" is the only major indicator that distinguishes Hallucinogens from CNS Stimulants, and "Reaction to Light" is a relatively subtle clue. For this reason, it can be very difficult to differentiate between these two categories.
	f. Pulse Rate: <b>up</b>	
	g. Blood Press: <b>up</b>	Thank the "volunteer" for thier
	h. Body Temp: <b>up</b>	help with Hallucinogens. Pick another volunteer to help with PCP.
	4. Phencyclidine	
•	a. HGN: present	
	b. VGN: present	i.e., at high doses; however, it is more common to see Vertical Gaze Nystagmus in someone under the influence of PCP.
	c. Lack Conv: <b>present</b>	
	d. Pupil Size: normal	
	e. React Light: normal	
	f. Pulse Rate: up	

Aides	Lesson Plan	Instructor Notes
	g. Blood Press: <b>up</b>	
	h. Body Temp: <b>up</b>	Thank the "volunteer" for their help with PCP.
		Select another volunteer to help with Narcotic Analgesics.
	5. Narcotic Analgesics	
	a. HGN: none	
	b. VGN: none	
	c. Lack Conv: <b>none</b>	
	d. Pupil Size: constricted	
	e. React Light: little or none visible	
	f. Pulse Rate: down	
	g. Blood Press: down	Thank the "volunteer" for their help with Narcotic Analgesics.
	h. Body Temp: down	
	6. Inhalants	Select another volunteer to help with Inhalants. Remind the volunteer that, with Inhalants, many of the effects noted on the major indicators will depend upon the specific substance inhaled.
	a. HGN: present	The vast majority of Inhalants will cause HGN; but it is possible that HGN would not be observed with a few specific Inhalants.
	b. VGN: present	High dose for that individual

Lesson Plan	Instructor Notes
c. Lack Conv: present	
d. Pupil Size: normal but may be dilated	
e. React Light: slow	
f. Pulse Rate: <b>up</b>	
g. Blood Press: up/down	The Volatile Solvents and the Aerosols usually cause blood pressure to be above normal; but the Anesthetic Gases can cause blood pressure to be below normal, even though they elevate the pulse rate.
h. Body Temp: up/down/normal	Some Inhalants leave body temperature within the normal range; others may elevate the temperature.  Thank the "volunteer" for their help with Inhalants. Select another volunteer to help with Cannabis.
Cannabis	
a. HGN: none	
b. VGN: none	
c. Lack Conv: <b>present</b>	
d. Pupil Size: dilated or possibly normal	
e. React Light: normal	
f. Pulse Rate: up	
	c. Lack Conv: present d. Pupil Size: normal but may be dilated e. React Light: slow f. Pulse Rate: up g. Blood Press: up/down  h. Body Temp: up/down/normal  Cannabis a. HGN: none b. VGN: none c. Lack Conv: present d. Pupil Size: dilated or possibly normal e. React Light: normal

Aides	Lesson Plan	Instructor Notes
	g. Blood Press: up h. Body Temp: normal	Thank the "volunteer" for their help with Cannabis.  Solicit students' comments or questions about the drug categories.  REFER STUDENTS TO the addendum at the end of this session is a small portion of the available scientific literature dealing with drug influence symptomatology. The sources are considered to be reliable sources of drug symptomatology.  Stress that not all symptoms associated with a drug category will be observed in all subjects in all cases. The excerpts from the references are consistent with DRE instruction and experience.

#### **Session XXII**

Overview of Signs and Symptoms



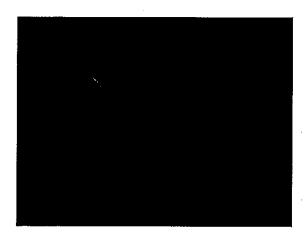
### Overview of Signs and Symptoms

Upon successfully completing this session, the participant will be able to:

- Name the possible effects that may be observed in each major indicator of drug impairment
- Identify the effects that will most likely be observed with suspects under the influence of each drug

Drug Eveluation & Classification Training

AAH U



#### COMPARISON OF DRE SYMPTOMATOLOGY WITH CROSS SECTION OF DRUG

#### SYMPTOMATOLOGY SOURCES

#### CNS DEPRESSANTS:

DRE Symptomatology:

Nystagmus

decreased pulse

decreased blood pressure

uncoordinated

disoriented

sluggish

thick slurred speech

drunk-like appearance

The Pharmacological Basis of Therapeutics, Seventh Edition, Gilman, A.; Goodman, I.; MacMillan Publishing Co. 1985, Barbiturates, pages 546-547:

Nystagmus

Strabismus

difficulty in visual accommodation

vertigo

ataxia gait

positive Romberg sign

Hypotonia

Dysmetria

Diplopia

sluggishness

difficulty in thinking

slowness, slurring of speech

poor comprehension faulty judgement

poor memory

emotional lability

A Primer of Drug Action, Julien, Robert M. W.H. Freeman and Company, New York, 6th Ed. 1992, pp. 61-63.

<u>Drug and Alcohol Abuse, A Clinical Guide to Diagnosis and Treatment</u>, (3rd Ed., Schuckit, M.D., Mark A. Plenum Medical Book Co, New York 1989, p.19.

Encyclopedia of Drug Abuse, O'Brien, Robert; Cohen, Sydney. M.D. Facts on File, INC New York (1984), page 36: barbiturates effects like alcohol (staggering, poor motor control).

<u>Drug Abuse and Dependence</u>, Grinspoon, Lester, MD; Bakalar, James B., Harvard Medical School Mental Health Review No. 1 (1990), page 11: sedative hypnotics same as alcohol and other depressants

<u>Drugs of Abuse</u>, Giannini, A. James, M.D.; Slaby, Andrew E. M.D., Ph.D. Medical Economics Books, Oradell, New Jersey (1989), page 72: Benzodiazepines same as barbiturate effects; pages 247; 292): Barbiturates:

Nystagmus

depressed pulse

depressed blood pressure

diminished concentration

incoordination

decreased reaction time

Manual of Drug and Alcohol Abuse, Guidelines for Teaching in Medical and Health Institutions, ed Arif, Awni. M.D., Westermeyer, Joseph, M.D.. Ph.D..D Plenum Medical Book Company, New York (1988), p. 135.

<u>Diagnostic and Statistical Manual of Mental Disorders</u> (Third Ed, Revised), American Psychiatric Association (1987), p. 159

Maladaptive behavioral changes, e.g., disinhibition of sexual or aggressive impulses, mood lability, impaired judgment, impaired social or occupational functioning.

slurred speech

incoordination

unsteady gait

impairment in attention or memory

#### CNS STIMULANTS:

DRE Symptomatology:

dilated pupils

increased temperature

body tremors

excited

111 11

talkative

anxiety redness to nasal area

loss of appetite

increased alertness

increased pulse rate

increased blood pressure

restlessness

euphoric

exaggerated reflexes

grinding teeth

runny nose

insomnia

The Pharmacological Basis of Therapeutics, Seventh Edition,

Gilman, A,; Goodman, I.; MacMillan Publishing Co. 1985, Cocaine 551-554

Medical Toxicology-Diagnosis and Treatment of Human Poisoning, Ellenhorn, Matthew J., Barceloux, Donald G. Elsevier Science Pub. Co. 1988, Amphetamines, Page 634:

Mild influence:

Mydriasis restlessness irritability

hyperreflexia talkativeness insomnia flushing

tremor Diaphoresis

combativeness

nausea

vomiting

pallor

dry mucous membranes

Moderate:

hyperactivity hypertension confusion Tachypnea

Tachycardia

premature ventricular contraction

chest discomfort

vomiting

abdominal pain

Profuser Diaphoresis

mild temperature

elevation

impulsivity hallucinations

repetitive behavior

panic reactions

Serious:

delirium

marked Hypertension/Tachycardia

Hyperreflexia

convulsions

Hypotension

coma

Cocaine, page 650-659

Early Stimulation:

euphoria excitement irritable behavior sudden headache

Garrulity apprehension Mydriasis

vomiting

nausea dizziness

twitching of small muscles tremor

tics ierks

Cocaine Psychosis

hallucinations

elevation of pulse

increased respiration

Advanced:

convulsions

Hyperreflexia

decreased consciousness

increased pulse and blood pressure

Later Stages:

Hypotension

Hypothermia

Dyspnea et al

A Primer of Drug Action, Julien, Robert M. W.H. Freeman and Company, New York, 1992, pages 120-123: Amphetamines and cocaine (CNSS):

dilation of pupils

increased blood pressure

slight tremor

restlessness

agitation

possibly hallucinations

<u>Drug and Alcohol Abuse, A Clinical Guide to Diagnosis and Treatment,</u> (3rd Ed., Schuckit, M.D., Mark A. Plenum Medical Book Co, New York 1989, page 99: CNSS cause:

dilation of pupils

rapid heart rate

elevation of blood pressure

tremor in hands

increased body temperature

restlessness

Encyclopedia of Drug Abuse, O'Brien, Robert; Cohen, Sydney. M.D. Facts on File, INC New York (1984), pages 25, 121: Amphetamine:

dilation of pupils

increase heart rate

blood pressure

flushing

teeth grinding tremors

dry mouth lack of coordination

pages 64, 100, 121:

dilation of pupils

increased heartbeat

increased temperature

similar to Amphetamine

<u>Drug Abuse and Dependence</u>, Grinspoon, Lester, MD; Bakalar, James B., Harvard Medical School Mental Health Review No. 1 (1990), pages 8 and 10 Cocaine and Amphetamine:

dilated pupils

increased pulse

increased blood pressure

vasoconstriction

agitation tremors

increased temperature

<u>Drugs of Abuse</u>, Giannini, A. James, M.D.; Slaby, Andrew E. M.D., Ph.D. Medical Economics Books, Oradell, New Jersey(1989), page 29 Amphetamines:

pupil dilation (Mydriasis)

increased pulse rate

elevated blood pressure talkative

hyperactive irritable

restless

Anorexia

tremors

urinary retention

teeth grinding (Bruxism) illogical, loose thoughts

fidgety, jerky, random motions

Page 295: Cocaine:

dilated pupils

Tachycardia

increased blood pressure

vasoconstriction

Hyperpyrexia

Manual of Drug and Alcohol Abuse, Guidelines for Teaching in Medical and Health Institutions, ed Arif, Awni. M.D., Westermeyer, Joseph, M.D.. Ph.D..D Plenum Medical Book Company, New York (1988) page 142: Amphetamine:

increased pulse

increased blood pressure

possibly increased temperature

increased wakefulness

general increase in psychomotor

activity

page 145: Cocaine

Mydriasis (dilated pupils);

may cause psychosis

euphoria

agitation

<u>Diagnostic and Statistical Manual of Mental Disorders</u> (Third Ed, Revised), American Psychiatric Association (1987), p. 142.

#### COCAINE:

Maladaptive behavioral changes, e.g., euphoria, fighting, grandiosity, hyper-vigilance, psychomotor agitation, impaired judgment, impaired social or occupational functioning.

pupillary dilation

Tachycardia

 $elevated\ blood\ pressure$ 

perspiration or chills

nausea or vomiting

visual or tactile hallucinations

#### **AMPHETAMINE**

Maladaptive behavioral changes, e.g., fighting, grandiosity, hyper-vigilance, psychomotor agitation, impaired judgment, impaired social or occupational functioning.

pupillary dilation

elevated blood pressure

nausea or vomiting

Tachycardia

perspiration or chills

#### HALLUCINOGENS:

DRE Symptomatology:

dilated pupils

increased blood pressure

poor perception of time/distance

dazed appearance Synesthesia

paranoia nausea

difficulty in speech

increased pulse rate increased temperature

body tremors hallucinations

uncoordinated disoriented

perspiring

The Pharmacological Basis of Therapeutics, Seventh Edition, Gilman, A.; Goodman, I.; MacMillan Publishing Co. 1985, LSD and Related Drugs, page 564

pupillary dilation

increased blood pressure

Tachycardia

Hyperreflexia

tremor

nausea

Piloerection

muscular weakness

increased body temperature

hallucinations

Hyper vigilance

loss of boundaries

Synesthesia

Medical Toxicology-Diagnosis and Treatment of Human Poisoning, Ellenhorn, Matthew J., Barceloux, Donald G. Elsevier Science Pub. Co. 1988, LSD, pages 667-669:

pupillary dilation

increased heart rate

increased body temperature

Piloerection tremor

weakness Hyperreflexia

Ataxia

hallucinations

depersonalization

poor judgment

mood swings

A Primer of Drug Action, Julien, Robert M.; W. H. Freeman and Company, New York, 1992

<u>Drug and Alcohol Abuse, A Clinical Guide to Diagnosis and Treatment,</u> (3rd Ed.), Schuckit, M.D., Mark A. Plenum Medical Book Co, New York 1989 page 160:

dilated pupils

increased blood pressure

increased awareness

faltered body images

sensory input

fine tremor

flushed face

increased body temperature

Encyclopedia of Drug Abuse, O'Brien, Robert; Cohen, Sydney. M.D. Facts on File, Inc New York (1984), pages 100; 115 120, 153): Hallucinogens:

dilated pupils

increased heart rate

increased blood pressure

increased temperature

profuse perspiration

loss of appetite

hallucinations

<u>Drug Abuse and Dependence</u>, Grinspoon, Lester, MD; Bakalar, James B., Harvard Medical School Mental Health Review No. 1 (1990)

<u>Drugs of Abuse</u>, Giannini, A. James, M.D.; Slaby, Andrew E. M.D., Ph.D. Medical Economics Books, Oradell, New Jersey (1989), page 218: LSD:

Ataxia

high blood pressure

Hyperreflexia

incoordination

Tachycardia

Manual of Drug and Alcohol Abuse, Guidelines for Teaching in Medical and Health Institutions, ed Arif, Awni. M.D., Westermeyer, Plenum Medical Book Company, New York (1988)

<u>Diagnostic and Statistical Manual of Mental Disorders</u> (Third Ed, Revised), American Psychiatric Association (1987), p. 145.

Maladaptive behavioral changes, e.g., marked anxiety or depression, ideas of reference, fear of losing one's mind, paranoid ideation, impaired judgment, impaired social or occupational functioning.

Perceptual changes occurring in a state of full wakefulness and alertness, e.g., subjective intensification of perceptions, depersonalization, derealization, illusions, hallucinations, Synesthesia

pupillary dilation

Tachycardia

sweating

palpitations

blurring of vision

tremors

incoordination

#### PHENCYCLIDINE

DRE Symptomatology:

Nystagmus

increased blood pressure

perspiring

blank stare "moon walking"

incomplete responses

repetitive speech

cyclic behavior

hallucinations

increased pulse

increased temperature

warm to the touch

early onset of nystagmus

difficulty in speech repetitive response

increased pain threshold

confused, agitated

possibly violent and combative

The Pharmacological Basis of Therapeutics, Seventh Edition, Gilman, A,; Goodman, I.; MacMillan Publishing Co. 1985, PCP, page 565-567

Nystagmus

elevated blood pressure

staggering gait

numbness of extremities

muscular rigidity

drowsiness

repetitive movements

elevated heart rate feeling of intoxication

slurred speech

sweaty

blank stare

hostile behavior

Medical Toxicology-Diagnosis and Treatment of Human Poisoning, Ellenhorn, Matthew J., Barceloux, Donald G. Elsevier Science Pub. Co. 1988, PCP 768-777:

Nystagmus

depressed light reflexes

diminished pain

tremors

slurred speech

increased pulse rate

Amnesia

body image distortion

depersonalization

hallucinations

Miosis

blurred vision

Ataxia

muscle weakness

drowsiness

increased blood pressure

anxiety/agitation

euphoria

disordered thought processes

A Primer of Drug Action, Julien, Robert M. W.H. Freeman and Company, New York, 1992, page 262: PCP:

increased blood pressure

blank stare

disinhibition

mood swings

muscle rigidity

agitation

delirium excitement

disorientation

hallucinations

analgesia

speech difficulty

pain tolerance

elevated blood pressure

Drug and Alcohol Abuse, A Clinical Guide to Diagnosis and Treatment, (3rd Ed.), Schuckit, M.D., Mark A. Plenum Medical Book Co, New York 1989 p. 178

sweating

muscle rigidity

fever convulsions

increased blood pressure

Encyclopedia of Drug Abuse, O'Brien, Robert; Cohen, Sydney. M.D. Facts on File, INC New York (1984), page 100, 208: PCP:

Nystagmus

increased blood pressure

increased pulse rate

flushing

mood swings

hallucinations speech difficulties

changes in body awareness

violent behavior

decreased responsiveness

Drug Abuse and Dependence, Grinspoon, Lester, M.D.; Bakalar, James B., Harvard Medical School Mental Health Review No. 1 (1990), page 25: PCP:

body image distortions

increased blood pressure

Nystagmus

muscle rigidity

loss of muscle control

incoherent speech

memory loss drooling

blank stare

Drugs of Abuse, Giannini, A. James, M.D.; Slaby, Andrew E. M.D., Ph.D. Medical Economics Books, Oradell, New Jersey (1989) page 296: PCP:

Nystagmus

disorientation

hallucination

extreme agitation

loss of motor control automated speech

disassociation from

Nystagmus at rest

environment

Manual of Drug and Alcohol Abuse, Guidelines for Teaching in Medical and Health Institutions, ed Arif, Awni. M.D., Westermeyer, Joseph, M.D. Ph.D.D Plenum Medical Book Company, New York (1988), page 156: PCP:

Ataxia

tremors

muscular hypertonicity

Hyperreflexia

Ptosis

Tachycardia Horizontal Gaze, Vertical Gaze

and Rotary Nystagmus

elevated blood pressure

mood swings

Diagnostic and Statistical Manual of Mental Disorders (Third Ed, Revised), American Psychiatric Association (1987), p. 155.

Maladaptive behavioral changes, e.g., belligerence, assaultiveness, impulsiveness, unpredictability, psychomotor agitation, impaired judgment, impaired social or occupational functioning.

Vertical or Horizontal Gaze Nystagmus increased blood pressure or heart rate numbness or diminished responsiveness to pain. Ataxia Dysarthria (slurred speech) muscle rigidity seizures Hyperacusis

#### NARCOTICS:

DRE Symptomatology:

constricted pupils

decreased blood pressure

droopy eyelids

drowsiness

low, raspy speech

facial itching

fresh puncture marks

decreased pulse rate decreased temperature (Ptosis) "on the nod" depressed reflexes

dry mouth euphoria

The Pharmacological Basis of Therapeutics, Seventh Edition, Gilman, A.; Goodman, I.; MacMillan Publishing Co. 1985, Opiods page 541-545

Medical Toxicology-Diagnosis and Treatment of Human Poisoning, Ellenhorn, Matthew J., Barceloux, Donald G. Elsevier Science Pub. Co. 1988; Heroin, pages 702-703. See also Methadone, Demerol, etc.:

A Primer of Drug Action, Julien, Robert M. W.H. Freeman and Company, New York, 1992, page 196-198: Morphine:

constructed pupils

decreased blood pressure

 ${\bf drowsiness}$ 

Dysphoria

mental clouding

sedation

depressed respiration

Analgesia

euphoria

<u>Drug and Alcohol Abuse, A Clinical Guide to Diagnosis and Treatment,</u> (3rd Ed., Schuckit, M.D., Mark A. Plenum Medical Book Co, New York 1989

Decrease pain (p.6)

Encyclopedia of Drug Abuse, O'Brien, Robert, Cohen, Sydney. M.D. Facts on File, INC New York (1984) page 100, 120, 123, 124: Narcotics:

constricted pupils

reduced heart rate

Analgesia

depressed appetite

euphoria

going "on the nod"

<u>Drug Abuse and Dependence</u>, Grinspoon, Lester, MD; Bakalar, James B., Harvard Medical School Mental Health Review No. 1 (1990), page 14: Narcotics:

constricted pupils

"nodding off"

dreamy state

pain suppression

euphoria

<u>Drugs of Abuse</u>, Giannini, A. James, M.D.; Slaby, Andrew E. M.D., Ph.D. Medical Economics Books, Oradell, New Jersey (1989) page 293 - 294:

Miosis (constricted pupils)

Bradycardia

Hypothermia

(decreased heart beat)

decreased temperature)

euphoria/dysphoria

drowsiness lethargy

confusion

flaccid muscle tone

depressed respiration

Analgesia

Manual of Drug and Alcohol Abuse, Guidelines for Teaching in Medical and Health Institutions, ed Arif, Awni. M.D., Westermeyer, Joseph, M.D.. Ph.D..D Plenum Medical Book Company, New York (1988), page 132

Miosis (constricted pupils)

low blood pressure

itching

flushing sweating

<u>Diagnostic and Statistical Manual of Mental Disorders</u> (Third Ed, Revised), American Psychiatric Association (1987), p. 152.

Maladaptive behavioral changes, e.g., initial euphoria followed by apathy, dysphoria, psychomotor retardation, impaired judgment, impaired social or occupational functioning.

pupillary constriction

drowsiness

slurred speech

impairment in attention or memory

#### INHALANTS:(Toluene)

DRE Symptomatology:

Nystagmus

increased pulse rate

increased blood pressure

residue around nose

odor on mouth

nausea disorientation

slurred speech

confusion

The Pharmacological Basis of Therapeutics, Seventh Edition, Gilman, A.; Goodman, I.; MacMillan Publishing Co. 1985, Inhalants, page 567

Drug and Alcohol Abuse, A Clinical Guide to Diagnosis and Treatment, (3rd Ed., Schuckit, M.D., Mark A. Plenum Medical Book Co, New York 1989, p. 185

decreased inhibitions

floating sensation

drowsiness

light sensitivity

sneezing runny nose

Encyclopedia of Drug Abuse, O'Brien, Robert; Cohen, Sydney. M.D. Facts on File, INC New York (1984)

lowered inhibitions

restlessness

incoordination confusion

disorientation

nausea

impaired judgment

<u>Drug Abuse and Dependence</u>, Grinspoon, Lester, MD; Bakalar, James B., Harvard Medical School Mental Health Review No. 1 (1990)

Drugs of Abuse, Giannini, A. James, M.D.; Slaby, Andrew E. M.D., Ph.D. Medical Economics Books, Oradell, New Jersey (1989), pages 265, 272, 297: Toluene:

Nystagmus

mental dulling

tremors cerebellar

Ataxia

rambling speech light headedness irritability

CNS depression that mimics

tremors Ataxia

Narcotic Analgesics

blank stare euphoric mood

Manual of Drug and Alcohol Abuse, Guidelines for Teaching in Medical and Health Institutions, ed Arif, Awni. M.D., Westermeyer, Joseph, M.D.. Ph.D..D Plenum Medical Book Company, New York (1988)

brief euphoria giddy intoxication, similar to alcohol CNS depression (volatile solvents/toluene) dizziness Vertigo

Diagnostic and Statistical Manual of Mental Disorders (Third Ed, Revised), American Psychiatric Association (1987), p. 149.

Maladaptive behavioral changes, e.g., belligerence, assaultiveness, apathy, impaired judgment, impaired social or occupational functioning.

Nystagmus

dizziness

incoordination

slurred speech

unsteady gait

lethargy

depressed reflexes

psychomotor retardation

tremor generalized muscle

blurred vision or diplopia

stupor or coma

weakness

euphoria

#### **CANNABIS**

DRE Symptomatology:

dilated pupils

odor of Marijuana

body tremors

relaxed inhibitions

paranoia

marked reddening of conjunctivae debris in mouth

eyelid tremors

increased appetite

disorientation

impaired perception of time and distance

The Pharmacological Basis of Therapeutics, Seventh Edition, Gilman, A.; Goodman, I.; MacMillan Publishing Co. 1985, Cannabis, pages 559-561

euphoria

temporal disintegration

information processing impairment

dry mouth

short term memory impairment balance and stance impairment

increased hunger additive to alcohol

Lower doses

affects perception, impairing well beyond when subject subjectively feels effects; alters all information processing; relatively simple motor skills unaffected

High doses:

anxiety

increased heart rate

marked reddening of Conjunctiva

hallucinations

increased systolic blood pressure

simple motor skills affected

Medical Toxicology-Diagnosis and Treatment of Human Poisoning, Ellenhorn, Matthew J., Barceloux, Donald G. Elsevier Science Pub. Co. 1988; Cannabis, page 678-681

reddening of Conjunctiva

motor coordination impairment

relaxation

temporal distortion

(time slows)

impairment of motor tasks and

reaction times requires higher dosages

loss of short term memory

systematic thinking impaired

dry mouth

alteration in mood

euphoria

sleepiness

decrease in balance, steadiness and

muscle strength

elective attention stimulated appetite

<u>A Primer of Drug Action</u>, Julien, Robert M. W.H. Freeman and Company, New York, 1985: page 178, Marijuana

reddening of Conjunctiva increased blood pressure dry mouth altered sensory perception Drug and Alcohol Abuse, A Clinical Guide to Diagnosis and Treatment, (3rd Ed., Schuckit, M.D., Mark A. Plenum Medical Book Co, New York 1989, page 145: Cannabis:

red Conjunctiva

euphoria

relaxation

dry mouth

increased heart rate

possibly Nystagmus short term memory

time distortion impairment in ability to do

tremors

multi-step tasks

decrease level of motor coordination

Encyclopedia of Drug Abuse, O'Brien, Robert; Cohen, Sydney. M.D. Facts on File, INC New York (1984), pages 100, 120: Marijuana:

red eye

increased appetite

increased heart beat

time and space distortions

dryness of mouth and throat

increased heart rate

increased pulse rate

lack of coordination

Drug Abuse and Dependence, Grinspoon, Lester, MD; Bakalar, James B., Harvard Medical School Mental Health Review No. 1 (1990).page 19: Marijuana:

increased appetite

faster heartbeat

bloodshot eyes

confusion

agitation

incoordination

hallucinations

Drugs of Abuse, Giannini, A. James, M.D.; Slaby, Andrew E. M.D., Ph.D. Medical Economics Books, Oradell, New Jersey (1989), page 296: Cannabis:

red Conjunctiva

increased appetite

pleasant relaxation

intensification of sensations

slowed time

passivity

apathy

Tachycardia (increased heart rate)

problems with motor coordination

Manual of Drug and Alcohol Abuse, Guidelines for Teaching in Medical and Health Institutions, ed Arif, Awni. M.D., Westermeyer, Joseph, M.D., Ph.D., Plenum Medical Book Company, New York (1988), page 147: Cannabis:

red Conjunctiva

increased hunger

changes in time sense

short-term memory loss

memory

dry mouth

coordination

Tachycardia (rapid heart beat)

balance and stance

elevated systolic pressure affected

<u>Diagnostic and Statistical Manual of Mental Disorders</u> (Third Ed, Revised), American Psychiatric Association (1987), p. 140.

Maladaptive behavioral changes, e.g., euphoria anxiety, suspiciousness, or paranoid ideation, sensation of slowed time, impaired judgment, social withdrawal.

red Conjunctiva

increased appetite

Tachycardia (rapid heart)

dry mouth

Fifty Minutes

#### SESSION XXIII

RESUME PREPARATION AND MAINTENANCE

#### SESSION XXIII RESUME PREPARATION AND MAINTENANCE

Upon successfully completing this session, the participant will be able to:

- o Describe and discuss the purpose of the resume.
- o Identify the elements of a Drug Recognition Expert's resume.
- o Prepare a basic resume summarizing their relevant training, education, experience and accomplishments to date.
- o Update and extend the resume, as their relevant achievements continue to expand.

# Content Segments A. Purpose of the Resume o Instructor Led Presentations B. Preparation for Court Qualification o Group Work session C. Resume Content o Reading Assignments D. Guidelines for Resume Preparation and Maintenance

Aides	Lesson Plan	Instructor Notes
, 33 ,	RESUME PREPARATION AND MAINTENANCE	Total Session Time: Approximately 50 Minutes
		Session title on wallchart.
XXIII-O		Overview session objectives, content segments and learning activities.
(Objectives)	•	
	A. Purpose of the Resume	
10 Minutes		
XXIII-1	1. The basic purpose of the resume is to record education, training and experience in a single document for use in establishing qualifications when testifying in court.	
·	2. Generally a witness can testify only to personal knowledge.	Point out that this generally consists of facts which they observed or witnessed.
	3. Witness cannot give an opinion on a matter.	<u>Point out</u> that opinions are allowed only if the witness is qualified as an expert.
XXIII-2	4. Basic rule is that a person skilled in some art, trade, science or profession, having a knowledge of matters not within the knowledge of persons of average education, learning and experience, may assist the jury in arriving at a verdict by expressing an opinion on a state of facts shown by the evidence and based upon his or her special knowledge.	(People vs. Willis, 70 Cal APP. 465)

		•	·
Aides		Lesson Plan	Instructor Notes
	5.	A witness is not qualified as an expert witness unless it is shown he or she is familiar with the subject upon which he or she is asked to give an opinion.	(People vs McLean, 56 Cal 2d 660)
	6.	Only the court can determine whether a witness is qualified to testify as an expert.	
	7.	Where a witness is qualified to give expert testimony, any question as to degree of knowledge goes to weight rather than admissibility.	(People vs Perry, 44 Cal 2d 861)
XXIII-3	8.	Witnesses' qualification is achieved through <u>Voir Dire</u> <u>Examination</u> .	Voir Dire - literally, French for "to see, to say"; loosely translated as "to seek the truth").
5 Minutes	В.	Preparation for Court Qualification	
o minutes	1.	Being qualified as an expert may be as simple as stating your occupation, or take several hours of exhausting questioning by both the prosecutor and the defense attorney.	
	2.	Although knowledge only	

greater than what the public has is required to qualify as an expert, your testimony will carry much more "weight" if you

information is essential for an

give his or her qualifications as

officer who is called upon to

have good credentials.

an expert in any field.

3. Accurate, up to date

Point out that it is imperative that each officer maintain an ongoing resume to establish their credentials as an expert.



- 4. Drug Recognition Experts will base their expertise on the following areas:
  - a. Formal education and training
  - b. Relevant Experience
  - c. Outside readings and studies
- C. Resume Content
  - 1. Formal education.
    - a. High school(s) attended
    - b. Colleges and Universities attended.
    - c. Specialized College or University level courses.
  - 2. Formal training.
    - a. Police Academy (recruit training)

- list dates highlight classes which provided knowledge in the area of drugs.
- o list dates, major, degree, etc. highlight classes which provided knowledge in the area of drugs.
- o list dates, instructor, subject(s) covered, credits, etc.
- o list dates, length, major topics covered, etc. Highlight classes which provided knowledge or skills in the area of drugs.



XXIII-5

Aides	Lesson Plan	Instructor Notes	
	b. Specialized police training or in-service training.	o list dates, length, instructor(s), subject(s) covered, etc. Highlight training which provided knowledge or skills in the area of drugs.	
	<ul><li>c. Other specialized training:</li><li>o military training</li><li>o lectures and seminars</li></ul>	o list dates, length, instructor(s), subject(s) covered, etc. Highlight training which provided knowledge or skills in the area of drugs.	
3.	Experience  a. Job experience - years	o list dates, division, duties, etc., include loans to specialized units.	
	b. Assignments	o list agencies, dates, assignments, etc.	
	c. Prior law enforcement experience		
	d. Other job related experience	o list employer, dates, duties, assignments, etc. which provided experience in the area of drugs.	
	<ul> <li>e. Drug enforcement/ evaluation experience:</li> <li>o total vehicle stops</li> <li>o total DWI investigations</li> <li>o total DWI arrests</li> <li>o total drug evaluations</li> <li>o total filings</li> <li>o total convictions</li> </ul>	Point out that it is important to maintain accurate records of all enforcement activities; documentation of the ratio of stops of investigations and investigations to arrests is essential. Not all stops are investigated and not all investigations result in arrests; demonstrates that officer is fair and impartial and that each case is decided on individual merits.	

Aides	Lesson Plan	Instructor Notes
	f. Prior testimony:	o list date, court, judge, charge, area qualified, etc.
	o municipal court	charge, area quanneu, ew.
	o superior court	
	o number of times qualified as an expert in drug cases	·
·	o number of times qualified as an expert in other cases	. *
	4. Outside readings and studies	
	a. Drug related texts read	o list title(s), author(s), subject(s), etc.
·.	b. Departmental training bulletins	
	c. Journals	
	d. Research papers	
	e. Drug related films viewed	
-	5. Training or research conducted (if applicable)	o list classes, briefings, training officer assignments, etc. where you served as an instructor or coach, etc. or conducted or participated in research, e.g. Alcohol workshop.
	6. Publications (if applicable)	o list all writings that were published, including departmental briefing papers, etc.
		,

Aides	Lesson Plan	Instructor Notes
15 Minutes	D. Guidelines for Resume Preparation and Maintenance	
	<ol> <li>List information in chronological order.</li> <li>Review and update resume frequently and record date of review.</li> </ol>	Refer students to sample resumes in their manuals and review steps for preparing the resume and keeping it up to date.  Review the sample resumes briefly with the students.

# **Session XXIII**

# Resumé Preparation and Maintenance



### Resumé Preparation and Maintenance

Upon successfully completing this session, the participant will be able to:

- · Describe and discuss the purpose of the resumé
- Identify the elements of a Drug Recognition Expert's
- Prepare a basic resumé summarizing their relevant training, education, experience and accomplishments to date
- Update and extend the resumé, as their relevant achievements continue to expand

Drug Evaluation & Classification Training

XXIII-0

#### Witness

- Generally can testify only to personal knowledge--facts which they observed or witnessed
- · Cannot give an opinion

Drug Evaluation & Classification Training

XXIII-

# **Expert Witness**

- Basic rule--person skilled in some art, trade, science, or profession, having knowledge of matters not within knowledge of persons of average education, learning, and experience
- May assist jury in arriving at a verdict by expressing an opinion on a state of facts shown by the evidence and based upon his or her special knowledge
- Only the court can determine whether a witness is qualified to testify as an expert

Drug Evaluation & Classification Training

XXIII.2

#### Voir Dire:

To seek the truth (literally, "to see, to say")

Date Evaluation & Classification Training

XXIII-3

# **Expertise/Qualifications**

These are based on:

- · Formal Education and Training
- Experience
- Outside readings and studies

Drug Evaluation & Classification Training

XXIII-

# Resumé Content

- Formal education
- Formal training
- Experience
- · Prior testimony
- · Outside readings and studies
- Training/research conducted
- Publications

Drug Evaluation & Classification Training

XXIII-5



# SAMPLE RESUME NUMBER ONE

# SHELTON POLICE DEPARTMENT

Traffic Division

The Resume of:

SERGEANT DAVID CARROLL REGAN Certified Drug Recognition Technician

Latest update: 3/17/XX

# Sgt. David C. Regan

### Introduction

Sergeant David Carroll Regan is a supervisor in the Traffic Division, Shelton Police Department. He currently commands the special Impaired Driving Enforcement Activities Squad (IDEAS), a unit he was instrumental in forming. Sgt. Regan is a 15 year veteran of law enforcement. Prior to joining the Shelton Police Department ten years ago, he served for five years as a deputy with the Fairfield County Sheriff's Department.

Sergeant Regan has been assigned to the Traffic Division since his promotion to sergeant on 11/18/YY. His duties have included coordination of speed and DWI enforcement activities, the Joint Shelton-Derby Task Force for Sobriety Checkpoints, the Officer Friendly Program, the Motorcycle Safety Education Project, and general supervision of Traffic Division officers. He also serves as the Department's principal instructor for radar speed measurement, Standardized Field Sobriety Testing and Drug Recognition Expert training.

Sergeant Regan holds a Bachelor's Degree in the Administration of Justice from Fairfield University, and currently is a candidate for a Master's Degree in Police Science and Administration at the University of Stratford. He also holds an Instructor Certificate from the State Law Enforcement Training Board.

Sergeant Regan has served on two committees of the Governor's Task Force to Prevent Drunk Driving: The Standardized Field Sobriety Tests Committee and The Paperwork Reduction Committee. The one page Standard Notetaking Guide for Field Sobriety Testing that is employed by all departments statewide was designed by him.

#### Law Enforcement Experience

11/18/YY to Present Sergeant, Traffic Division

Shelton Police Department Supervisor, IDEAS Unit

Drug Recognition Expert Program Coordinator

7/8/ZZ to 11/17/YY Patrol Officer First Class

Training and Operations Shelton Police Department

Unit Supervisor, Traffic Law Enforcement Training Branch

9/11/XX to 7/7/ZZ Patrol Officer

Third Precinct, Motorcycle Shelton Police Department

# Sgt. David C. Regan

# Law Enforcement Experience (continued)

11/5/MM to 9/10/XX

Patrol Officer

First Precinct

Shelton Police Department

10/10/NN to 11/4/MM

Deputy

Traffic Patrol

Fairfield County Sheriff's Department

### Special Police Training

10/XX National Highway Traffic Safety Administration

**DRE Instructor Training** 

(Certified as a DRE Instructor on 11/12/XX)

8/XX Drug Enforcement Administration

**Drug Interdiction Seminar** 

11/YY National Highway Traffic Safety Administration

Drug Evaluation and Classification Training: DRE School

(Certified as a DRE on 1/28/XX)

10/YY National Highway Traffic Safety Administration

Drug Evaluation and Classification Training: PRE School

3/YY Southeastern University Institute of Police Technology

Special Conference: Managing DWI Squads

4/ZZ International Association of Chiefs of Police

Instructor Training in Horizontal Gaze Nystagmus and

**Divided Attention Field Sobriety Tests** 

10/MM University of Stanford, Northern Police Institute

Standardized Field Sobriety Testing

6/NN Acme Scientific Instruments, Inc.

(Certified to perform inspection and repair of the Intoxotector J2Z

breath testing instrument on 6/22/NN)

# Sgt. David C. Regan

### Court Qualification Record

8/VV Qualified as Drug Recognition Expert in a case involving

Phencyclidine impairment. (Judge Sally Grey, 8th District)

11/WW Qualified as Drug Recognition Expert in a case involving a

combination of CNS Stimulant and Narcotic Analgesic. (Judge Lewis

Buchanan, Superior Court)

3/WW Qualified as Drug Recognition Expert in a case involving Cannabis

impairment. (Judge Sally Grey, 8th District)

9/UU Qualified as Drug Recognition Expert in a case involving Narcotic

Analgesic impairment. (Judge Jerome Byrnes, 8th District)

# Specialized Readings

<u>Title</u> <u>Author</u>

Drug and Alcohol Abuse Marc A. Schuckit, M.D.

A Primer of Drug Action Jerome Jaffee, Robert Petersen and Ray

Hodgson

The Practitioner's Guide to Ellen L. Bassuk, M.D. and

Psychoactive Drugs Stephen C. Schoonover, M.D.

Drug Abuse: A Manual for Law Smith, Kline & French (pub.)

Enforcement Officers

Licit and Illicit Drugs Edward M. Brecher

Chocolate to Morphine Andrew Weil, M.D. and Winifred Rosen

Cocaine Addiction U.S. Department of Health and Human

Services

Marijuana Alert Peggy Mann

# SAMPLE RESUME NUMBER TWO

TRUMBULL POLICE DEPARTMENT

The Resume of:

OFFICER ANN MARIE REED Certified Drug Recognition Technician

Latest Update: 4/25/YY

#### Officer Ann M. Reed

#### Introduction

Officer Ann Marie Reed is an eight year veteran with the Trumbull Police Department. She is currently assigned to the Special Operations Branch of the Administrative Division, where she serves as a Narcotics Enforcement Officer. Previously, she has served in the same Branch as a Vice Enforcement Officer, and as a patrol officer in the Department's first and second precincts.

Officer Reed is a graduate of Monroe College, with the Bachelor's Degree in Police Science and Administration. She is currently a candidate for the JD Degree at the Law School of the University of Bridgeport.

т	T1 C	T3 )
law	Enforcement	Experience
11U 11	111101 CCIII CIII	TIMPOLICITOO

5/12/VV to Present Narcotics Enforcement Officer and Drug Recognition Expert

Special Operations Branch Trumbull Police Department

3/26/WW to 5/11/VV Vice Enforcement Officer Special Operations Branch

Trumbull Police Department

9/23/XX to 3/25/WW Patrol Officer

First Precinct

Trumbull Police Department

8/28/NN to 9/22/XX Patrol Officer

Second Precinct

Trumbull Police Department

5/15/NN to 8/25/NN Trainee

Fairfield County Regional Police Academy

(Graduated 8/25/NN)

Special Police Training

2/YY University of Norwalk, Police Science Institute

Seminar: Packaging and Transport of Illicit Drugs

10/VV University of Norwalk, Police Science Institute

Seminar: Suppression of Drug-related Crime

3/VV National Highway Traffic Safety Administration

Drug Evaluation and Classification Training: DRE School

(Certified as a DRE on 5/22/VV)

#### Officer Ann M. Reed

Special Police Training (Continued)

2/VV

Fairfield County Regional Police Academy

Drug Evaluation and Classification Training: PRE-School

10/WW

Fairfield County Regional Police Academy Standardized Field Sobriety Testing

Publications Authored

Reed, Ann M. and Cockroft, Robert S., "Narcotics Enforcement Tactics for the Medium-sized Department"; The Police Chief. January 17, 19XX.

Reed, Ann M., Procedures for Requesting Drug Recognition Expert Services; Training Bulletin for the Trumbull Police Department. 6/VV.

Reed, Ann M., Recognizing the Heroin Addict; Training Bulletin for the Trumbull Police Department. 1/VV.

Court Qualification Record

Qualified as an expert witness for identification of Heroin 11/WW

impairment. (Judge Michael Adkins, 7th District)

3/WW Qualified as a Drug Recognition Expert in a case involving a

combination of CNS Stimulant and Narcotic Analgesic. (Judge

Author

Roberta Mayer, 7th District)

Qualified as an expert witness for identification of "track" marks. 9/ZZ

(Judge Charles Peltier, 7th District)

Specialized Readings

<u>Title</u>

Barbara McVan, M.D. Signs and Symptoms Handbook

Drugs From A to Z Richard R. Lingeman

Richard Seymour and David E. Smith, M.D. Guide to Psychoactive Drugs

Robert M. Julien, M.D. Addictions: Issues and Answers

Report on Synthetic China

White: Fentanyl Det. James Miller, LAPD

One Hour and Fifty Minutes

SESSION XXIV
DRUG COMBINATIONS

### SESSION XXIV DRUG COMBINATIONS

Upon successfully completing this session, the participants will be able to:

- o Explain the prevalence of polydrug use among drug impaired suspects and identify common combinations of drug abused by those suspects.
- o Explain the possible effects that combinations of drugs can produce on the major indicators of drug impairment, and define the terms "Null", "Overlapping", "Additive" and "Antagonistic" as they relate to polydrug effects.
- o Identify the specific effects that are most likely to be observed in persons under the influence of particular drug combinations.

# A. The Prevalence of Polydrug Use o Instructor Led Presentations B. Possible Effects of Drug Combinations o Interactive Discussions

0

C. Identifying Expected Indicators of Specific Combinations

Content Segments

- Workbook Exercise
- o Video Presentations

<u>Learning Activities</u>





XXIV-O (Objectives)



10 Minutes



XXIV-1 ("Prevalence of Polydrug Use")

HS 172 R9/02

DRUG COMBINATIONS

- A. The Prevalence of Polydrug Use.
  - 1. Polydrug use means having two or more drugs in your body at the same time.
  - 2. It is actually more common for a DRE to encounter polydrug users than single drug users.
    - a. In the Los Angeles Field Study (1985), 72% of the suspects had two or more drugs in them.
    - b. In that study, alcohol was often found in combination with one or more other drugs.
    - c. But even if we discount alcohol, nearly half (45%) of the Field Study suspects had two or more other drugs in them.
    - d. During Certification
      Training in New York City
      in early 1989, two-thirds
      (67%) of the suspects
      evaluated had two or more
      drugs other than alcohol
      in their urine.

XXIV-1

Total Lesson Time: Approximately 110 Minutes

Session title on wallchart.

Briefly review the objectives, content and learning activities of this session.

Point out that 81 of the 173 suspects (47%) in the Los Angeles Field Study had alcohol in combination with one or more other drugs.

Aides	Lesson Plan	Instructor Notes
XXIV-2 ("Common Combina- tions")	3. Common combinations of drug	s.
	<ul><li>a. Cocaine and Cannabis.</li><li>b. Cocaine and Heroin.</li><li>c. PCP and Cannabis.</li></ul>	Point out that virtually any possible drug combinations may be encountered by the DRE.
	4. Many of the suspects you examine will be exhibiting the effects of two or more drugs acting together.	Solicit students' comments and questions about the prevalence of polydrug use.
65 Minutes	<ul> <li>B. Possible Effects of Drug Combinations.</li> <li>1. Let us examine the possible ways in which two drugs migh interact.</li> </ul>	NOTE: AT THIS TIME DRAW THE FOLLOWING MATRIX ON THE CHALKBOARD:
Pupil Size	Possible Effects of Drug Number 1  normal dilated constricted	Possible Effects of Drug Number 2  normal dilated constricted
XXIV-3 ("Neither Affects")	2. Our specific example will focus on pupil size; there are four situations that could occur.  a. Situation #1: Neither drug affects pupil size.	

Aides	Lesson Plan	Instructor Notes
4.	o drug #1 leaves pupil size in the normal range.	
	o drug #2 also leaves pupil normal.	
	o the combination also will leave pupil size normal.	
		Point out a general principle: If neither drug affects a Major indicator, the combination of those two drugs also will not affect that indicator.
XXIV-4 ("Null Effect")	b. Situation #1 is called the <b>Null Effect</b> .	Clarification of "Null Effect": The combination of no action plus no action equals no action.
	c. Specific examples of the Null Effect:	
	o Pupil Size: Neither PCP nor Valium affects pupil size; the combination of PCP and Valium will not affect pupil size.	
	o Body Temp: Neither Alcohol nor Marijuana usually affects body temperature; the com- bination of Alcohol and Marijuana usually leaves body temperature normal.	
	o HGN: Neither Cocaine nor Heroin will cause Nystagmus; the combination of Cocaine and Heroin also will not cause Nystagmus.	

Aides	Lesson Plan	Instructor Notes	
		Ask students to suggest a specific combination of drugs that will exhibit the Null Effect on Horizontal Gaze Nystagmus.  Solicit students' questions about the Null Effect.  Redirect the students' attention to our example of pupil size: point to the matrix on the chalkboard or flipchart.	
XXIV-5 ("One affects, one doesn't")  XXIV-6 ("Overlapping Effect")	d. Situation #2: one drug affects pupil size, but the other does not.  o one possibility: drug #1 dilates pupils, drug #2 leaves pupil size alone.  o another possibility: drug #1 leaves pupils, drug #2 constricts pupils, drug #1 leaves pupil size alone.  e. Situation #2 is called the Overlapping Effect.  o One example: PCP doesn't affect pupil size, but Cocaine dilates pupils; a suspect who has taken a combination of PCP and Cocaine will usually exhibit dilated pupils.	Clarification of "overlapping Effect": action plus no action equals action.	

### Lesson Plan

# Instructor Notes

- o Another example:
  Valium won't affect pupil size, but heroin will constrict pupils; a suspect under the combined influence of Valium and Heroin usually will have constricted pupils.
- f. Other examples of the "Overlapping Effect":

Ask a student to give an example of a specific combination of drugs that will produce an "Overlapping Effect" on Horizontal Gaze Nystagmus.

- o Alcohol will cause HGN, but Marijuana will not affect HGN; a person under the combined influence of alcohol and Marijuana will usually cause HGN.
- o Xanax will not affect temperature, but Demerol will lower temperature; a suspect impaired by a combination of Xanax and Demerol usually will have a lower temperature.

Ask a student to give an example of a specific combination of drugs that will produce an "Overlapping Effect" on body temperature.

Redirect the students' attention to the example of pupil size. Point to the matrix on the chalkboard.



XXIV-7 ("they affect in the same way") g. Situation #3: The two drugs affect pupil size in the same way.

Aides	Lesson Plan	Instructor Notes
. ;	o One possibility: both drugs <b>dilate</b> the pupils.	Example: Both Methamphetamine and LSD will dilate the pupils. Therefore, a person under the combined influence of Methamphetamine and LSD will have dilated pupils.
	o Another possibility: both drugs <b>constrict</b> the pupils.	Example: Both Morphine and Demerol are Narcotic Analgesics, so both constrict the pupils; someone under the combined influence of Morphine and Demerol will have constricted pupils.
XXIV-8 ("Additive Effect")	h. Situation #3 is called the Additive Effect.	Clarification of the "Additive Effect": <b>action plus the same</b> <b>action reinforces the action</b> .
	o One example: a CNS Stim- ulant plus an Hallucinogen will produce an additive effect on pupil size.	Ask a student to give an example of a drug combination that will cause an additive effect on Nystagmus.
·	o Example: a CNS Depressant plus PCP will cause an additive effect on HGN.	Ask a student to give an example of a drug combination that will produce an additive effect on blood pressure.
	o Example: PCP plus Cannabis will produce an additive effect on blood pressure.	Redirect students' attention to our example of pupil size; point to the matrix on the chalkboard.



XXIV-9 ("They produce opposite effects")



XXIV-10 ("Antagonistic Effect")

- i. Situation #4: The two drugs affect pupil size in exactly opposite ways.
  - o Either drug #1 constricts the pupils while drug #2 dilates them.
  - o Or, drug #1 dilates the pupils while drug #2 constricts them.
- Situation #4 is called the Antagonistic Effect.
- k. When two drugs produce an "Antagonistic Effect", they tend to try to cancel each other out.
  - o possibility #1: the
    effects might actually
    cancel out; e.g., the
    speedballer's pupils
    might be normal of size,
    as the Heroin's
    constriction cancels out
    the Cocaine's dilation.
  - o possibility #2: the
    Heroin might be
    exerting the stronger
    effect at some given
    moment; in this case,
    the pupils might be
    constricted, but possibly
    not as much as they
    would be if the Cocaine
    were not present.

Ask students for an example of a drug combination in which one drug dilates while the other constricts.

Clarification of "Antagonistic Effect": action versus opposite action: can't predict the outcome.

Example: When a suspect takes a "speedball" (Heroin plus Cocaine), the two drugs try to cancel out their effects on pupil size.

Aides	Lesson Plan	Instructor Notes	
,	o possibility #3: the Cocaine might be exerting the stronger effect, and the pupils might be dilated, but maybe not as much as if the Heroin weren't present.		
	o With an "Antagonistic Effect", we just can't predict what we will see.	Solicit students' questions about the Null, Overlapping, Additive and Antagonistic Effects.	
XXIV-11 ("Effects of Drug Combi- nations")	3. To summarize, when two or more drugs are taken together, they tend to produce a combination of Null Effects, Overlapping Effects, Additive Effects and Antagonistic Effects.		
XXIV-12	4. A specific Example: Consider a person who is under the influence of a combination of Cannabis and a CNS Stimulant.	Display <b>only the title</b> of <b>XXIV-12</b> ("Cannabis and a Stimulant in Combination"); you will reveal this visual one line at a time.	
		Ask students: "Will you see HGN with this particular combination?"	
	a. Neither Cannabis nor a Stimulant causes HGN.	Reveal the first line of the Visual.	
,	o This is a case of <b>no</b> action plus no action equals no action.	Point out that the combination of Cannabis and Stimulant produces a Null Effect on HGN.	
	o We will not see HGN with this combination	Ask students: "Will we see Vertical Gaze Nystagmus?"	

		_		
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### Lesson Plan

#### Instructor Notes



- b. Neither Cannabis nor a stimulant causes Vertical Gaze Nystagmus.
  - o This is another Null Effect.
  - o We won't see Vertical Gaze Nystagmus.
- c. Cannabis causes Lack of Convergence; a CNS Stimulant does not.
  - o This is a case of action plus no action equals action.
  - o We will see Lack of Convergence with this combination.
- d. CNS Stimulants dilate pupils; Cannabis either dilates pupils or leaves them alone.
  - o This may be a case of action plus no action equals action.
  - o Or it may be a case of action plus same action reinforces action.
  - o In either case, we should see dilated pupils with this combination.
- e. CNS Stimulants slow the pupils' reaction to light; Cannabis usually doesn't affect the pupils' reaction.

Reveal the second line of the Visual.

Ask students: "Will we see a Lack of Convergence?"

Reveal the third line of the Visual.

Point out that the combination of Cannabis and Stimulant produces an Overlapping Effect on Lack of Convergence.

Ask students: "What will we see when we examine pupil size?"

Reveal the fourth line of the Visual.

Point out that the combination of Cannabis and Stimulant produces either an Additive Effect or an Overlapping Effect on pupil size.

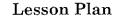
Ask students: "What should we see when we examine the pupils' reaction to light?"

Reveal the fifth line of the Visual.





Aides	Lesson Plan		Instructor Notes
; ·		o Here we have another Overlapping Effect.	
		o We should observe a slowed reaction of the pupils.	Ask students: "What should we see when we measure this person's pulse rate?"
	f.	Both Cannabis and CNS Stimulants usually elevate pulse rate.	Reveal the sixth line on the Visual.
·		o This is an Additive Effect.	
		o We will see a pulse rate higher than normal.	Ask students: "What should we see when we measure this person's blood pressure?"
	g.	Cannabis usually causes blood pressure to be above normal; so does a CNS Stimulant.	Reveal the seventh line on the Visual.
		o This is another Additive Effect.	
		o We should see a higher than normal blood pressure.	Ask students: "What can we expect to find when we check this person's temperature?"
	h.	Cannabis usually does not affect body temperature. But CNS Stimulants usually elevate temperature.	Reveal the eighth line on the Visual.
		o This is another case of action plus no action equals action.	Point out that Cannabis in combination with CNS Stimulant produces an —
		o We can expect to see an elevated temperature	Overlapping Effect on body temperature.
·		with this combination.	Solicit students' comments and questions about the Cannabis/CNS Stimulant combination.



# Instructor Notes

5. Another specific example: PCP and Heroin.

Point out that this particular combination produces no Antagonistic Effects.

Consider a person under the influence of a combination of

Display only the title on XXIV-13 ("PCP and Heroin")

a. PCP causes HGN, Heroin does not.

Ask students: "What will we see when we examine this person for HGN?"

This is an Overlapping Effect.

Reveal the first line of the Visual.

We can expect to see HGN with this suspect. Ask Students: Can we expect to see Vertical Gaze Nystagmus?

b. PCP may cause Vertical Gaze Nystagmus, especially at high doses; Heroin will not cause Vertical Gaze

Reveal the second line of the Visual.

Nystagmus.

This is another Overlapping Effect.

We may see Vertical

Ask students: "Can we expect to see a Lack of Convergence?"

Gaze Nystagmus in this suspect.

> Reveal the third line of the Visual.

- PCP causes Lack of Convergence; Heroin doesn't.
  - Effect.

Ask students: "What are we likely to see when we check the size of this suspect's pupils?"

- Another Overlapping
- We can expect to see Lack of Convergence.



Aides	Lesson Plan	Instructor Notes	
	d. PCP doesn't affect pupil size, but Heroin <b>constricts</b> pupils.	Reveal the fourth line of the Visual.	
	o This is yet another Overlapping Effect.		
	o We can expect to see constricted pupils with this suspect.	Ask students: "What are we likely to observe when we check the reaction of this suspect's pupils to light?"	
	e. PCP doesn't affect pupils' reaction to light; but Heroin usually produces "little or none visible" reaction to light.	Reveal the fifth line of the Visual.	
	o This, too, is an Overlapping Effect.  o We can expect "little or none visible" reaction in this suspect's pupils.	Point out that the combination of PCP and Heroin produces Overlapping Effects on all major eye indicators of drug impairment.  Ask students: "What can we are to find when we check	
		expect to find when we check this suspect's pulse rate?"	
	f. PCP usually causes pulse rate to be above normal; Heroin usually produces a below normal pulse rate.	Reveal the sixth line of the Visual.	
	o This is our first Antagonistic Effect.		
	o We cannot predict what this suspect's pulse rate will be.		

The pulse rate could be above normal, or below normal, or within the

normal range.

# Lesson Plan

# Instructor Notes

- This suspect's pulse rate will depend on many factors, including:
  - How much of each drug was taken.
  - How and when each drug was taken.
  - How tolerant the suspect is of each drug.
- h. PCP usually elevates blood pressure; Heroin usually lowers blood pressure.
  - This is another Antagonistic Effect.
  - We can't predict what the blood pressure will be.
  - It could be above normal, below normal or within the normal range.
- PCP usually elevates temperature; Heroin usually lowers it.
  - This, too, is an Antagonistic Effect.
  - The temperature could be above normal, or below normal or within the normal range.

Ask students: "What are we likely to find when we check this suspect's blood pressure?"

Reveal the seventh line of the Visual.

Ask students: "What are we likely to find when we check this suspect's temperature?"

Reveal the eighth line of the Visual.

Point out that the combination of PCP and Heroin produces Antagonistic Effects on all three vital signs.

Solicit students' comments and questions about the combination of Heroin and PCP.



Aides	Lesson Plan	Instructor Notes	
		Show the video tape of suspects under the influence of specific drug combinations. Point out the Null, Overlapping, Additive and Antagonistic Effects exhibited by those suspects.	
35 Minutes	C. Identifying Expected Indicators of Specific Combinations.	Direct the students' attention to the Cumulative Drug Symptomatology Matrix, found in Section XXIV of their Student's Manual. A copy also appears at the end of these lesson plans, for your reference.	
	<ol> <li>Cumulative Drug Symptomatology Matrix.</li> </ol>		
	a. The Matrix outlines the  expected results of the  drug recognition  examination for each category.	Remind students that we "never say never": and we "always avoid saying always" when it comes to signs and symptoms of drugs. The Matrix summarizes what we usually see but doesn't guarantee we will always see exactly that.	
	b. We will refer to the Matrix to help us interpret what we are likely to see when we examine drug combinations.		
	2. Worksheet Exercises	Assign the students to work in three-member teams.	
	a. Worksheet #1: PCP and Hallucinogen	Direct the students' attention to the three worksheets in their Student's Manual.	
2	b. Worksheet #2: Cannabis and CNS Depressant	Instruct the teams that they have only 15 minutes to fill out all three worksheets (5 minutes per worksheet).	

		•		
Aides	Lesson Plan	Instructor Notes  Solicit students' questions about this assignment.		
	c. Worksheet #3: CNS Depressant and CNS Stimulant			
		Tell the teams to start working. Terminate their work after fifteen minutes.		
	3. Discussion of Worksheets	For each worksheet, select a team to lead the discussion. Critique and correct the students' analyses of the drug combinations, as appropriate.		
		Solicit students' comments and questions about drug combinations.		
		·		

# **Session XXIV**

# **Drug Combinations**







# **Drug Combinations**

Upon successfully completing this session, the participants will be able to:

- Explain the prevalence of polydrug use among drug-impaired suspects and identify common combinations of drugs abused by those suspects
- Explain the possible effects that combinations of drugs can produce on the major indicators of drug impairment

Drug Evaluation & Classification Training

XXIV-0A

# **Drug Combinations**

Upon successfully completing this session, the participants will be able to:

- Define the terms "Null", "Overlapping", "Additive" and "Antagonistic" as they relate to polydrug effects
- Identify specific effects that are most likely to be observed in persons under the influence of particular drug combinations

Drug Evaluation & Classification Training

BO-VIXX

# Prevalence of Polydrug Use

In the Los Angeles Field Validation Study (1985):

- 72% of suspects had two or more drug categories in them (including alcohol)
- · 45% had two or more drugs other than alcohol

In New York City Certification Training (1989), 67% of suspects had two or more drug categories other than alcohol

Orug Evaluation & Classification Training

VYIII.

# **Common Combinations of Drugs**





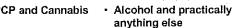


Cocaine and Cannabis . Cocaine and Heroin









Drug Evaluation & Classification Training

XXIV-2

# Two Drugs in Combination: How Do they Affect Pupil Size?

#### Situation #1

- · Neither drug affects pupil size
- Example: PCP and Valium
   Neither one affects the size of the pupils
- The combination will also not affect pupil size

Drug Evaluation & Classification Training

XXIV-

#### **Null Effect**

- · No action plus no action equals no action
- If neither drug affects a particular indicator of impairment, their combination also will not affect that indicator

Drug Evaluation & Classification Training

XXIV-

# Two Drugs in Combination: How Do They Affect Pupil Size?

#### Situation #2

- One drug affects the pupil size, but the other does not
- Example: PCP and Cocaine
   Cocaine dilates pupils, PCP doesn't affect pupils
- · The combination will affect pupil size

Drug Evaluation & Classification Training

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# **Overlapping Effect**

- · Action plus no action equals action
- If one drug affects a particular indicator of impairment, and another drug has no effect on that indicator, the combination of those two drugs will affect the indicator, in the same way as the first drug alone

Drug Evaluation & Classification Training

XXIV-

# Two Drugs in Combination: How Do They Affect Pupil Size?

#### Situation #3

- The two drugs affect pupil size in the same way
- Example: LSD and Cocaine
   Cocaine dilates pupils, and so does LSD
- · The combination will affect pupil size

Drug Evaluation & Classification Training

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### **Additive Effect**

- Action plus the <u>same action</u> produces reinforced action
- If two drugs independently affect some indicator in the same way, their use in combination will also affect the indicator, and the effect may be reinforced

Drug Evaluation & Classification Training

8-VIXX

# Two Drugs in Combination: How Do They Affect Pupil Size?

#### Situation #4

- The two drugs affect pupil size in exactly opposite ways
- Example: Heroin and Cocaine
   Cocaine dilates pupils, Heroin constricts pupils
- We can't predict how the combination will affect pupil size

Drug Evaluation & Classification Training

XXIV-9

# Antagonistic Effect

- <u>Action</u> versus <u>opposite action</u>: can't predict the outcome
- If two drugs affect some indicator in exactly opposite ways, their use in combination could affect that indicator in any possible way

Drug Evaluation & Classification Training

XXIV-10

# The Effects of Drug Combinations

- Null Effects
- Overlapping Effects
- · Additive Effects
- · Antagonistic Effects

Drug Evaluation & Classification Training

XXIV-1

# Cannabis and Stimulant in Combination

Impairment Indicator	Effect Due to Cannabls	Effect Due to Stimulant	Type of Combined Effect	What will We See?
HGN	None	None	Null	Коле
VGN	None	None	Nell	None
Lack of Convergence	Present	None	Overlapping	Present
Pupil Size	Dilated (1)	Dilated	Overlapping or Additive	Dilated
Reaction to Light	Hormal	Slow	Overlapping	Slow
Pulse Rate	Up	Up	Additive	_Up
Blood Pressure	Up	Up	Additive	Uρ
Body Temperature	Normal	Uр	Overlapping	Uρ

(1) Pupil size possibly normal

Drug Evaluation & Classification Training

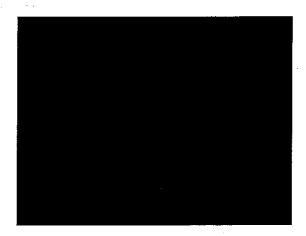
XXIV-

# Phencyclidine and Heroin in Combination

Impairment Indicator	Effect Due to Phericyclidine	Effect Due to Heroin	Type of Combined Effect	What will We See?
KGN	Present	None	Overlapping	Present
VGN	Present	Nane	Overtopping	Present
Lack of Convergence	Present	None	Overlepping	Present
Pupil Size	Normal	Constricted	Ovatimpping	Constituted
Reaction to Light	Notros	Little of None Visible	Overlapping	Little or None Visible
Puise Rate	Up	Down	Antagonistic	Down/Normal/Up
Blood Pressure	Up	Down	Antegonistic	Down/Normal/Up
Body Temperatura	Üρ	Down	Antegonistic	Down/Normal/Up

Drug Eveluation & Classification Training

XXIV-1



# CANNABIS AND STIMULANT IN COMBINATION

IMPAIRMENT INDICATOR	EFFECT DUE TO CANNABIS	EFFECT DUE TO STIMULANT	TYPE OF COMBINED EFFECT	WHAT WILL WE SEE
HORIZONTAL GAZE NYSTAGMUS	NONE	NONE	NULL	NONE
VERTICAL GAZE NYSTAGMUS	NONE	NONE	NULL	NONE
LACK OF CONVERGENCE	PRESENT	NONE	OVERLAPPING	PRESENT
PUPIL SIZE	DILATED OR NORMAL	DILATED	OVERLAPPING OR ADDITIVE	DILATED
REACTION TO LIGHT	NORMAL	SLOW	OVERLAPPING	SLOW
PULSE RATE	UP	UP	ADDITIVE	ÜP
BLOOD PRESSURE	UP	UP	ADDITIVE	UP
BODY TEMPERATURE	NORMAL	UP	OVERLAPPING	UP

# PHENCYCLIDINE AND HEROIN IN COMBINATION

IMPAIRMENT INDICATOR	EFFECT DUE TO PHENCYCLIDINE	EFFECT DUE TO HEROIN	TYPE OF COMBINED EFFECT	WHAT WILL WE SEE
HORIZONTAL GAZE NYSTAGMUS	PRESENT	NONE	OVERLAPPING	PRESENT
VERTICAL GAZE NYSTAGMUS	PRESENT	NONE	OVERLAPPING	PRESENT
LACK OF CONVERGENCE	PRESENT	NONE	OVERLAPPING	PRESENT
PUPIL SIZE	NORMAL	CONSTRICTED	OVERLAPPING	CONSTRICTED
REACTION TO LIGHT	NORMAL	LITTLE OR NONE VISIBLE	OVERLAPPING	LITTLE OR NONE VISIBLE
PULSE RATE	UP	DOWN	ANTAGONISTIC	DOWN/ NORMAL/UP
BLOOD PRESSURE	UP	DOWN	ANTAGONISTIC	DOWN/ NORMAL/UP
BODY TEMPERATURE	UP	DOWN	ANTAGONISTIC	DOWN/ NORMAL/UP

# INDICATORS CONSISTENT WITH DRUG CATEGORIES

	DEPRESSANT	CNS	HALLUCINOGEN	PCP	NARCOTIC	INHALANT	CANNABIS
HORIZONTAL GAZE NYSTAGMUS	PRESENT	NONE	NONE	PRESENT	NONE	PRESENT	NONE
VERTICAL GAZE NYSTAGMUS	PRESENT (HIGH DOSE)*	NONE	NONE	PRESENT	NONE	PRESENT (HIGH DOSE)*	NONE
LACK OF CONVERGENCE	PRESENT	NONE	NONE	PRESENT	NONE	PRESENT	PRESENT
PUPIL SIZE	NORMAL (1)	DILATED	DILATED	NORMAL	CONSTRICTED	NORMAL (4)	DILATED (6)
REACTION TO LIGHT	SLOW	MOTS	NORMAL (3)	NORMAL	LITTLE OR NONE VISIBLE	SLOW	NORMAL
PULSE RATE	DOWN (2)	UP	UP	UP	DOWN	UP	UP
BLOOD PRESSURE	DOWN	UP	UP	an.	DOWN	UP/DOWN (5)	UP
BODY TEMPERATURE	NORMAL	UP	UP	UP	DOWN	UP/DOWN/ NORMAL	NORMAL

*high dose for that particular individual

# FOOTNOTE:

These indicators are those most consistent with the category, keep in mind that there may be variations due to individual reaction, dose taken and drug interactions.

- SOMA, Quaaludes usually dilate pupils.
  - Quaaludes and ETOH may elevate.
- Certain psychedelic amphetamines cause slowing. નું જું છું
  - Normal but may be dilated.
- Down with anesthetic gases, up with volatile solvents and aerosols. 4. 70. 6
  - Pupil size possible normal

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	Marked reddening of conjunctiva Odor of Marijuana Marijuana Marijuana Marijuana Marijuana Marijuana debris in mouth Body tremors Eyelid tremors Relaxed inhibitions Increased appetite Impaired perception of time & distance Disorientation Possible paranoia	2-3 hours - exhibits effects (Impairment may last up to 24 hours, without awareness of effects.)	Smoked Oral	Fatigue Paranoia
O E IV O IV I IIV	Residue of substance around nose & mouth Odor of substance Possible nausea Slurred speech Disorientation Confusion Bloodshot, watery eyes Lack of muscle control Flushed face Non-communicative Intense headaches intense headaches ar Nones and aerosols cause above normal blood pressure; volatile solvents and aerosols cause above normal blood pressure	6-8 hours for most volatile solvents Anesthetic gases and aerosols - very short duration.	Insufflated (Historically, have been taken orally.)	Coma
NARCOTIC	Droopy eyelids ("ptosis") "On the nod" Drowsiness Depressed reflexes Low, raspy, slow speech Dry mouth Facial itching Euphoria Fresh puncture marks Nausea Track marks Nausea Track marks Nittle psychomotor impairment.	Heroin: 4-6 hours Methadone: Up to 24 hours Others: Vary	Injected Oral Smoked Insufflated	Slow, shallow breathing Clammy skin Coma Convulsion
aya	Perspiring Warm to the touch Blank stare Very early angle of HGN onset Difficulty in speech Incomplete verbal responses Repetitive speech Increased pain threshold Cyclic behavior Confused agitated Hallucinations Possibly violent & combative Chemical odor "Moon walking"	Onset: 1-5 minutes Peak Effects: 15-30 minutes Exhibits effects up to 4-6 hours	Smoked Oral Insufflation Injected Eye drops	Long intense "trip"
CANCILLAN	Dazed appearance Body tremors Synesthesia Hallucinations Paranoia Uncoordinated Nausea Disoriented Disoriented Perspiring Poor perception of time & distance Memory loss Disorientation Flashbacks  Note: With LSD, piloerection may be observed (goose bumps, hair standing on end)	Duration varies widely from one hallucinogen to another.	Oral Insuffation Smoked Injected Transdermal	Long intense "trip"
CNS STIMILI ANTS	8	Cocaine: 5-90 minutes Amphetamines: 4-8 hours Methamphetamines: 12 hours	Insuffiation (snorting) Smoked Injected Oral	Agitation Increased body temperature Hallucinations Convulsions
CNS	Uncoordinated Disoriented Sluggish Thick, slurred speech Carl ataxia Drowsiness Droopy eyes Fumbling *Note: With Wethaqualone, pulse will be elevated and body tremors will be evident. Alcohol and Quaaludes elevate pulse. Soma and Quaaludes dilate pulse. Soma and Quaaludes dilate	Barbiturates: 1-16 hours Tranquilizers: 4-8 hours Methaqualone: 4-8 hours	Oral Injected (occasionally)	Shaltow breathing Cold, clammy skin Pupils dilated Rapid, weak pulse Coma
MAJOR	GENERAL	DURATION OF EFFECTS	USUAL METHODS OF ADMINISTRATION	OVERDOSE SIGNS

### WORKSHEET #1

### PCP AND HALLUCINOGENS

IMPAIRMENT INDICATOR	EFFECT DUE TO PCP	EFFECT DUE TO HALLUCINOGEN	TYPE OF COMBINED EFFECT*	WHAT WILL WE SEE
HORIZONTAL GAZE NYSTAGMUS				
VERTICAL GAZE NYSTAGMUS				
LACK OF CONV.				
PUPIL SIZE				
REACT LIGHT				
PULSE RATE				
BLOOD PRESSURE				
BODY TEMP		·		

^{*}Null; Overlapping; Additive; or, Antagonistic

### WORKSHEET #2

### CANNABIS AND DEPRESSANT

IMPAIRMENT INDICATOR	EFFECT DUE TO CANNABIS	EFFECT DUE TO DEPRESSANT	TYPE OF COMBINED EFFECT*	WHAT WILL WE SEE
HORIZONTAL GAZE NYSTAGMUS		7		
VERTICAL GAZE NYSTAGMUS				
LACK OF CONV.				
PUPIL SIZE				
REACT LIGHT				
PULSE RATE				
BLOOD PRESSURE				
BODY TEMP				·

^{*}Null; Overlapping; Additive; or, Antagonistic

### WORKSHEET#3

### STIMULANT AND DEPRESSANT

IMPAIRMENT INDICATOR	EFFECT DUE TO STIMULANT	EFFECT DUE TO DEPRESSANT	TYPE OF COMBINED EFFECT*	WHAT WILL WE SEE
HORIZONTAL GAZE NYSTAGMUS				
VERTICAL GAZE NYSTAGMAS				
LACK OF CONV.				
PUPIL SIZE				
REACT LIGHT				
PULSE RATE				
BLOOD PRESSURE		•		
BODY TEMP				

^{*}Null; Overlapping; Additive; or, Antagonistic

### SESSION XXV

PRACTICE: TEST INTERPRETATION

### SESSION XXV PRACTICE: TEST INTERPRETATION

Upon successfully completing this session, the participant will be able to:

- o Analyze the results of a complete Drug Evaluation and Classification Examination and identify the category or categories of drugs affecting the individual examined.
- o Articulate the basis for the drug category identification.

### Content Segments

- A. Interpretation Demonstrations
- B. Interpretation Practice

### Learning Activities

- o Instructor Led Demonstrations
- o Small Group Practice
- o Participant Led Presentations

Aides	Lesson Plan	Instructor Notes
3.7.7.1 s	PRACTICE: TEST INTERPRETATION	Total Lesson Time: Approximately 45 Minutes
		Point out the "Test Interpretation" wall chart.
XXV-O (Objectives)		Briefly review the objectives, content and activities of this session.
	A. Interpretation Demonstrations	
20 Minutes		
	1. Case #1: "Subject Fief"	Direct students to review the "Subject Fief" exemplar in Section XXV of their manual.
	a. Preliminary Examination.	Review the results of the Preliminary Examination of Subject Fief.
		Ask students: "What category or categories of drugs would produce preliminary examination results consistent with this exemplar?" Probe to draw out the bases for students' responses.
	b. Eye Examinations.	Review the results of the Eye Examinations of Subject Fief.
		Ask students to discuss the category or categories of drugs that would cause these eye examination results.
	c. Psychophysical Tests.	Review the results of the Psychophysical Tests of Subject Fief.

Aides	Lesson Plan	Instructor Notes
		Ask students to discuss the category or categories of drugs that would produce these psychophysical test results.
	d. Vital Signs Examinations.	Review the results of the Vital Signs Examinations of Subject Fief.
		Ask students to discuss the category or categories of drugs that would produce these results.
	e. Dark Room Examinations.	Review the results of the Dark Room Examinations of Subject Fief.
		Ask students to discuss the category or categories of drugs that would produce these results.
	f. Other evidence.	Review the results of the examinations for injection sites and muscle rigidity, and of the final interview of Subject Fief.
		Ask students to comment on the category or categories of drugs that would be consistent with all of the evidence on this exemplar.
	g. Opinions of evaluator.	Point out that the evidence indicates that Subject Fief is under the influence of Cannabis.
į		Solicit students' questions concerning this demonstration.
	2. Case #2: "Subject Powel".	Direct students to review the "Subject Powel" exemplar.

Aides	Lesson Plan	Instructor Notes
	a. Preliminary Examination.	Review the results of the Preliminary Examination of Subject Powel.
		Ask students: "What category or cateories of drugs would produce preliminary examination results consistent with this exemplar?" Probe to draw out the bases for students' responses.
	b. Eye Examinations.	Review the results of the Eye Examinations of Subject Powel.
		Ask students to discuss the category or categories of drugs that would cause these eye examination results.
	c. Psychophysical Tests.	Review the results of the Psychophysical Tests of Subject Powel.
		Ask students to discuss the category or categories of drugs that would produce these psychophysical test results.
	d. Vital Signs Examinations.	Review the results of the Vital Signs Examinations of Subject Powel.
		Ask students to discuss the category or categories of drugs that would produce these results.
	e. Dark room examinations.	Review the results of the Dark Room Examinations of Subject Powel.

Aides	Lesson Plan	Instructor Notes
		Ask students to discuss the category or categories of drugs that would produce these results.
	f. Other evidence.	Review the results of the examinations for injection sites and muscle tone, and of the final interview of Subject Powel.
		Ask students to comment on the category or categories of drugs that would be consistent with all of the evidence on this exemplar.
	g. Opinions of evaluator.	Point out that the evidence indicates that Subject Powel is under the influence of Inhalants.
		Solicit students' questions concerning this demonstration.
	B. Interpretation Practice	
25 Minutes		
	1. Team practice.	Assign students to work in teams of 3 or 4 members.
		Tell teams that they are to review three exemplars (Subjects Jacobs, LaSalle, and Lyons). Team members are to discuss the evidence among themselves and reach a conclusion concerning the category or categories of drugs, if any.
		Teams will present their conclusions to the entire class.

Aides	Lesson Plan	Instructor Notes
	a. Review and discussion of exemplars by teams.	
The second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second secon	<ul> <li>b. Feedback of results.</li> <li>o Subject Jacobs</li> <li>o Subject LaSalle</li> <li>o Subject Lyons</li> </ul> 2. Session wrap up.	Allow teams approximately 15 minutes to review the three exemplars and reach their conclusions.  Poll the teams to determine their conclusions concerning the category or categories of drugs present in each subject.
		Offer appropriate comments concerning the teams' performance.
		Solicit students' comments and questions concerning this practice session.
		·

### DRUG CATEGORIES FOR INTERPRETATION PRACTICE

SUBJECT

CATEGORY(IES)

Fief

Cannabis

Jacobs

Inhalants

Powel.

PCP  $\underline{\text{and}}$  Cannabis

Lasalle

Narcotic Analgesic

Lyons

Hallucinogen

### **Session XXV**

Practice: Test Interpretation



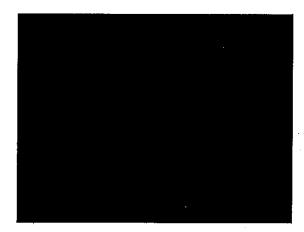
### **Practice: Test Interpretation**

Upon successfully completing this session, the participant will be able to:

- Analyze the results of a complete drug evaluation and classification examination and identify the category or categories of drugs affecting the individual examined
- Articulate the basis for the drug category identification

Drug Evaluation & Classification Training

vvi o



**Drug Influence Evaluation** Rolling Log No. DRE No Evaluator PICLIARDSON XXV-1 SANDY Recorder/Witness Crash: □ None O Fatal □ Property Injury DOB Arrestee's Name (Last, First, MI) Sex Race Arresting Officer (Name, ID No.) M 9-03-61 دن QUIGLEY. GARY BILL e Examined/Time/Location Breath Results: □ Refused Chemical Test Urine □ Blood Instrument# □ Refused 3-21.01 0.00 2330 VALLEY TRAFFIC DIVIS 012838 Miranda Warning Given: What have you eaten today? )≪Yes □ No Have you been drinking? When? How much? Time of last drink? COOKIES "FOU HRS AGO" NA SOME N/A RICHARDSON TEX When did you last sleep? How long? Are you sick or injured? □Yes ¤(No Are you diabetic or epileptic? Time now? I DON'T REMEMBER □ Yes 10 No YESRERDAY ... LAST NIGHT" (DOA" Do you have any physical defects? ☐ Yes KNo Do you take insulin? □ Yes K No Are you under the care of a doctor or dentist? □ Yes Attitude COOPERATIVE, BUT SLOW TO RESPOND Are you taking any medication or drugs? Tyes (No Coordination DISORIENTED UNSTEADY DISINITERESTED Breath STALE MOSMA Eyes: 

Reddened Conjunctiva Tracking: Speech Blindness: □ Normal Bloodshot □ Watery None □LEye. □REye 🗶 Equal 🗆 Unequal SLOW Corrective Lens: Pupil Size: Equal Unequal (explain) Able to follow stimulus: Yes □ No Eyelids: None □ Glasses □ Contacts, if so □ Hard □ Soft Normal □ Droopy LÊG TREMORS Pulse & Time HGN Left Eye Right Eye Vertical Nystagmus? ☐ Yes M No One Leg Stand 1. 112 1 2335 Lack of Smooth Pursuit Convergence NO 100Right Eye 12347 Max. Deviation ここ 20 8. 112 12357 Angle of Onset ろうひぐ Nove Romberg Balance Walk and Turn Test Cannot Keep Balance 🍛 LOWER BODY TREMORS Starts Too Soon Approx. Approx. 1st Nine 2nd Nine Sways While Balancing Stops Walking Misses Heel-Toe 7/Uses Arms to Balance Steps Off Line □ Hopping Raises Arms JJJענג Puts Foot Down Actual Steps Taken 9 Internal Clock Cannot Do Test (Explain) Describe Turn Type of Footwear 43 Estimated At 30 Sec. SAUDALS 11/1 AS NUSTRUCTED BUT SCOLL Room Light Nasal Area Pupil Size Darkness Direct EYELIB TRAMORS O Right 🛆 Left SWAYING Left Eye 5.5 Draw lines from spot touched 7.0 5.0 Oral Cavity B೬೦೦೦೧ 514, Right Eye GREEN COATIVE OU TONELE 5.5 7.0 5.0 Hippus □ Yes Rebound Dilation Reaction To Light ⋉No □ Yes XNo NORMAL 100 Blood Pressure 140 Muscle Tone: 

Normal □ Flaccid □ Rigid Attach Photos Of Fresh Puncture Marks Comments: What Medicine or Drug Have You Been Using? How Much? Time of Use? Where Were The Drugs Used? (Location) NO AUSINER NA NO ANSWER コンス・ドイド Time DRE Notified .e/Time of Arrest **Eval Start Time** Time Completed 2315 2330 3·71-01 3250 0010 3-57-01 Member Signature (Include Rank) ( N473A ID No. Reviewed By: 12-13-3 o PCP Opinion of Evaluator: □ Iohalant □ Rule Out □ Alcohol □ Stimulant O Medical □ Depressant □ Hallucinogen □ Narcotic Analgesic □ Cannabis

0753		·			
	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>			
LOG NO.	DRE: Sandy Richardson	ARRESTEE: Gary Fief			
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / DNS 6. MEDICAL PROBLEMS 7. PSYCHO I 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS			
1. LOCATION: Examin	1. LOCATION: Examination of Gary Fief took place in the DRE room, Valley Traffic Division, LAPD				
2. WITNESS: Arresting	Officer Bill Quigley, NHSP				
3. BREATH TEST: Qu	rigley administered breath test to Fief the re	esult was 0.00 and 0.00.			
4. NOTIFICATION / I	NTERVIEW of ARRESTING OFFICER	: Writer was contacted by radio and advised			
to return to Valley Traffic Division to conduct a DRE evaluation. Quigley stated he had observed the subject					
driving very slowly ((	@20 mph) without headlights and impeding	traffic.			
5. INITIAL OBSERVA	ATIONS: Writer observed the subject seated	1 int he breath testing room. Subject appeared			
passive, quiet, and se	emed uninterested in what was going on are	ound him. However, he was cooperative and			
responsive when I talked with him.					
6. MEDICAL PROBLI	EMS: None noted or stated				
7. PSYCHOPHYSICA	L TESTS: Romberg Balance: Subject swa	yed approximately 2" in a circular motion, and			
exhibited eyelid tremors, and estimated 43 seconds as 30 seconds. Walk and Turn: Subject lost his balance during the					
instructions and raised	l his arms for balance. One Leg Stand: Sub	ject raised his arms, swayed, and put his foot down.			
Finger to Nose: Subje	Finger to Nose: Subject swayed, exhibited eyelid tremors, and missed the tip of his nose.				
8. CLINICAL INDICA	TORS: Subject's pulse and blood pressure	were above the normal range. His pupils were			
dilated, there was laci	k of convergence, and reddening of the conj	unctiva.			
9. SIGNS of INGESTI	ON: Subject had a brownish - green colorat	ion on his tongue.			
10. STATEMENTS: Su	bject denied using any medication or drugs.				
11. OPINION of EVAL	UATOR: In my opinion Gary Fief is unde	r the influence of			
and unable to operate a vehicle safely.					
12. TOXICOLOGICAL	SAMPLE: Subject agreed to provide a urr	ne sample.			
13. MISCELLANEOUS	S: Throughout the evaluation subject exhibit	ted eyelid and muscle tremors.			
		,			

Evaluator ( 600 C)	Drug Influence Evaluati	
3444CBS BOB	0501 00-13-4	101
Recorder/Witness T. UNSWORTH	Crash: None Proj	
Arrestee's Name (Last, First, MI)		esting Officer (Name, ID No.)
e Examined/Time/Location	H-16-MI M Breath Results: 0.00	VSWORTH J #1811 Phoenix PD Pa Refused   Chemical Test   Surine   Blood
18-21-00 2300 Ph	DENIX PD   Instrument # 12=	∃ Cartused
Miranda Warning Given: Wes No What By: B SPARKS	have you eaten today? When? Have you ReSPONSC NO	been drinking? How much? Time of last drink?
Time now? When did you last sleep? How long? RESPONSE EAT? Some Hot dogs	Are you sick or injured?	
Do you take insulin?	Do you have any physical defects?  \( \text{Yes} \)	No Are you under the care of
NO Kesponse  Are you taking any medication or drugs? • Yes No	Didn't drink Anyth	
Are you taking any medication of drugs: 6 1es parto	NON KESPONSING PAS	Sive Staggering Stumpling
Myswers "NO" Very Slow	Odor of MARISUan	a SWEATY BLANK Stare
Speech SLOW DRAWN JOUT	Eyes:   Reddened Conjunctiva  Normal Bloodshot Watery	Blindness: Tracking:  None □ L Eye □ R Eye   Equal □ Unequal
Corrective Lens: PNone Pupil	Size: F Equal	
	qual (explain)	Able to follow stimulus: <b>S</b> Yes □ No Eyelids: SNormal □ Droopy
Pulse & Time HGN	Left Eye Right Eye Vertical Nystagmus?	Yes One Leg Stand
1. 108,2307 Lack of Smooth Pursuit	Ves Ves Convergence Right Eye	Left Eye
2. 110 12318 Max. Deviation 3. 108 12329 Angle of Onset	Yes Yes .	
Romberg Balance Walk and Turn Tes	34 134 14 4 3	
Арргох. Арргох. 3	Starts Too	1st Nine 2nd Nine Sways While Releasing
	Misses Heel	
orms Rich	Steps Medical Steps Medical Arms	intopio
The state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the s	m m m m m new Est Agryal Step.	s Taken 9 9 M/Puts Foot Down
Internal Clock Describe Turn		
Estimated At 30 Sec. FRONT F	OOT Stationary NA	
HAD TO BE ORIght A Left	Left Fue	Darkness Direct Nasal Area
Reminded Draw lines from spot touched	Right Eye 5.5	7.5 50-7.5 Oral Cavity Small Bits
TOURN SIGN	5.5	7.5 50-7.5 OF Green Leafy Dilation Reaction To Light
Lower, Arms.	• No • Yes	NORMAL
2	RIGHT ARM	LEFT ARM
		BLEM
OPana Barril Picil	<u>X6</u>	- Da
ARMS VERY RIGID  Blood Pressure 148 / 102 Temp	99.8	
Muscle Tone: ONormal OFlaccid Rigid Comments: VERY RIGIA	<del></del>	Photos Of Fresh Puncture Marks
What Medicine or Drug Have You Been Using?	~   ~	Vere The Drugs Used? (Location)
Nesponse Blank	DRE Notified Eval Sta	JOT Telling Yoll rt Time Time Completed
Member Signature (Inghide Rank)	DRE Notified ARREST Eval Sta	rt Time Time Completed
Opinion of Evaluator: □ Rule Out □	Alcohol	□ Inhalant
□ Medical □	Depressant Depressant Narcot	ic Analgesic 🗆 Cannabis

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	The second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second second secon	as a gradual of the first of	
	DRUG INFLUENCE EVALUATION	Page <u>2_of_2</u>	
LOG NO.	DRE: Sgt. Bob Sparks	ARRESTEE: Jerome Powel	
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / ONS 6. MEDICAL PROBLEMS 7. PSYCHO N 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS	
1. LOCATION: Exami	nation of Jerome Powel took place in the Di	RE room, Traffic Office, Phoenix Police Dept.	
2. WITNESS: Officer J	ames Unsworth, #1811 PPD		
3. BREATH TEST: W	riter administered breath test to Powel the	result was 0.00%.	
4. NOTIFICATION /	INTERVIEW of ARRESTING OFFICEF	R: Writer was present at the time of arrest.	
5. INITIAL OBSERV	ATIONS: Writer was supervising a sobriety	check point and Officer Unsworth approach a	
vehicle and initiate a	a conversation with the subject. When the su	abject exited his vehicle, he was unsteady on his feet,	
and very slow in res	ponding to Officer Unsworth's questions an	d instructions.	
6. MEDICAL PROBL	EMS: None noted or stated		
7. PSYCHOPHYSICA	AL TESTS: Romberg Balance: Subject swa	ayed approximately 3" side to side, and	
estimated 55 second	s as 30 seconds. Walk and Turn: Subject los	t his balance during the instructions, stepped off the	
line, missed heel to t	toe, stopped walking, raised his arms for bal	ance, and turned improperly.	
One Leg Stand: Subj	ect raised his arms, swayed and put his foot	down. On the second legs he could not maintain his	
balance and the test	was terminated for his safety. Finger to Nos	se: Subject missed tip of his nose each time, and kept	
his finger in contact	with the face on every trial.	·	
8. CLINICAL INDICA	ATORS: Subject had HGN, Vertical Nystag	mus and Lack of Convergence. His pulse, blood	
pressure, and tempe	erature were all elevated. His pupils were di	lated in near total darkness and exhibited rebound	
dilation.			
9. SIGNS of INGEST	ION: Subject's breath had an odor of mariji	nana and there was vegetable material on his teeth.	
10. STATEMENTS: S	ubject denied using any medication or drugs	3.	
11. OPINION of EVA	LUATOR: In my opinion Jerome Powel is	under the influence of a	
and unable to opera	ate a vehicle safely.		
12. TOXICOLOGICA	L SAMPLE: Subject agreed to provide a u	rine sample.	
13. MISCELLANEOUS:			

Drug Influence Evaluation			
Evaluator Aporthur, Randy App	DRE No 0459 Rolling Log No		
Authier, Ken Tops	Crash. A None	pperty	
Arrestee's Name (Last, First, MD)  Jacobs, 306	DOB   Sex   Race   Ai	resting Officer (Name, ID No.) Authier, Ken 5212 Sca	
		Chemical Test Urine #Blood	
	Jard County Breath Results Of Instrument # 12		
By: Authier, Ken Piz	era ventember co	u been drinking? How much? Time of last drint	
Time now? When did you last sleep? How long?  GPM Castnight, all night	1	'No Are you diabetic or epileptic?  □ Yes #No	
Do you take insulin? □ Yes → No	Do you have any physical defects? • Yes	No Are you under the care of a doctor or dentist?   Yes You	
Are you taking any medication or drugs?   Yes   T	lo Attitude Withdrawn, passi	Coordination Stumbling	
	Breath Chemical Odor	Face Flushed	
Slow, Slurred, raspy	Eyes:   Reddened Conjunctiva  Normal Bloodshot Watery	Blindness: Tracking:  ★None □ L Eye □ R Eye   ★Equal □ Unequa	
Corrective Lens: None Pupi	l Size: 考Equal nequal (explain)	Able to follow stimulus: Yes \( \text{No} \) No Eyelids:	
Pulse & Time HGN	Left Eye Right Eye Vertical Nystagmus	Ma Yes □ No One Leg Stand	
1. 102 , 0210 Lack of Smooth Pursuit		CO E	
2. 104 , 0222 Max. Deviation	Yes Yes Right Eye	Left Eye	
3. 104 , 0232 Angle of Onset	350 350	Washing To t	
Romberg Balance Walk and Turn	est Cannot Ko Starts Too	ep Balance	
2" 2" 7" 7	Stops Wall	14 Nine 2 Nine   D Sways While Balancir	
Circular instruct	Hons! Misses Her		
Sway Det	Steps Off I Raises Arn	ns Hopping	
in in	M M & S Actual Ste	188 1888 1 /// /	
Internal Clock Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn / Describe Turn /	ery slow, as Cannot Do Test (B		
PISTIOGE	dand stiff Pupil Size Room Light	Darkness Direct Nasal Area Runny nose	
○ Right △ Left		Daintsmears on fai	
Draw lines from spot touched	3.0	6.5 4.5 Oral Cavity 6.5 4.5 Clear	
N 5/40)		d Dilation Reaction To Light	
	No © Yes	NNO NORMHL	
2	RIGHT ARM	LEFT ARM	
	13 Paint Smears	- Paint Somear	
(5)	<u>\</u>	Paint Smear Smear	
Swaying			
Blood Pressure 142 , 98 Temp	, 98.8 .		
Muscle Tone: ★Normal □ Flaccid □ Rigid Comments:	Attach	Photos Of Fresh Puncture Marks	
' \ / / / / .	i	Were The Drugs Used? (Location)	
othing No ans	e DRE Notified Eval St	NO Answer  art Time Time Completed	
Member Signatur (Include Pank)	I II) No I Review		
	□ Alcohol □ Stimulant □ PCP	Jour Pacy John Dinhalant	
		tic Analgesic □ Cannabis	

0131		
	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>
LOG NO.	DRE: F/Sgt. Bill Tower	ARRESTEE: Bob Jacobs
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / ONS 6. MEDICAL PROBLEMS 7. PSYCHOI N 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS
1. LOCATION: Examin	nation of Bob Jacobs took place in the DRE	room, Howard County Police Dept.
2. WITNESS: Sgt. Ken	Authier	
3. BREATH TEST: Sg	gt. Ken Authier administered breath test to J	acobs the result was 0.00%.
4. NOTIFICATION / I	INTERVIEW of ARRESTING OFFICER	: Writer was the evaluating officer.
5. INITIAL OBSERVA	ATIONS: Sgt. Authier was at residence whe	en awaken by loud shouts and arguing voices.
Through window, wri	iter observed four individuals standing on th	ne front lawn. Three were young males, and one
female, they were sho	outing at and pushing each other. The arrest	tee was standing passively several yards away. Upon
turning on the outside	: light and exiting my residence, the two ma	les and one female fled. The arrestee subject
remained standing on	the lawn. He appeared dazed and confused	l. There was a strong chemical odor emanating from
him.		
6. MEDICAL PROBLI	EMS: None noted or stated	
7. PSYCHOPHYSICA	L TESTS: Romberg Balance: Subject swa	yed approximately 2" in a circular motion, and
estimated 90 seconds	as 30 seconds. When asked, "how long he	had been instructed to keep his eyes closed." He
stared straight ahead	for a few seconds and then said, "what? wha	at did you say?" When the question was repeated
he slowly shrugged a	nd said, "I don't know?" Walk and Turn: S	ubject lost his balance during the instructions,
stopped walking, rais	ed his arms for balance, and missed heel to	toe and stepped off the line. On several occasions
he asked,"What do y	ou want me to do next?" One Leg Stand: S	ubject could not maintain his balance and the test
was stopped for his s	afety. Finger to Nose: Subject missed tip of	f his nose each time, and kept opening his eyes.
8. CLINICAL INDICA	TORS: Subject had HGN, Vertical Nystagr	nus and Lack of Convergence. His pulse and
blood pressure were	above the normal range.	
9. SIGNS of INGESTIC	ON: Subject's breath had a strong chemical	odor. He had what appeared to be paint smears on
his nostrils, lips and r	right hand.	
10. STATEMENTS: Su	bject denied using any medication or drugs.	
11. OPINION of EVAL	UATOR: In my opinion Bob Jacobs is und	ler the influence of a
and unable to operate	e a vehicle safely.	
12. TOXICOLOGICAL	SAMPLE: Subject agreed to provide a blo	ood sample.
13. MISCELLANEOUS	S:	

**Drug Influence Evaluation** DRE No 5601 Evaluator FOOTBE/ Simms Rolling Log No. 0210 Recorder/Witness None Crash T. Herndon (HCSO) □ Fatal □ Property □ Injury Arrestee's Name (Last, First, M)
-a a le, lav Arresting Officer (Name, ID No.)

Milstead, F. Race Sex 01-21-75 W m te Examined/Time/Location Breath Results 0.00 Refused Chemical Test **∠**Urine Central Intox Instrument# □ Refused 1930 10-2-2000 1234 Miranda Warning Given: Yes 🗆 No What have you eaten today? y? When? This morning Have you been drinking? How much? Time of last drink? By: F. Milstead Corn Flakes. Nothing at all Time now? | When did you last sleep? How long? Are you sick or injured? Are you diabetic or epileptic? □ Yes ≽ No Midnight I don't remember Feelstomach sick Do you take insulin? □Yes %tNo Do you have any physical defects? 

Yes No Are you under the care of a doctor or dentist? □ Yes 🗶 No Attitude Cooperative, but slow Coordination Stumbling Are you taking any medication or drugs? Yes Do Breath appears Norma □ Reddened Conjunctiva Blindness: Eyes: Tracking: slow low rasp XNormal □ Bloodshot □ Watery **≭**None □ L Eye □ R Eye # Equal □ Unequal Pupil Size: KEqual Able to follow stimulus: 🗷 Yes 🗆 No Eyelids: Droopy □ Glasses □ Contacts, if so □ Hard □ Soft □ Unequal (explain) □ Normal One Leg Stand
Test stopped on
both attempts Left Eye Pulse & Time HGN Right Eye Vertical Nystagmus? □ Yes ■ No 1. 56 Νo Lack of Smooth Pursuit NO Convergence Right Eye Left Eye N.O Max. Deviation 2005 3.56 Angle of Onset None Walk and Turn Test Romberg Balance Cannot Keep Balance 🗸 🗸 Z Approx. Starts Too Soon □ Sways While Balancing Stops Walking 🔏 Uses Arms to Balance Misses Heel-Toe Steps Off Line □ Hopping Raises Arms Actual Steps Taken Internal Clock

Stimated At 30 Sec. Describe Turn Lost balance Cannot Do Test (Explain) Type of Footwear Work boots staggered to the right. Nasal Area clear Pupil Size Room Light Darkness Direct ○ Right △ Left Left Eye 1.5 1. 5 1.5 Draw lines from spot touched Oral Cavity Right Eye Clear 1.5 1.5 Hippus □ Yes Rebound Dilation Reaction To Light LiH/e or ₽ No □ Yes KNo none Visible scar MRA THEM (2)scabs 6 Temp 97.5. uncture wounds 110 60 Blood Pressure Attach Photos Of Fresh Puncture Marks Muscle Tone: □ Normal PFlaccid □ Rigid Neck rubbery Time of Use Hones What Medicine or Drug Have You Been Using? Where Were The Drugs Used? (Location) How Much? in clean now. Im notusing now. Im clean clean Time Completed 2010 Time DRE Notified 0/2/2000 ID No. 5211 Member Signature (Include Rank Reviewed By: DC METRO Opinion of Evaluator: Rule Out □ Alcohol □ Stimulant □ PCP □ Inhalant □ Medical □ Depressant O Hallucinogen □ Narcotic Analgesic □ Cannabis

00759		
·	DRUG INFLUENCE EVALUATION	Page 2_of _2
LOG NO.	DRE: Sgt. William Niles	ARRESTEE: Paul LaSalle
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / ONS 6. MEDICAL PROBLEMS 7. PSYCHOI 1 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS
1. LOCATION: Examin	nation of Paul LaSalle took place in the hold	ling area NRB
2. WITNESS: Arresting	Officer Frank Milstead #4443 PPD	
3. BREATH TEST: Of	fficer Milstead administered breath test to L	aSalle the result was 0.00%.
4. NOTIFICATION / I	INTERVIEW of ARRESTING OFFICER	: Writer was assisting members of the Phoenix
Police Department of	conduct a drug surveillance at Compton Terr	ace, prior to a 'Graceful Chickens' concert. Officer
Milstead had receive	ed information, that there was a very drunk i	ndividual seated near the entrance to Compton
Terrace. The subject	t appeared very sleepy and was very unstead	y while walking, even while being supported.
5. INITIAL OBSERVA	ATIONS: Writer observed subject seated in	a chair his head was flopped down against his chest
and he appeared to b	e sleeping. As he walked, he was very unst	eady unsteady and stumbling. His pupils were
constricted and his ve	oice was low, slow, and raspy.	
6. MEDICAL PROBLE	EMS: Subject indicated some nausea.	
7. PSYCHOPHYSICA	L TESTS: Romberg Balance: Subject swa	yed approximately 1" side to side, 2" front to
back, and estimated 5	8 seconds as 30 seconds. Walk and Turn: S	ubject lost his balance during the instructions,
stopped walking, miss	sed heel to toe, stepped off the line, and used	I his arms for balance. One Leg Stand: Subject was
unable to perform tes	et, and it was terminated for his safety. Fing	er to Nose: Subject missed tip of his nose each time,
His movements were	very slow, and his head was leaning forwar	d towards his chest.
8. CLINICAL INDICA	TORS: Subject had constricted pupils. His	pulse, blood pressure and body temperature were
below the normal ran	ge.	
9. SIGNS of INGESTI	ON: Subject had several old track marks on	both arms, and fresh puncture wounds on his left
hand. All three of th	ese were oozing clear fluid.	
10. STATEMENTS: Su	bject made several statements about being "	clean" and "not using now."
He repeatedly answe	ered "not sick" to questions concerning the t	se of medication. He also failed to respond to a
couple of the questi	ons	
11. OPINION of EVAL	.UATOR: In my opinion Paul LaSalle is un	der the influence of a
and unable to operat	e a vehicle safely.	
12. TOXICOLOGICAI	SAMPLE: Subject agreed to provide a uri	ne sample.
13. MISCELLANEOUS	S:	
	AU-ON-C-C-C-C-C-C-C-C-C-C-C-C-C-C-C-C-C-C-	

Drug Influence Evaluation						
Evaluator TOOPERAND STEVE	DRE No OS	9 <b>)</b> 1	Rolling Log No	124		
Recorder/Witness	Crash: NO	None     □ Injury		operty		-
Arrestee's Name (Last, First, MI)	DOB	Sex I	Race Ar	resting Officer (	Name, ID No.)	
YONS J.B.	6-1-52		WL	ewis	MARK	
e Examined/Timé/Location	WbD	Breat Instr	th Results:0-0 ument#1み	O□ Refused	Chemical Test  © Refus	<b>≇</b> Urine □ Blood sed
	have you eaten to	day? When?	Have yo	u been drinking:	? How much?	Time of last drink?
By STEVE TOLAND MI	IKSHAKE	Noo		ا سسن د	on4drink M	LA NA
Time now? When did you last sleep? How long?	Are you sick o		□ Yes 📽	No	Are you diabetic □ Yes <b>©</b> No	
Midnight Today The Do you take insulin? Tyes PNo		ny physical defect	Fine s? • Yes	иVo	Are you under th	
					a doctor or dentis	st? □ Yes 🍃 No
Are you taking any medication or drugs? • Yes • No	RAPID EIN	iotional ch	vanges La	Wating !	VERY POOR	Stumbling
G		RMAL	<u> </u>	I was a	Flushe	
Mumbled, INCoherent SHOUTIN	□ Normal □	ddened Conjuncti Bloodshot ロWat こくとみ くみだ		Blindness: -%None □ L	Еуе 🗆 R Еуе	Tracking: <b>⊕</b> Equal □ Unequal
Corrective Lens: None Pupil	Size: Equal equal (explain)	2780 571	TRE	Able to follow	stimulus: 🏕 Yes 🗆 No	Eyelids: Wide
Colleges Contacts, it so Chart I both College	squar (explain)	· · · · · · · · · · · · · · · · · · ·			· · · · · · · · · · · · · · · · · · ·	<u>OPEN</u>
Pulse & Time HGN	Left Eye Ri	ight Eye Vertic	al Nystagmus?	'□Yes & No	One Leg St	tand 1PST 1
1. 116/2110 Lack of Smooth Pursuit	NO	R	rgence ight Eye	Left Eye		S COPPE
2. <u>/08 / 2130</u> Max. Deviation	NO	NO (	<b>→</b>	(-v)	Q	
3. 1/2 / 3/45   Angle of Onset   Romberg Balance   Walk and Turn Te		one one	Cannot Ke	ep Balance V		JOARH V
Romberg Balance Approx. Approx. Approx. COULD			Starts Too	Soon	L R	Sways While Balancing
	~ TOE	رفاست	Stops Wall	_	<u> </u>	
		,	Misses Hee Steps Off I		<del>-  </del>	Uses Arms to Balance
- 1et ped		man.	Raises Arn	ıs		Hopping
5101		···	Actual Ste	<u> </u>	7	Puts Foot Down
Internal Clock    N A Estimated At 30 Sec.   Describe Turn				ixplain) LOST Nearly F	BALANCE Type of For	
TEST SYOPPED O Right \( \Delta \) Left			Room Light		Direct Nasal Area	AFERS
STRGGERE Braw lines from spot touched		Left Eye	6.5	8,5		Lear
CRYING! Draw lines nom spot touched	Luma	Right Eye	6.5		Oral Cavit	ear
nearly Oilo		Hippus □ Yes	s Reboun	d Dilation R	eaction To Light	
Fell (1)	Ŋ,	<b>₽</b> No	□ Yes	ØNo	NORMA	<u>L</u>
		AIGHT .	ARM		LEFT ARM	
				)		
4	<u>/3</u> \			$\sim \mathcal{N}_{\mathcal{O}}$		
(5)	<u>6</u>				Via.	
	Yes closed	€			RIPI	
Blood Pressure 156 / 102 Temp	Oct o	<u></u>			"For	05
Muscle Tone: Mormal   Flaccid   Rigid   Attach Photos Of Fresh Puncture Marks						
What Medicine or Drug Have You Been Using?	How Much?	Time of Use?	1 .	Were The Drugs	Used? (Location)	
usting Uncontrolled	RESPONSE DRENOTIFIED 2055	LAUGHIN	B   b	AUGHI	in G	
Member Signature (Include Rank)	2055	ID No	~ (	art Time SIOO	Time Comple	eted 50
XX/06~	Alcohol	ID No. 1586	Reviewe	su Dy. C . (	Of the	
	Depressant	□ Hallucinogen		tic Analgesic	□ Cannabis	

D	DRUG INFLUENCE EVALUATION	Page 2_of _2		
LOG NO. DRE	E: Sgt. Steve Toland	ARRESTEE: J.B. Lyons		
5. INITIAL OBSERVATIONS 6		NTERVIEW ARRESTING OFCR. PHYSICAL 8. CLINICAL INDICATORS N 12. TOXICOLOGY SAMPLE 13. MISC.		
1. LOCATION: Examination	of J.B. Lyons took place in the DRE I	room, Mesa P.D. Hdqtrs		
2. WITNESS: Arresting Offic	cer Mark Lewis #4196 MPD			
3. BREATH TEST: Officer	Lewis administered breath test to Lyc	ons the result was 0.00%.		
4. NOTIFICATION / INTER	RVIEW of ARRESTING OFFICER	: Writer was contacted by radio and advised to		
return to Hdqtrs to conduct	a DRE evaluation. Officer Lewis info	ormed me that the subject had nearly been involved		
in a head on accident.				
5. INITIAL OBSERVATION	NS: Writer observed subject seated in	the breath test room at Hdqtrs. He was talking to		
himself and laughing unco	ontrollably.			
6. MEDICAL PROBLEMS:	: None noted or stated			
7. PSYCHOPHYSICAL TES	STS: Romberg Balance: Subject swa	yed approximately 2" front to back, and 4" side to .		
side. The test was terminat	ated for the subjects safety. Walk and T	urn: Subject was unable to complete, test terminated		
stopped for the subjects saf	fety. One Leg Stand: Subject was una	ble to complete, test was terminated for the subjects		
safety. Finger to Nose: Sul	bject was unable to complete.			
8. CLINICAL INDICATOR	RS: Subject's pupils were dilated, and	his pulse, blood pressure and temperature were		
above the normal range.				
9. SIGNS of INGESTION: 1	None noted			
10. STATEMENTS: Subject	stated he had not used any drugs since	e the 60's		
11. OPINION of EVALUAT	OR: In my opinion J.B. Lyons is und	er the influence of a		
and unable to operate a ve	ehicle safely.			
12. TOXICOLOGICAL SAN	12. TOXICOLOGICAL SAMPLE: Subject agreed to provide a urine sample.			
13. MISCELLANEOUS:				
		· ·		

Fifty Minutes

## SESSION XXVI $\begin{array}{c} \text{REPARING THE NARRATIVE REPORT} \end{array}$

### SESSION XXVI PREPARING THE NARRATIVE REPORT

Upon successfully completing this session, the participants will be able to:

- o Discuss the essential elements of the Drug Evaluation report.
- o Prepare a clear and concise narrative description of the results of the Drug Evaluation.

### Content Segments

- A. Purpose of the Narrative Report
- B. Drug Evaluation Report Format
- C. Sample Report

### **Learning Activities**

- o Instructor Led Presentations
- o Interactive Discussion

Aides	Lesson Plan	Instructor Notes
XXVI-O (Objectives)	PREPARING THE NARRATIVE REPORT	Total Lesson Time: Approximately 50 Minutes Session title on wallchart. Briefly review session objectives, content and learning activities.
	A. Purpose of the Narrative Report	
10 Minutes	<ol> <li>Successful prosecution depends on how clearly, completely and convincingly the DRE presents their observations, measurements and conclusions.</li> <li>A well written, clear and convincing drug evaluation report increases the likelihood that the suspect will be convicted.</li> </ol>	
	a. Prosecutor is more likely to press the charge if the evidence is organized, clearly documented and compelling.	Point out that prosecutor's decision generally is based on the offense/arrest report and, consequently, if they cannot find the information they need, they are more likely to plea bargain or dismiss the charge.
	b. Defense is less likely to contest the charge when the report is descriptive, detailed and complete.	Point out that evidence gathered during the drug evaluation is rarely challenged because it is well documented on the evaluation form, backed up by a narrative report.



3. The standard Drug Influence Evaluation Face Sheet is <u>part</u> of your drug evaluation report; but it is <u>not</u> the entire report.

Aides	Lesson Plan	Instructor Notes
	a. The Face Sheet contains some very important information.	Point out some of the key information on the sample Face Sheet.
		Examples: o Suspect's pulse rate was below normal on the last two measurements.
		o Suspect had some evidence of Nystagmus, but no onset angle was found.
		o Suspect's eyes failed to converge.
		o Suspect's pupils were constricted.
	b. But the Face Sheet does not contain all of the important information that is available concerning this suspect.	Ask students to suggest some important information that might be available that wouldn't ordinarily appear on the Face Sheet.  Examples: o Information obtained during the interview of the arresting officer.  o Elaborate or lengthy state-
		ments made by the suspect.  o Paraphernalia found in
		suspect's possession.
4	Most importantly, the standard Drug Influence Evaluation Face Sheet is a <u>Technical Document</u> .	
	a. Trained DREs know how to complete and interpret the Face Sheet.	

### **Instructor Notes**

- b. But many prosecutor, judges, and jurors won't know how to interpret it.
- 5. It is up to you to take all of the information you work so hard to obtain, and to put it into a clear, plain English, written report so that the prosecutor, the judge and the jury will understand what you observed and what it means.
  - a. As a DRE, you have a special ability to secure powerful, scientific evidence that can make the difference between success and failure in court.
  - b. It would be a shame to waste that special ability by submitting an inadequate written report.
- 6. To ensure that the information contained on the Face Sheet is systematic and standardized the results of the tests should be recorded as follows:

### Lack of Convergence

a. A dot should be made where the pupil is and draw an arrow to indicate the movement and where the pupil stops.

### Romberg

a. The first figure indicates the sway from front to back and should be estimated in inches from center. Remind students of the K.I.S.S. principle- (Keep It Simple Stupid). While using very technical terminology is OK, the DRE must remember that it does no good to have a report that no one but them can understand.

Show the students an example. Remind them that in their student manuals is a complete description of the correct way to mark their evaluations.

Show the students an example. Remember to have them put the approximate number of inches from center the subject sways on either end of the arrows.

		т		
A	ı	d	e	S

### Lesson Plan

### **Instructor Notes**

- b. The second figure indicates the sway from side to side and is estimated in inches from center.
- c. Record actual elapsed time.

### Walk and Turn

- a. The first two categories, cannot keep balance and starts too soon, are observed during the instruction stage.
  - o On the lines indicate the number of times each cue is observed.
- b. Indicate by a check the number of times the suspect stops, misses heel to toe, steps off line or raises arms.
- c. Record the actual number of steps taken.
- d. If the suspect stops walking, indicate where with a vertical slash mark and an "S" under that mark.
- e. If the suspect steps off the line, indicate with half of a slash mark at an angle in the direction the step was off the line.
- f. If the suspect misses heelto-toe, indicate with a vertical slash mark and an "M" under that mark.
- g. Describe turn.

Demonstrate how each cue is to be documented using flipcharts or chalkboard

Aides	Lesson Plan	Instructor Notes	
	One Leg Stand  a. Indicate above the feet the number they were counting when they put their foot down.	Demonstrate how each cue is to be documented using flip charts or chalk board.	
	b. Check marks should be made to indicate the number of times the suspect swayed, used arms, hopped or put foot down.		
	c. Indicate how far the subject counted in 30 seconds in the top area of the box above the foot raised.		
	Finger to Nose  a. A line should be drawn to the appropriate triangle or circle to indicate where the suspect touched their nose.	Demonstrate how each cue is to be documented using a flip chart or chalk board.  Instructor's Note: Suggestion: If the DRE draws the line from the place where the suspect touches to the triangle it enables them to draw a straighter line.  Solicit students' comments and questions about the Purpose of the Narrative Report.	
20 Minutes	B. Drug Evaluation Report Form		
XXVI-2A ("First Seven Items")	<ol> <li>The typical drug evaluation report contains 13 major items.</li> <li>First item: the Location (i.e., where the evaluation was conducted).</li> </ol>		
Items")			

Aides		Lesson Plan	Instructor Notes
	3.	Second item: Witnesses.	
		a. List the person who served as the evaluator and the recorder with the complete agency name spelled out.	
į		b. Other officers who helped to conduct the evaluation.	
		c. Others who observed the evaluation.	Include any instructors who witnessed the evaluation
	4.	Third item: The Breath Alcohol Test.	
		a. Indicate BAC.	
		b. Who administered the breath alcohol test.	
		c. Time the test was administered.	
	5.	Fourth item: The Notification and Interview of the Arresting Officer.	
		a. When were you first notified of the request for a drug evaluation?	
		b. Summarize the information you were given at that time.	
		c. Document any information provided by the arresting officer.	
		d. Summary of your interview with the arresting officer	

and other witnesses.

- 6. Fifth item: Initial Observation of the Suspect.
  - a. Where you first saw the suspect.
  - b. Noteworthy aspects of your initial observations.
  - c. Findings of the Preliminary Examination of the Suspect.
- 7. Sixth item: Medical Problems and Treatment.
  - a. Your observations of any apparent injury or illness affecting the suspect.
  - b. Suspect's statements of injury or illness.
  - c. Summary of any medical treatment provided to the suspect.
- 8. Seventh item: Psychophysical Indicators of Impairment.
  - a. Briefly summarize performance of the Romberg, Walk and Turn, One Leg Stand and Finger to Nose tests.
  - b. Include any relevant behaviors on the tests that are not included on the face sheet.
- 9. Eighth item: Clinical Indicators of Impairment.



- a. Eye signs.
  - o Briefly summarize your observations of HGN, Vertical Gaze Nystagmus, Lack of Convergence, pupil size, reaction to light and appearance of the suspect's eyes.
  - o Document any observations of eyelid tremors
- b. Vital signs.
  - o Briefly summarize the suspect's pulse rate, blood pressure and temperature.
- c. Document if there were any body, leg or eyelid tremors present.
- 10. Ninth item: Signs of Ingestion.
  - a. Results of examinations of oral and nasal cavities.
  - b. Results of examinations for injection marks.
  - c. Odors detected on suspect's breath, hands, clothing, etc.
  - d. Physical debris of drugs or drug paraphernalia found on suspect's person.
- 11. Tenth item: Suspect's Statements.

Aides	Aides Lesson		Instructor Notes	
	a.	"Miranda" waiver and responses.	Remind students to contact their local DA's office for information on when to give	
	b.	Volunteered or spontaneous statements.	Miranda during the evaluation.	
	c.	Statements made as a result of your interview.		
		o Include admission or denial of drug use, time and location drugs were used, statements relating to the suspect's perception of their impairment if applicable.		
	12. El	eventh item: DRE's Opinion.		
	a.	State the category or categories of drugs that you believe is/are affecting the suspect.	Note: Anytime you have a positive BAC reading, you must list alcohol as part of the opinion.	
	b.	State your opinion concerning the suspect's ability to operate a motor vehicle safely, if applicable to this case.		
		velfth item: Toxicologic ample.	Note: Show students a copy of a toxicology request form that they will be using.	
	a.	State the type of sample (urine, blood, etc.) obtained from the suspect.	Remind the students that if they have a tracking number on the toxicology request form, that they should also include	
· · ·	b.	State who drew the sample or observed the collection of the sample.	that number in the report.	
	c.	State where the sample was taken and to whom it was given.		

Aides	Lesson Plan	Instructor Notes
	d. If the suspect refused to provide a sample, state that fact.  14. Thirteenth item: Miscellaneous.	
	a. Any other pertinent information such as, drugs or drug paraphernalia found in the suspect's possession, or possibly which hand the suspect uses.	
20 Minutes	C. Sample Report	Direct the students' attention to the Sample Drug Evaluation Report (Richardson) in Section XXVI of their Student Manual.  A copy of this report is found at the end of these lesson plans, for your reference.  Briefly review all thirteen items of the report with the students.  Solicit their comments and questions about the report.
		questions about the report.

#### **Session XXVI**

#### **Preparing the Narrative Report**



#### **Preparing the Narrative Report**

Upon successfully completing this session, the participants will be able to:

- Discuss the essential elements of the drug evaluation report
- Prepare a clear and concise narrative description of the results of the drug evaluation

Drug Evaluation & Classification Training

XXVI-0

#### Sample Drug Influence Evaluation Face Sheet

#### "INSERT NEW EXEMPLAR"

Drug Evaluation & Chasilization Training

XXVI-1

#### Items on the Drug Evaluation Report

- 1. The location
- 2. Witnesses
- 3. The breath alcohol test
- 4. Notification and interview of arresting officer
- 5. Initial observations of the suspect
- 6. Medical problems and treatment
- 7. Psychophysical indicators of impairment

Drug Evaluation & Classification Training

XXVI-2A

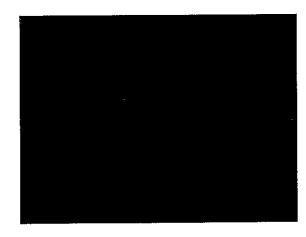
### Items on the Drug Evaluation Report

(continued)

- 8. Clinical indicators of impairment
- 9. Signs of ingestion
- 10. Suspect's statements
- 11. DRE officer's opinion
- 12. Toxicological sample
- 13. Miscellaneous

Drug Evaluation & Classification Training

XXVI-2B



Drug Influence Evaluation					
Evaluator Jim Brown, LAPD	DRE No	Rolling J	og No.		
Recorder/Witness DERALD GAUTIER, DPD		≼ None □ Injury	□ Property		
stee's Name (Last, First, MI)  LICHAROSON, JOHN M	DOB   S	Race W	Arresting Officer (N	Iame, ID No.) Гоно ≠3691	LAPD
	COVIER	Breath Resu Instrument #	ts:.00 Refused	Chemical Test	≱Urine □Blood d
Miranda Warning Given: ■ Yes □ No What	have you eaten today?	When? F	Have you been drinking?		Time of last drink?
Time now? When did you last sleep? How long?  The LAST NIGHT & YHM.	Are you sick or injur		Yes ≱4,No	Are you diabetic or □ Yes 🕱 No	epileptic?
Do you take insulin?	Do you have any phy	ysical defects? 🛛 Y	es ANo	Are you under the a doctor or dentist	
Are you taking any medication or drugs? The XNo	Attitude COOPENAT Breath	NOE/ALITE	E WITHDRAWN		TRY POOR LELY STAND
				PALE	
Speech Low, 520w, RASPY	Eyes: □ Reddene □ Normal <b>#</b> Bloods	ed Conjunctiva shot 🗆 Watery	Blindness:		Tracking: ≰Equal □ Unequal
	Size: 🕦 Equal equal (explain)		Able to follow	stimulus: <b>≠</b> Yes □ No	Eyelids: □ Normal 🙇 Droopy
Pulse & Time HGN	Left Eye Right E	* -	agmus? □ Yes □ No	One Lee Sta	6 P.O
1. 60 / 7 Lack of Smooth Pursuit 2. 58 / 9 53 P Max. Deviation	YES YES	Convergence Right Ey	ze Left Eye		
3. 58 / /0:07/2 Angle of Onset	NO NO			· 12	30 15/36
Romberg Balance Approx. Approx.  Walk and Turn To	RAISED ARM.	معجب کی۔			Sways While Balancing
		Sto	ps Walking sses Heel-Toe	- COUTIN	USUSCY RAISED USes Arms to Balance ARMS
He40		~~~~~	ps Off Line	- 0 G F	Iopping
DROPPED	m m		tual Steps Taken q	9	uts Foot Down
Internal Clock 52 Estimated At 30 Sec. Describe Turn PIVOTED	NEARLY FEZ		Test (Explain)	0,1	U 5440E5 17 <i>E</i> D
○ Right △ Left		ipil Size Room I		Direct Nasal Area	EPP
Draw lines from spot touched	\ <i>1</i>	th Eye Z.		2.0 Oral Cavity 2.0 DQY	CLEAR JPS
y olto,	Hir	ppus □Yes ¤No		eaction To Light LITT	EGR
		RIGHT ARM			TAKLES
	\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\				TO NEST
4	<del>\frac{\gamma_3}{\alpha}</del>		- N	The state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the s	3 FRESH Punctions
5 SWITCHED HANDS ON	<b>5</b>			-	
	97.8				TRACK
Comments: ARMS FET COOL TO TOO	ICH		Attach Photos Of Fi		ks Meics
at Medicine or Drug Have You Been Using?  I DONT DO DAUGS		ne of Use?	N/.	A	
	e DRE Notified		Eval Start Time 9:3	OP Time Comple	10:14 <i>P</i>
Member Signature (Include Rank) Jum Brown		No. 4731	Reviewed By:		
Opinion of Evaluator: Aule Out  Medical		Stimulant Hallucinogen	□ PCP Narcotic Analgesic	□ Inhalant □ Cannabis	

#### DRUG INFLUENCE EVALUATION NARRATIVE

- LOCATION: Evaluation was conducted in the DRE room at Jail Division -Parker Center.
- 2. WITNESSES: Evaluation was recorded by Sgt. Derald Gautier, DPD and witnessed by Officer Clark John, LAPD
- 3. BREATH ALCOHOL TEST: Officer Clark John obtained an 0.00 BrAC reading from the defendant at 9:25P.
- 4. THE NOTIFICATION AND INTERVIEW OF THE ARRESTING OFFICER: At approximately 9:30P Officer John requested that writer conduct a DRE evaluation on defendant John M. Richardson (DOB 09/06/60). Richardson had been arrested by Officer John for DUI. Impairment was not consistent with the 0.00 BrAC obtained from the defendant. Officer John also stated he stopped the defendant after observing him commit numerous traffic violations. Officer John stated that Richardson appeared sleepy, "on the nod," and that his voice was low in volume, raspy in tone and slow in tempo. The defendant failed to perform the psychomotor tasks of the SFSTs as demonstrated.
- 5. INITIAL OBSERVATION OF SUSPECT: Writer first observed Richardson in the DRE room at approximately 9:35P. Defendant walked very slowly, staggered and stumbled without falling. Richardson swayed constantly while standing and having the handcuffs removed. He responded to all questions in a low, slow, raspy, voice. His eyelids were droopy, and his pupils appeared constricted. The first pulse was 60 BPM.
- 6. MEDICAL PROBLEMS AND TREATMENT: Suspect claimed no illness or injury. No evidence of injury or illness observed.
- 7. PSYCHOPHYSICAL: Richardson exhibited impairment throughout all portions of the psychophysical exams. Romberg: He exhibited a 2" sway front to back and a 3" sway side to side. His head repeatedly dropped forward throughout the test. Had a slow internal clock; estimated 52 seconds as 30 seconds. Walk and Turn: He lost balance during the instruction stage, missed heel to toe 3 times during the first nine steps and 3 times during the second nine steps. Turned incorrectly, pivoting and nearly fell. Defendant raised his arms almost continuously throughout the test. One Leg Stand: Richardson counted very slowly throughout the test, 1000-12 in 30 seconds while standing on his left foot and 1000-15 in 30 seconds while standing on his right foot. He put is foot down 3 times while standing on his left foot and 2 times while standing on his right foot. Additionally, he swayed while trying to balance and used his arms almost continuously to balance. Finger to Nose: Richardson responded to commands very slowly, used the wrong hands on #5 and #6. He did not touch the tip of his nose on any of the six attempts.

- 8. CLINICAL INDICATORS: EYES: Lack of smooth pursuit was observed in both eyes. No additional Horizontal Gaze or Vertical Gaze Nystagmus clues were observed. Lack of Convergence was observed. Defendant's pupils were constricted in all three light conditions, little or no visible reaction to light and the eyelids were droopy. VITAL SIGNS: Richardson's last two pulse readings (58 and 58 bpm), were below the normal range. His systolic blood pressure reading (114/78) was also below the normal range.
- 9. SIGNS OF INGESTION: Three fresh puncture wounds were found on the back of the Defendant's left forearm. Numerous scar lines (track marks) were also observed on the left forearm. Muscle tone was flaccid and Richardson's arms felt cool to the touch.
- 10. SUSPECT'S STATEMENTS: Richardson denied any drug usage. He stated he was right-handed and the puncture wounds on back of his left forearm were from thorns scratches he received while gardening.
- 11. DRE'S OPINION: It is my opinion that Richardson is under the influence of a Narcotic Analgesic and is unable to operate a vehicle safely.
- 12. TOXICOLOGICAL SAMPLE: A urine sample was obtained from Richardson at 10:30P, witnessed by writer and Officer John. Sealed sample was delivered to the Evidence Property Room, to be forwarded to the Forensic Laboratory for analysis.
- 13. MISCELLANEOUS: Three syringes with needles were found by Officer John in Richardson's vehicle.

One Hour and Thirty Minutes

#### SESSION XXVII

PRACTICE: TEST ADMINISTRATION

#### SESSION XXVII PRACTICE: TEST ADMINISTRATION

Upon successfully completing this session, the participants will be able to:

- o Administer selected portions of the battery of examinations that constitute the Drug Evaluation and Classification Process.
- o Articulate the examination procedures.
- o Document the results of the evaluations.

$\underline{\text{Cont}}$	ent Segments	Lear	ning Activities
A.	Procedures for This Session	0	Participants' Hands On Practice
В.	Hands On Practice	0	Instructor Led Coaching
C. *	Session Wrap Up	0.	Participant Led Coaching

#### Lesson Plan

#### **Instructor Notes**





(Objectives)



15 Minutes

PRACTICE: TEST ADMINISTRATION

A. Procedures for This Session

1. Participants will work in two or three member teams.

- a. At any given time, one member of the team will be engaged in conducting and recording examinations of another member.
- b. The third member of the team will help coach and critique the student who is conducting the examinations.
- Students will take turns serving as test administrator, test subject and coach.
- 2. For this practice session, each student will conduct a <u>complete</u> Drug Evaluation and Classification Examination.

Total Lesson Time: Approximately 90 Minutes

Point out "Practice Session" wall chart.

Briefly review the objectives, content and activities of this session.

NOTE: Three member teams are preferable. However, no four member teams should be constructed. Thus, for example, if the class has 25 students, assign 7 three member teams and 2 two member teams.

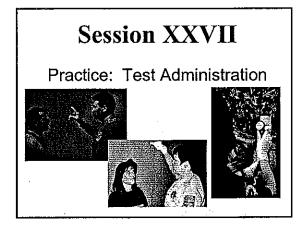
Make team assignments.

Emphasize that students can help each other learn by pointing out errors of omission or commission.

<u>Instruct</u> students to review the Standardized Drug Evaluation and Classification Form in their student's manuals.

Aides	Lesson Plan	Instructor Notes
	a. Begin with the Pr Examination. o <u>Ask</u> all of the	
	questions.  o <u>Conduct</u> the i check of the e	
	o <u>Check</u> the pub first time.	lse for the  Point out that the student who is "coaching" should simultaneously take the subject's pulse along with the test administrator.
	b. Conduct the tests Horizontal Gaze Nystagmus, Verti Nystagmus and I Convergence.	conducting the HGN test, the "coach" should check the
	<ul> <li>c. Administer the foattention psychop tests.</li> <li>o Walk and Turo One Leg Stano Romberg Balao Finger to Nos</li> </ul>	necessary for the student (subject) actually to perform these tests (except for Finger to Nose). It will suffice for the student (administrator) simply to give the test instructions
	d. <u>Check</u> the pulse for second time.	or the
	e. Check the vital si	gns.
	o Blood Pressur o Temperature o Check the pul third time.	

Aides		Lesson Plan	Instructor Notes
		f. Conduct the dark room examinations.	Point out that, for this practice session, these examinations will not actually be given in the dark.
		g. Check for muscle rigidity.	
		h. Examine the student (subject's) neck and arms for signs of injection.	Solicit students' questions concerning procedures for this practice session.
	В.	Hands On Practice	Instruct students to begin their practice.
60 Minutes			Monitor the teams, and offer encouragement and constructive criticism, as appropriate.
			Make sure each student serves as the test administrator for at least one complete drug evaluation and classification examination.
15 Minutes	C.	Session Wrap Up	Offer appropriate comments and observations about the students' performance.
			Solicit students' comments concerning this practice session.
·			



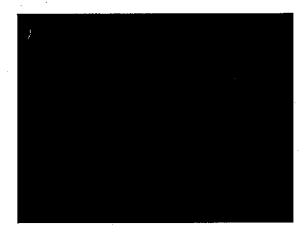
#### Practice: Test Administration

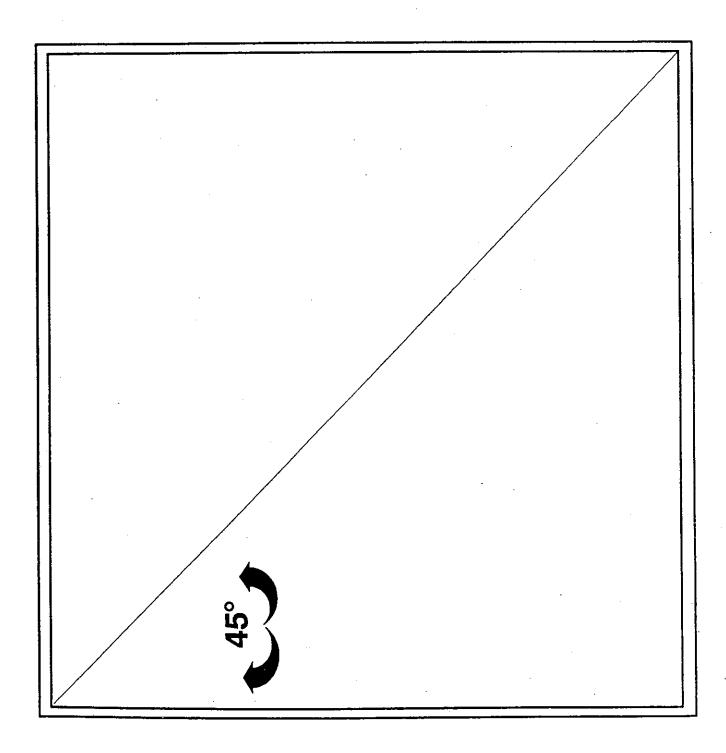
Upon successfully completing this session, the participants will be able to:

- Administer selected portions of the battery of examinations that constitute the drug evaluation and classification process
- · Articulate the examination procedures
- · Document the results of the evaluations

Drug Eveluation & Classification Training

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One Hour and Thirty Minutes

## SESSION XXVIII CASE PREPARATION AND TESTIMONY

#### **SESSION XXVIII**

#### CASE PREPARATION AND TESTIMONY

Upon successfully completing this session, the participant will be able to:

- o Conduct a thorough pre-trial review of all evidence and prepare for testimony.
- o Provide clear, accurate and descriptive direct testimony concerning Drug Evaluation and Classification examinations.
- o Respond effectively and appropriately to cross examination in Drug Evaluation and Classification cases.

## Content Segments Learning Activities A. Guidelines for Case Preparation o Instructor Led Presentations B. Guidelines for Direct Testimony o Instructor Led Demonstrations C. Typical Defense Tactics o Reading Assignments

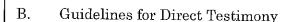
Aides	Lesson Plan	Instructor Notes
4,31	CASE PREPARATION AND TESTIMONY	Total Session Time: Approximately 90 Minutes
		Session title on wallchart.
XVIII-O		Overview session objectives, content segments and learning activities.
(Objectives)		
	A. Guidelines for Case Preparation	
10 Minutes	1. Preparation	·
	a. Preparation to present your case in court begins during your initial investigation.  o The quality of your investigation and documentation will ultimately determine your ability to	Point out: That it is especially important to take complete and accurate notes of your investigation and observations. Complete documentation of this information is essential.
	accurately present information during trial.	
·	b. When you receive the trial notice you should:	Schedule a pre trial conference with the prosecutor.
	o Review all records and reports associated with the case.	
	o Review all evidence and your conclusion.	
	o Review notes with arresting officer.	
	o Review any weak areas.	
	o Clarify or resolve any discrepancies.	

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5

#### Lesson Plan

#### Instructor Notes

- Review questions the prosecutors will be asking.
- Review tactics the prosecutors expects the defense to use.
- Review your resume and credentials.
- 2. If a pre trial conference is not possible, identify the main points of the case and discuss them with the prosecutor during the few minutes before the trial.
- Contact the DRE Agency Coordinator to discuss any new findings regarding drug categories.



- Direct testimony
  - Although knowledge only greater than what the public has is required to qualify as an "expert", your testimony will carry much more weight if you have good credentials.

Note: It is very important to meet with prosecutors that have never been exposed to the DRE program before trial to explain that it can not be treated like a typical DUI trial. You must explain that there are different protocols for DUI versus DRE cases (see Phoenix Prosecutors Training Manual).



- b. Qualifications will be established during Voir Dire:
- Point out that officer's resume is invaluable in establishing credibility.

Voir Dire is a french expression literally meaning "to see, to say". Loosely, this would be rendered in English as "To seek the truth", or "to call it as you see it". In a law or court context, one application of voir dire is to question a witness to assess his or her qualifications to be considered an expert in some matter pending before the court.

Highlight fact that you were selected to attend specialized DRE training, not just assigned randomly.

Being qualified as an expert in the past does not automatically qualify you as an expert in particular court or case.

When testifying, relate

training and experience

to the type of arrest

PCP, Cocaine, etc.)

being tried (e.g. DWI,

Point out that officers should document all previous cases where they were qualified as an expert.

If possible, do not allow defense to stipulate that you are an expert.

Point out that if your credentials are good you should always try to get your specific qualifications in front of the jury.

Document and record all evaluations conducted. Establish ratio of evaluations that resulted in a finding that subject was not under the influence.

Point out that if evaluation is properly conducted officers will be able to determine source of impairment accurately.

It is essential to demonstrate to the jury that you are fair and impartial, and that you look at each case individually.

- o Highlight the number of times you have seen a person under the influence of the drug(s) in question and have observed the symptomatology, etc.
- o Ability to answer specific questions with confidence, skill and exactness will bolster professional image in the eyes of judge and/or jury.

#### 2. New Scientific Principle

- o The scientific principles are unfamiliar to the jury or judge.
- o Your task is to establish that your hard work through training will be acceptable in the court.
- o The landmark case "Frye vs. U.S." 293F 1013 (D.C. Cir. 1923).
- o Frye requires that the scientific principle or theory used to support "evidence" be in conformity with a generally accepted explanatory theory, if the "evidence" is to be admissible.

#### 2. General guidelines.

a. Basic job is to prove that suspect was under the influence of a drug or some combination of drugs. Point out that this is critical in establishing credibility.

Point out that minor details are important.

Point out that they aren't really new just not within the common realm of knowledge of the average person.

Point out it is not enough that qualified experts testify that a particular scientific technique is valid. The technique must be generally accepted by the relevant scientific community.

Keep this in mind at all times.

Aides	Lesson Plan	Instructor Notes
	b. Don't be afraid to say "I don't know".	Point out that officer is not expected to be an expert on <u>all</u> aspects of <u>all</u> drugs. Testify to only what you know. Remember, an expert witness can rely on hearsay to develop his or her expertise.
	c. Avoid contact with the defense attorney if possible.	
	d. Don't be upset if prosecutor and defense attorney appea friendly to each other.	
	e. Jury focuses on an officer's demeanor more than content of testimony.	Point out that an officer should be polite and courteous during testimony. Do not become agitated as a result of defense questions. Do not take personal issue with defense statements, stick to the facts.
	f. Do <u>not</u> bring manuals or articles into court for reference.	Review materials before court to become familiar with contents.
	g. Explain technical terms in layman's language.	For example, HGN means an involuntary jerking of the eyes occurring as the eyes gaze to the side.
	h. Pay attention to what evidence or testimony can be and is excluded.	Point out that if officer testifies on subject matter that was excluded, it could result in a mistrial.
	<ul> <li>i. When describing suspect's performance on SFST's,</li> </ul>	Point out that the terms "pass" or "fail" should not be used.

Describe actual performance.

The defense will try to trip you

up on this point...there are no passing or failing marks.

explicitly describe exactly

what the suspect did or

neglected to do: <u>don't</u> use the terms "pass" or "fail."

Aides	Lesson Plan	Instructor Notes
·		Results of suspect's performance are describable evidence.
		Be sure to emphasize that <u>all</u> evidence is taken into account before forming an opinion.
	j. If defense attorney asks a "why" question, take the opportunity to explain in great detail if appropriate.	Point out that this suggestion does not mean that the officer should embellish his or her testimonybe careful not to open any doors for the defense.
		Note: See attachment for typical defense questions.
	C. Typical Defense Tactics	
45 Minutes	The defense relies on several factors to "impeach" or discredit your testimony.	Point out that the defense attorney's job is to try to create a "reasonable doubt". Don't take it personally.
	a. By impeaching your credibility:	
	o inconsistencies	Arresting officer's and examining officer's testimony must be complimentary. Any differences <u>must</u> be explained.
		Get your facts straight and stick to them.
	o comparison with past testimony	Try to get copies of transcripts of previous trials to review your strong/ weak points. If possible, review your testimony with the prosecutor.
	o testimony that is at odds with other established experts	Do your homeworkreview the literature. Explain any differences if possible.

Aides	Lesson Plan	Instructor Notes
	o lack of recall	Try to be prepared, but don't be afraid to say "I don't know". Be honest.
	2. By exposing the court to alternative conditions which account for your observations.	
	o another drug, o alcohol, o sickness, o injury, o other	
	3. Defense will challenge your credentialsa bona fide expert has both formal training resulting in a high degree of knowledge and experience in applying that knowledge, resulting in a skill.	Point out that if the defense can discredit your training and/or experience your testimony will have little "weight" with the jury.
	a. By directly deprecating formal training and experience.	The trial tactic is to show that the officer does not have the expertise to accurately diagnose the cause of intoxication/impairment because of inadequate formal training which lessens the value of his field experience and increases likelihood that he is mistaken in his conclusion.
	b. By demonstrating the officer lacks depth of knowledge in the drug field by contrasting his or her knowledge with the defense expert's knowledge.	
	c. By demonstrating that the officer incorrectly performed part of the evaluation, resulting in an erroneous conclusion.	Point out that the evaluation should be performed "by the book" each and every time it is conducted.

- 4. Role of defense expert.
  - To impeach credibility of the arresting officer and/or the prosecution expert.
  - b. To present alternative conditions and states that could have produced the same or similar symptoms.
- 5. Typical defense questions.
  - a. Pupillary examination in a drug case:
    - o Where the examination took place.
    - o How dark was the examining room.
    - o The size or power of the flashlight.
    - o Where the defendant was placed in relationship to the examiner.
    - o Where the flashlight was directed during the examination.
    - o Where the defendant was looking during the examination.
    - o How many times each pupil was checked.
  - b. Describe the difference between a fresh puncture site; up to 10 hours old; a 24 hour old site; a 36 hour old site; and, a 48 hour old site.

My expert v. your expert. Usually they are 180 degrees apart in their opinions.

The instructor should develop this section based on his or her personal experiences. The sample questions concerning a heroin case are based on "How To Use The Expert Witness In A Narcotic Case" by Donald M. Trookman, MD. It may be beneficial to conduct a role play cross examination to demonstrate typical questions.

Point out that the maximum is about 10 hours.

Solicit students' comments and questions concerning case preparation and testimony.

Aides	Lesson Plan	Instructor Notes
	<ul> <li>c. Are there any physical illnesses or conditions that manifest the same signs as heroin intoxication, and describe a few.</li> <li>d. How long does an occasional heroin user remain under the influence of the drug after injection?</li> </ul>	Point out that the list of possible answers is almost interminable.  SUGGESTED ROLE PLAY TO DISCUSS THE FOLLOWING QUESTIONS.  What is a DRE?  What is involved in the training program?  How do you properly identify the categories or category?  How do you explain the opinion?  What are the components of an evaluation?  Tell the students to open their Manuals to the end of Session XXVIII, and find the "Self-Test for Review". Point out that it is very similar in content and format to the Knowledge Examination they will take on the last day of this school. Also point out that the answers to the "Self-Test" appear in the Manual, on the pages immediately following the test.  Emphasize that the students do not have to "take" this "Self-Test": the decision is strictly up to them. But point out that they may find it to be a useful study aid to prepare for the final examination.

#### **Session XXVIII**

Case Preparation and Testimony



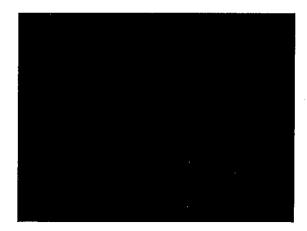
#### **Case Preparation and Testimony**

Upon successfully completing this session, the participant will be able to:

- Conduct a thorough pre-trial review of all evidence and prepare for testimony
- Provide clear, accurate and descriptive direct testimony concerning drug evaluation and classification examinations
- Respond effectively and appropriately to crossexamination in drug evaluation and classification cases

Drug Evaluation & Classification Training

XXVIII-0



#### DRE DEFENSE CROSS EXAMINATION QUESTIONS

The following are representative of questions the defense may use to challenge the Drug Recognition in court. (The Defendant is identified as Miss Alicia Ann Ace.)

#### Missing Symptoms/Normals

This line of questions attempts to elicit the fact that the defendant did not have all of the expected signs or symptoms of the drug (s) in question.

Officer, you were taught that bruxism or grinding of the teeth is a sign of CNS Stimulant influence, isn't it? Miss Ace didn't have that sign, did she?

The defense may also focus on those signs or symptoms that were normal, and were therefore, not consistent with the drug in question.

Officer, you learned the normal range of temperature in DRE training, didn't you? And that range is 98.6 plus or minus one degree, isn't it? What was Miss Ace's temperature? (98) 98 is within normal ranges, isn't it? Miss Ace's temperature was normal, wasn't it? CNS Stimulants cause elevated temperature, don't they? Miss Ace's was not elevated, was it?

#### Alternative Explanations

The defense elicits alternative explanations for the signs and symptoms of the drug (s) in question. These alternative explanations usually deal with medical conditions, stress, a traffic crash, etc.

Officer, an elevated pulse rate can be caused by things other than drugs, can't it? Excitement may cause it? Stress may cause it? Being involved in a traffic crash is stressful, isn't it? And being involved in a traffic crash may cause elevated pulse, right? Being interviewed in the early morning by three police officers is stressful? And that may also cause the pulse to be elevated, can't it?

#### **Defendant's Normals**

The defense attempts to emphasize the fact that nor everyone is so-called normal, that normal is subjective.

Officer, you were taught the normal range for pulse in DRE training, weren't you? And you agree that not all people fall in that normal range, don't you? That there are people with pulse rates above normal that aren't on drugs, right? A person's pulse changes over time, doesn't it? You don't know what Miss Ace's normal pulse is, do you? It could be in the normal range, right? But it could be above or below the normal range - normally for her, isn't that so?

#### **Doctor Cop**

The line of questioning challenges the credibility of the officer's teachers - that they are police officers, rather than medical professionals.

Officer, the teachers in this DRE school weren't doctors, were they? They weren't nurses either? Toxicologists? Pharmacologists? Paramedics? They were police officer, right?

#### Just a Cop

This line of questioning challenges the DRE's credentials - that they are "just a cop." This infers that the DRE evaluation is an ersatz medical evaluation that should be undertaken only by a medical professional.

Officer, you're not a doctor, are you? A toxicologist? A pharmacologist? A nurse? A physiologist? You don't have a degree in chemistry, do you? You're a police officer, right?

#### The Unknown

By causing the officer to state that they don't know how a sign or symptom is caused, the defense attacks the officer's credibility. This line of questioning challenges the officer's expertise, by implying that a real expert would know these things.

Officer, you don't know how CNS Stimulants dilate the pupil, do you? You don't know how alcohol supposedly causes Nystagmus, do you? You don't know how CNS Stimulants supposedly elevate the heart rate, do you?

#### Guessing Game

This tactic attacks the DRE opinion as a subjective guess, a belief, rather than objective. And guesses can be wrong.

Officer, your opinion in a DRE case is subjective, isn't it? It's a belief on your part? You've made these beliefs in DRE cases in the past, haven't you? A sometimes toxicology didn't find the drug you predicted, isn't that so? And, in fact, sometimes, toxicology didn't find any drug, isn't that so? And so, sometimes your opinion is not correct, right? Sometimes, you guess wrong?

Two Hours and Thirty Minutes

REVIEW OF THE DRE SCHOOL

#### REVIEW SESSION

The principal purpose of the Review Session is to help students prepare for the final written examination. The following questions and exercises can be posed to the class to cover all of the information that will be elicited on the final exam. Try to involve all of the students actively in these questions and exercises.

Aides	Lesson Plan	Instructor Notes
	1. HOW DO WE DEFINE THE TERM "DRUG" FOR DRE PURPOSES?	Key Points to Emphasize: o any substance
RS-1		o that impairs the ability to operate a vehicle
RS-2	2. BASIC DRUG STATISTICS:	
RS-3		
	a. What percentage of DWI arrests involve drugs other than alcohol?	LAPD Estimate: 10-20%
	b. What drug other than alcohol was found most frequently in the Los Angeles Field Validation Study?	Answer: PCP
	c. What does "polydrug use" mean?	Having two or more drug categories in your body at the same time.
RS-4	d. How common was polydrug use in the field validation study?	72% of the suspects had two or more drug categories in them.
	e. How good were the DREs in the Field Validation Study?	o Nearly 80% of the time when the DREs said a particular category of drugs was present, that category was found in the suspect's blood.
· .		o In more than 90% of the suspects, the DREs correctly identified at least one of the categories that were present.
	f. In the University of Tennessee Study, what percentage of	40% of those drivers had

evidence of other drugs in their

urine.

injured drivers had drugs other

than alcohol in them?

RS-5

Lesson Plan	Instructor Notes
3. REVIEW OF SYMPTOMATOLOGY	SOLICIT STUDENTS' QUESTIONS ABOUT DRUG STATISTICS  Prepare a "symptomatology matrix" on the chalkboard:
HGN VGN CONV PULSE	BP TEMP PUPILS
<ul> <li>a. Name six different CNS Depressants.</li> <li>b. Name four different CNS Stimulants.</li> <li>c. Name two naturally-occurring Hallucinogens.</li> <li>d. Name four different synthetic Hallucinogens.</li> <li>e. Name a major analog of PCP.</li> <li>f. Name the three sub-categories of Inhalants.</li> <li>g. What is the active ingredient in Cannabis?</li> </ul>	Ask students to "fill in" the matrix by stating how each category will affect these major indicators of impairment.  Write students' responses on the chalkboard.
	<ul> <li>a. Name six different CNS Depressants.</li> <li>b. Name four different CNS Stimulants.</li> <li>c. Name two naturally-occurring Hallucinogens.</li> <li>d. Name four different synthetic Hallucinogens.</li> <li>e. Name a major analog of PCP.</li> <li>f. Name the three sub-categories of Inhalants.</li> </ul>

Aides	Lesson Plan	Instructor Notes
:		SOLICIT STUDENTS' QUESTIONS ABOUT DRUG CATEGORIES & SYMPTOMATOLOGY.
RS-7	4. REVIEW OF VITAL SIGNS	
	a. Pulse Rate	
	(1) Define "Pulse".	Contraction and expansion of an artery, generated by the pumping action of the heart.
	(2) True or false: Pulse rate is measured in units of "millimeters of mercury".	FALSE: pulse rate is measured in "beats per minute".
RS-8	(3) Name three different pulse points, and indicate where they are located.	Make sure that students point out the Radial, Brachial and Carotid pulse points.
	(4) What is the "normal" range of adult human pulse rate, for DRE purposes?	60-90 beats per minute.
	b. Blood Pressure	
RS-9		
	(1) Define "Blood Pressure".	The force that the circulating blood exerts on the walls of the arteries.
• .	(2) Name the instrument used to measure blood pressure.	SPHYGMOMANOMETER: Ask a student to spell this, and write the correct spelling on the chalkboard.
	(3) When does blood pressure reach its highest value? What is the highest value called?	The <u>systolic</u> pressure is reached when the heart contracts and pushes blood into the arteries.

Aides	Lesson Plan	Instructor Notes
RS-10	(4) When does blood pressure reach its lowest value? What is the lowest value called?	The <u>diastolic</u> pressure is reached when the heart is fully expanded.
	(5) What is the "normal" range of adult human blood pressure, for DRE purposes?	Systolic: 120-140 Diastolic: 70-90
RS-11	(6) What does "Hg" stand for?	Chemical symbol for mercury ("Hydrargyrum", latin word for "Mercury"). Blood pressure is measured in millimeters of mercury.
		SOLICIT STUDENTS' QUESTIONS ABOUT VITAL SIGNS.
RS-12	5. REVIEW OF THE EYE EXAMINATIONS	
	a. Horizontal Gaze Nystagmus	
RS-13	(1) What are the three validated clues of impairment that have been established for HGN?	o Lack of Smooth Pursuit o Distinct Jerking at Maximum o Onset Within 45 Degrees
	(2) What formula expresses the approximate statistical relationship between BAC and onset angle?	BAC = 50 - Angle
	(3) What categories of drugs usually will cause HGN?	o CNS Depressants o PCP o Inhalants
	b. Vertical Gaze Nystagmus	

TRUE: All drugs that cause

Horizontal Gaze Nystagmus

Nystagmus, if the dose is large

will cause Vertical Gaze

enough.

RS-14

(1) True or False: any drug

cause Vertical Gaze

Nystagmus.

that causes HGN may also



(2) What category of drugs causes Vertical Gaze Nystagmus but not Horizontal Gaze Nystagmus?

Lack of Convergence

(1) True or false: any drug that causes nystagmus will also usually cause the eyes to be unable to converge.

(2) What category of drugs usually causes Lack of Convergence but does not cause nystagmus?

NO DRUG CAUSES VERTICAL GAZE NYSTAGMUS BUT NOT HGN.

TRUE: CNS Depressants, PCP and Inhalants usually cause the eyes to be unable to converge.

CANNABIS usually causes Lack of Convergence, but doesn't cause nystagmus.

SOLICIT STUDENTS' QUESTIONS ABOUT THE EYE EXAMINATIONS.



6. REVIEW OF THE DARKROOM **EXAMINATIONS** 

What are the three lighting conditions under which we must estimate the size of the suspect's pupils?

b. How long should we wait in the Darkroom before beginning to check the suspect's pupils?

Name the device that we use to estimate the size of the suspect's pupils.

d. What do the numbers on the Pupillometer refer to?

e. In what units of measurement are those number given?

f. For DRE purposes, what is the "normal" range of the size of an adult human's pupil?

o Room Light

o Near Total Darkness

o Direct Light

At least 90 seconds.



The diameters of the black circles.

The circles' diameters are given in millimeters.

The diameter of the pupil normally ranges from about 3.0 mm to about 6.5 mm.





RS-18

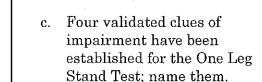
Aides	Lesson Plan	Instructor Notes
	g. What does the term "MIOSIS" mean?	"Miosis" means an abnormally small or constricted pupil, i.e., a pupil with a diameter smaller than 3.0 mm.
RS-19	h. What does the term "MYDRIASIS" mean?	"Mydriasis" means an abnormally large or dilated pupil, i.e., a pupil with a diameter larger than 6.5 mm.
RS-20	i. What category of drugs usually causes Miosis, or constricted pupils?	Narcotic Analgesics usually cause pupils to be constricted below the normal range.
	j. What categories usually cause Mydriasis, or dilated pupils?	CNS Stimulants and Hallucinogens usually cause pupils to be dilated above the normal range. Cannabis also may cause dilation. Some inhalants will also cause dilation.
	k. What is unique about the drug "Methaqualone" and SOMA?	Methaqualone and Soma is a CNS Depressant that causes dilation.
		SOLICIT STUDENTS' QUESTIONS ABOUT THE DARKROOM EXAMS.
RS-21 RS-22	7. REVIEW OF THE DIVIDED ATTENTION TESTS	
	a. Name the four Divided Attention Tests administered during the DRE Examination.	o Romberg Balance o Walk and Turn o One Leg Stand o Finger to Nose
	b. Why is the Romberg Balance always the first test administered?	<ul> <li>(1) For Standardization.</li> <li>(2) The test requires the suspect to estimate the passage of 30 seconds; thus, it should be administered before the One Leg Stand test, in which the suspect is instructed to count out</li> </ul>

30 seconds.

# Aides RS-24 RS-25RS-26

#### Lesson Plan

#### Instructor Notes



- d. How many times is One Leg Stand administered during the DRE Examination?
- e. Which foot must the suspect stand on the first time he or she performs One Leg Stand?
- f. How many validated clues of impairment have been established for the Walk and Turn test? Name them.

g. In what sequence is the suspect instructed to touch the index fingers to the nose on the Finger to Nose test?

- o Swaying
- o Raising the arms
- o Hopping
- o Putting the foot down

Twice.

The Left.

Eight validated clues.

- o Losing balance during the instructions
- o Starting to walk too soon
- o Raising arms while walking
- o Stepping off the line
- o Missing heel to toe
- o Stopping while walking
- o Wrong number of steps
- o Turning improperly

Left, Right, Left, Right, Right, Left.

SOLICIT STUDENTS'
QUESTIONS ABOUT THE
DIVIDED ATTENTION
TESTS.



#### 8. GENERAL REVIEW QUESTIONS

- a. What is the medical or technical term for "droopy eyelids"?
- b. What does "Piloerection" mean? What drug often causes piloerection?
- c. What is the medical or technical term for Heroin?

RS-7

PTOSIS.

"Piloerection" means "Hair Standing Up", or "Goose Bumps". It is often caused by LSD.

Diacetyl Morphine.

Aides	Lesson Plan	Instructor Notes
DC 99	d. Explain the terms "Null",  "Additive", "Antagonistic" and	"Null": neither drug affects some specific indicator.
RS-28	"Overlapping" Effect as they apply to polydrug use. Give examples.	"Additive": the two drugs produce some identical effects.
		"Antagonistic": the two drugs produce some directly opposite effects.
		"Overlapping": one drug affects some symptom that the other doesn't affect, and vice versa.
RS-29	e. What is the difference between "Hippus" and "Rebound Dilation"?	"Hippus" refers to pupils that pulsate rhythmically in size between fixed limits; usually, Hippus develops during withdrawal from Narcotic Analgesics.
		"Rebound Dilation" is a period of constriction followed by dilation with a change equal to or greater than 2 mm.
RS-30	f. What is the drug "Percobarb"?	It is a combination of the natural opiate Percodan with a barbiturate. Percobarb thus is a polydrug, a combination of a Narcotic Analgesic and a CNS Depressant.
	g. What does "Bruxism" mean?	Grinding the teeth.
RS-31	h. What does the number denoting the size of an hypodermic needle refer to?	The inside diameter of the needle.
	i. What does "Synesthesia" mean?	A mixing of senses, i.e., hearing colors or seeing odors.
	j. What is "Sinsemilla"?	A variety of marijuana with a high concentration of THC.

Aides

#### Lesson Plan

#### **Instructor Notes**



RS-32

k. What are the twelve major components of the DRE Examination?

List students' responses on the chalkboard.

- o Breath Alcohol Test
- o Interview of Arresting Officer
- o Preliminary Examination
- o Examinations of the Eyes
- o Divided Attention Tests
- o Vital Signs Examinations
- o Dark Room Examinations
- o Examination for Muscle Tone
- o Examination for Injection Sites.
- o Suspect's Statements
- o Opinions of the Evaluator
- o The Toxicological Exam

Ask students to describe each component briefly, and to clarify the kinds of information each component supplies.



9. REVIEW OF PHYSIOLOGY

a. Name the ten major body systems.

List students' responses on the chalkboard.

- o Muscular System
- o Urinary System
- o Respirator System
- o Digestive System
- o Endocrine System
- o Reproductive System
- o Skeletal System
- o Integumentary System
- o Nervous System
- o Circulatory System

We consciously control the Striated; we don't consciously control the Smooth.



b. What is the distinction between the "Smooth" muscles and the "Striated" muscles?

c. What do we call the chemicals that are produced by the Endocrine System?

Hormones.

Aides	Lesson Plan	Instructor Notes
	d. What is a neuron?	A nerve cell.
RS-35	e. What do we call the space between two nerve cells?	The synapse, or synaptic gap.
10.5 00	f. What do we call the chemicals that pass from one nerve cell to the next?	Neurotransmitters.
	g. What do we call the part of a nerve cell that sends out the neurotransmitter?	The axon.
RS-36	h. What do we call the part of a nerve cell that receives the neurotransmitter?	The dendrite.
	i. What do the Sensory Nerves do?	Carry messages to the brain, from the sense organs, pain sensors, etc.
	j. What do the Motor Nerves do?	Carry messages from the brain, to the muscles, etc.
RS-37	k. Name the two sub-divisions of Motor Nerves.	Voluntary (control striated muscles) and Autonomic (control smooth muscles).
	l. Name the two sub-divisions of Autonomic Nerves and describe their functions.	Sympathetic (command the body's response to fear, excitement, etc.), and Parasympathetic (promote the body's tranquil activities).
RS-38	m. What does it mean to say that a drug is "sympathomimetic"?	It means that the drug's effects mimic those caused by messages transmitted along sympathetic nerves (excite- ment, agitation, arousal, etc.).
	n. What does it mean to say that a drug is "parasympatho-mimetic"?	The drug's effects mimic those caused by messages transmitted along parasympathetic nerves (relaxation, calm, sleep, etc.).

P	1	i	d	le	9	2

#### Lesson Plan

#### **Instructor Notes**



RS-39

- o. Which two categories of drugs can most appropriately be called sympathomimetic?
- p. Which category can most appropriately be call parasympathomimetic?

Hallucinogens.

CNS Stimulants and

Narcotic Analgesics.

Clarification: Cannabis, PCP and Inhalants have some sympathomimetic characteristics, but not as many as do the CNS Stimulants and Hallucinogens. Depressants have some parasympathomimetic characteristics, but not as many as do the Narcotic Analgesics.



q. What is an artery?

Strong, elastic blood vessel that carries blood from the heart to the body's tissues and organs.

r. What is a vein?

Blood vessel that carries blood back to the heart from the tissues and organs.



RS-41

- s. What is the Pulmonary Artery, and what is unique about it?
- t. What are the Pulmonary Veins, and what is so special about them?

It is the artery that carries blood from the heart to the lungs. It is the only artery that carries blood depleted of oxygen.

They are the veins that carry blood back to the heart from the lungs. They are the only veins that carry blood rich in oxygen.

SOLICIT STUDENTS' QUESTIONS ABOUT PHYSIOLOGY.

SOLICIT ANY ADDITIONAL QUESTIONS THAT THE STUDENTS MIGHT HAVE.

Aides	Lesson Plan	Instructor Notes
		ADMINISTER QUIZ NUMBER FIVE TO THE STUDENTS. ALLOW 20 MINUTES FOR THE STUDENTS TO COMPLETE THE QUIZ. REVIEW THE QUIZ WITH THE CLASS, AND ALLOW THE STUDENTS TO RETAIN THE QUIZ FOR THEIR INDEPENDENT STUDY.
		THANK THE STUDENTS FOR ATTENDING THE OPTIONAL REVIEW SESSION.
	·	





# Review of the DRE School





R5-1

## How do we define the term "drug" for DRE purposes?

- · Definition must include:
  - Any substance
  - That impairs the ability to operate a vehicle safely

Drug Evaluation & Classification Training

RS-

#### **Basic Drug Statistics**

- What percentage of DWI arrests involve drugs other than alcohol?
  - LAPD Estimate: 10-20%
- What drug other than alcohol was found most frequently in the Los Angeles Field Validation Study?
  - PCF
- What does "polydrug use" mean?
  - Ingesting two or more distinct drug categories on the same occasion

Drug Evaluation & Classification Training

RS-3

#### **Basic Drug Statistics**

- How common was polydrug use in the Field Validation Study?
  - More than 70% of the suspects had two or more drug categories in them
- How good were the DREs in the Field Validation Study?
  - Nearly 80% of the time when the DREs said a particular category of drugs was present, that category was found in the suspect's blood.
  - In more than 90% of the suspects, the DREs correctly identified at least one of the categories that were present

Drug Evaluation & Classification Training

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## **Basic Drug Statistics**

- In the University of Tennessee Study, what percentage of injured drivers had drugs other than alcohol in them?
  - 40% of those drivers had evidence of other drugs in their urine

Drug Evaluation & Classification Training

RS-5

## Review of Symptomatology

- · Name six different CNS Depressants
- · Name four different CNS Stimulants.
- Name two naturally-occurring Hallucinogens.
- · Name four different synthetic Hallucinogens.
- · Name a major analog of PCP.
- Name the three sub-categories of Inhalants.
- · What is the active ingredient in Cannabis?

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#### **Review of Vital Signs**

- Pulse Rate
  - ~ Define "Pulse"
    - * Contraction and expansion of an artery, generated by the pumping action of the heart
  - True or false: Pulse rate is measured in units of "millimeters of mercury".
    - * FALSE: pulse rate is measured in "beats per minute"

Drug Evaluation & Classification Training

RS.

### Review of Vital Signs

- · Pulse Rate (cont..)
  - Name three different pulse points, and indicate where they are located.
    - * Radial, Brachial and Carotid pulse points
  - What is the "normal" range of adult human pulse rate, for DRE purposes?
    - * 60-90 beats per minute

Drug Evaluation & Classification Training

RS-8

#### **Review of Vital Signs**

- · Blood Pressure
  - Define "Blood Pressure".
    - * The force that the circulating blood exerts on the walls of the arteries
  - Name the instrument used to measure blood pressure.
    - Sphygmomanometer
  - When does blood pressure reach its highest value? What is the highest value called?
    - * The systolic pressure is reached when the heart contracts and pushes blood into the arteries

Drug Evaluation & Classification Training

RS-9

### **Review of Vital Signs**

- · Blood Pressure (cont.)
  - When does blood pressure reach its lowest value? What is the lowest value called?
    - * The <u>diastolic</u> pressure is reached when the heart is fully expanded
  - What is the "normal" range of adult human blood pressure, for DRE purposes?

* Systolic: 120-140mmHg * Diastolic: 70-90mmHg

RS-10

## **Review of Vital Signs**

- · Blood Pressure (cont.)
  - What does "Hg" stand for?
    - * Chemical symbol for mercury ("Hydrargyrum", Latin word for "Mercury"). Blood pressure is measured in millimeters of mercury

Drug Evaluation & Classification Training

RS-11

## Review of the Eye Examinations

- · Horizontal Gaze Nystagmus
  - What are the three validated clues of impairment that have been established for HGN?
    - * Lack of Smooth Pursuit
    - * Distinct Nystagmus at Maximum Deviation
    - * Angle of Onset of Nystagmus Prior to 45 Degrees

Drug Evaluation & Classification Training

#### Review of the Eye Examinations

- · Horizontal Gaze Nystagmus (cont.)
  - What formula expresses the approximate statistical relationship between BAC and the angle of onset of nystagmus?
    - * BAC = 50 angle
  - What categories of drugs usually will cause HGN?
    - * CNS Depressants
    - * PCP
    - * Inhalants

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RS-13

## Review of the Eye Examinations

- · Vertical Gaze Nystagmus
  - True or False: Any drug that causes HGN may also produce <u>Vertical</u> Gaze Nystagmus.
    - * TRUE: All drugs that cause Horizontal Gaze Nystagmus will cause Vertical Gaze Nystagmus, if the dose is large enough
  - What category of drugs causes Vertical Gaze Nystagmus but not Horizontal Gaze Nystagmus?
    - * NO drug causes Vertical Gaze Nystagmus but not HGN

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RS-1

#### Review of the Eye Examinations

- · Lack of Convergence
  - True or False: Any drug that causes nystagmus will also usually cause the eyes to be unable to converge.
    - * TRUE: CNS Depressants, PCP and Inhalants usually cause the eyes to be unable to converge
  - What category of drugs usually causes lack of convergence but does <u>not</u> cause nystagmus?
    - CANNABIS usually causes Lack of Convergence, but doesn't cause nystagmus

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RS-1

#### **Review of the Darkroom Examinations**

- What are the three lighting conditions under which we must estimate the size of the suspect's pupils?
  - Room Light
  - Near Total Darkness
  - Direct Light
- How long should we wait in the Darkroom before beginning to check the suspect's pupils?
  - At least 90 seconds

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RS-1

#### Review of the Darkroom Examinations

- Name the device that we use to estimate the size of the suspect's pupils.
  - The Pupillometer
- · What do the numbers on the Pupillometer refer to?
  - The diameters of the black circles
- · In what units of measurement are those numbers given?
  - The circles' diameters are given in millimeters

Drug Evaluation & Classification Training

RS-17

#### **Review of the Darkroom Examinations**

- For DRE purposes, what is the "normal" range of the size of an adult human's pupil?
  - The diameter of the pupil normally ranges from about 3.0 mm to about 6.5 mm
- What does the term "MIOSIS" mean?
  - "Miosis" means an abnormally small or constricted pupil, i.e., a pupil with a diameter smaller than 3.0 mm

Drug Evaluation & Classification Training

AS-

#### Review of the Darkroom Examinations

- · What does the term "MYDRIASIS" mean?
  - "Mydriasis" means an abnormally large or dilated pupil, i.e., a pupil with a diameter larger than 6.5 mm
- What category of drugs usually causes Miosis, or constricted pupils?
  - Narcotic Analgesics usually cause pupils to be constricted below the normal range

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RS-19

#### Review of the Darkroom Examinations

- What categories usually cause Mydriasis, or dilated pupils?
  - CNS Stimulants and Hallucinogens usually cause pupils to be dilated above the normal range. Cannabis also may cause dilation. Some inhalants will also cause dilation.
- What is unique about the drug "Methaqualone" and SOMA?
  - Methaqualone and Soma is a CNS Depressant that causes dilation

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RS-20

#### **Review of the Divided Attention Tests**

- Name the four Divided Attention Tests administered during the DRE Examination.
  - Romberg Balance
  - Walk and Turn
  - One Leg Stand
  - Finger to Nose

. Drug Evaluation & Classification Training AS-2

#### **Review of the Divided Attention Tests**

- Why is the Romberg Balance always the first test administered?
  - For Standardization
  - The test requires the suspect to estimate the passage of 30 seconds; thus, it should be administered <u>before</u> the One Leg Stand test, in which the suspect is instructed to count out 30 seconds.

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RS-22

#### **Review of the Divided Attention Tests**

- Four validated clues of impairment have been established for the One Leg Stand Test; name them.
  - Swaying
  - Raising the arms
  - Hopping
  - Putting the foot down

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RS-23

#### **Review of the Divided Attention Tests**

- How many times is One Leg Stand administered during the DRE Examination?
  - Twice
- Which foot must the suspect stand on the first time he or she performs One Leg Stand?
  - The Left

Orag Evaluation & Classification Traising

#### Review of the Divided Attention Tests

- How many validated clues of impairment have been established for the Walk and Turn test? Name them.
  - Eight Validated Clues
    - · Losing balance during the instructions
    - · Starting to walk too soon
    - · Raising arms while walking
    - · Stepping off the line
    - · Missing heel to toe
    - Stopping while walking
  - Wrong number of steps
  - · Turning improperly

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RS-25

#### Review of the Divided Attention Tests

- In what sequence is the suspect instructed to touch the index fingers to the nose on the Finger to Nose test?
  - Left, Right, Left, Right, Right, Left

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RS-2

#### **General Review Questions**

- · What is the medical or technical term for "droopy eyelids"?
  - PTOSIS
- What does "Piloerection" mean? What drug often causes piloerection?
  - "Piloerection" means "Hair Standing Up", or "Goose Bumps". It is often caused by LSD
- What is the medical or technical term for Heroin?
  - Diacetyl Morphine

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RS-27

#### **General Review Questions**

- Explain the terms "Null", "Additive", "Antagonistic" and "Overlapping" Effect as they apply to polydrug use. Give examples
  - "Null": neither drug affects some specific indicator
  - "Additive": the two drugs produce some identical effects
  - "Antagonistic": the two drugs produce some directly opposite effects
  - "Overlapping": one drug affects some symptom that the other doesn't affect, and vice versa

Drug Evaluation & Classification Training

RS-28

## **General Review Questions**

- What is the difference between "Hippus" and "Rebound Dilation"?
  - "Hippus" refers to pupils that pulsate rhythmically in size between fixed limits; usually, Hippus develops during withdrawal from Narcotic Analgesics
  - "Rebound Dilation" is a period of constriction followed by dilation with a change equal to or greater than 2 mm.

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RS-29

## **General Review Questions**

- · What is the drug "Percobarb"?
  - It is a combination of the natural opiate Percodan with a barbiturate. Percobarb thus is a polydrug, a combination of a Narcotic Analgesic and a CNS Depressant
- · What does "Bruxism" mean?
  - Grinding the teeth

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#### **General Review Questions**

- What does the number denoting the size of a hypodermic needle refer to?
  - The inside diameter of the needle
- · What does "Synesthesia" mean?
  - A mixing of senses, i.e., hearing colors or seeing odors
- · What is "Sinsemilla"?
  - A variety of marijuana with a high concentration of THC

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70.74

#### **General Review Questions**

What are the twelve major components of the DRE Examination?

- The Breath Alcohol Test
- Examination for Muscle Tone
- Interview of Arresting Officer
- Examination for injection Sites
- Preliminary Examination
- ______
- Fremiliary Examination
- Suspect's Statements
- Examinations of the Eyes
- Opinions of the Evaluator
- Divided Attention Tests
- The Toxicological Exam
- Vital Signs Examinations
- Dark Room Examinations

- Dark Nobili Examination

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PS.32

## **Review of Physiology**

Name the ten major body systems.

M is for Muscular System

U is for Urinary System

R is for Respiratory System

I is for Integumentary System

D is for Digestive System

N is for Nervous System
C is for Circulatory System

E is for Endocrine System

R Is for Reproductive System S is for Skeletal System

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RS.

## Review of Physiology

- What is the distinction between the "Smooth" muscles and the "Striated" muscles?
  - We consciously control the Striated; we don't consciously control the Smooth
- What do we call the chemicals that are produced by the Endocrine System?
  - Hormones
- · What is a neuron?
  - A nerve cell

Drug Evaluation & Classification Training

RS-3

## **Review of Physiology**

- · What do we call the space between two nerve cells?
  - The synapse, or synaptic gap
- What do we call the chemicals that pass from one nerve cell to the next?
  - Neurotransmitters
- What do we call the part of the nerve cell that sends out the neurotransmitter?
  - The axon

Drug Evaluation & Classification Training

RS-35

#### Review of Physiology

- What do we call the part of a nerve cell that receives the neurotransmitter?
  - The dendrite
- · What do the Sensory Nerves do?
  - Carry messages to the brain, from the sense organs, pain sensors, etc.
- What do the Motor Nerves do?
  - Carry messages from the brain, to the muscles, etc.

Drug Evaluation & Classification Training

#### Review of Physiology

- · Name the two sub-divisions of Motor Nerves.
  - Voluntary (control striated muscles) and Autonomic (control smooth muscles)
- Name the two sub-divisions of Autonomic Nerves and describe their functions.
  - Sympathetic (command the body's response to fear, excitement, etc.), and Parasympathetic (promote the body's tranquil activities)

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RS-37

## **Review of Physiology**

- What does it mean to say that a drug is "sympathomimetic"?
  - It means that the drug's effects mimic those caused by messages transmitted along sympathetic nerves (excitement, agitation, arousal, etc.)
- What does it mean to say that a drug is "parasympathomimetic"?
  - The drug's effects mimic those caused by messages transmitted along parasympathetic nerves (relaxation, calm, sleep, etc.)

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RS-38

#### Review of Physiology

- Which two categories of drugs can most appropriately be called sympathomimetic?
  - CNS Stimulants and Hallucinogens
- Which category can most appropriately be called parasympathomimetic?
  - Narcotic Analgesics
  - Clarification: Cannabis, PCP, and Inhalants have some sympathomimetic characteristics, but not as many as do the Stimulants and Hallucinogens.
     Depressants have some parasympathomimetic characteristics, but not as many as do the Narcotic Analgesics.

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RS-39

#### **Review of Physiology**

- · What is an artery?
  - Strong, elastic blood vessel that carries blood from the heart to the body's tissues and organs
- · What is a vein?
  - Blood vessel that carries blood back to the heart from tissues and organs

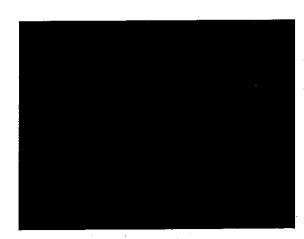
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RS-4

## **Review of Physiology**

- · What is the Pulmonary Artery, and what is unique about it?
  - It is the artery that carries blood from the heart to the lungs. It is the only artery that carries blood depleted of oxygen
- What are the Pulmonary Veins, and what is so special about them?
  - They are the veins that carry blood back to the heart from the lungs. They are the only veins that carry blood rich in oxygen.

Drug Evaluation & Classification Training



Four Hours

# SESSION XXIX CLASSIFYING A SUSPECT (ROLE PLAY)

## SESSION XXIX CLASSIFYING A SUSPECT (ROLE PLAY)

Upon successfully completing this session, the participants will be able to:

o Compile a complete, clear and accurate report documenting the conduct and results of a Drug Evaluation and Classification Examination.

#### Learning Activities Content Segments Scenarios: Simulated Examinations **Interviewing Practice** A. Note taking Practice В. Report Preparation Practice 0 Small Group Work session C. Report Review and Critique 0 Instructor led Presentations 0 Participant led Presentations 0 Participant led Critiques

Aides	Lesson Plan	Instructor Notes
· ·	CLASSIFYING A SUSPECT (ROLE PLAY)	Total Lesson Time: Approximately 240 Minutes Session title on wallchart.
XXIX-O (Objectives)		Briefly review the objectives, content and activities of this session.
	A. Scenarios: Simulated Examinations	
120 Minutes (Approxi- mately)		
	1. Team assignments	Assign the students to teams of 3-4 members.  Note: the total number of student teams should not be more than the number of "role players" participating in this session. Otherwise, one or more teams would be unoccupied during major portions of this segment.
	2. Procedures	Explain procedures to the students.
	a. Each team will examine as many as possible of the "role players", until the time scheduled for this segment elapses.	Solicit students' questions concerning the procedures.
	b. Each examination will be carried out <u>fully</u> : nothing will be omitted <u>except</u> for the breath alcohol test.	

#### Lesson Plan

#### **Instructor Notes**

- c. At certain points in the examination, the "role player" will inform the team what to record.
- d. All data will be recorded on the standard Drug Influence Evaluation Form.
- e. Some "role players" will be simulating the signs and symptoms of exactly one category of drugs.
- f. Some "role players" may be simulating the signs and symptoms of two or more categories in combination.
- g. It is possible that one or more "role players" may be simulating persons who are not under the influence of any drugs.
- h. At the completion of each examination, the team will discuss the evidence obtained and reach a consensus concerning the category or categories of drugs present.
- Subsequently, each team will be assigned the responsibility of preparing and presenting a complete narrative report on one "role player".

Example: The "role players" will instruct the teams concerning the evidence to be recorded from the Horizontal Gaze Nystagmus test.

Clarification: "Role player Alpha" might be simulating a person who is under the influence of a CNS Stimulant only. "Role Player Delta" might be simulating a person under the influence of an Inhalant only.

"Role Player Bravo" might be simulating someone who is under the influence of both PCP and Marijuana.

Aides	Lesson Plan	Instructor Notes
	j. All students will participate in critiquing the reports.	Verify that all teams understand the procedures.
		Make sure that teams have sufficient copies of the Drug Evaluation Form.
	3. Drug Evaluation and Classification practice.	Assign a "role player" to each team.
		Example:  "Alpha" to team #1  "Bravo" to team #2  "Charlie" to team #3, etc.
		As each team completes the entire evaluation, the team will hand over its "role player" to the next team. That is, team #1 hand off to team #2, team #2 to team #3, etc.
		Make sure that each team member fully participates, and conducts some portion of the evaluation of each "role player".
		Allow the practice to continue for approximately 2 hours, or until each team has completed the evaluation of at least three "role players" (whichever occurs <u>later</u> ).
	B. Report Preparation Practice	
60 Minutes		
	1. Team assignments	<u>Instruct</u> each team to prepare a report based on the <u>third</u> "role player" evaluated by the team.
		Verify that each team understands who is to be the subject of the report.

Aides	Lesson Plan	Instructor Notes
	2. Group writing exercise	Note: team members may divide the report writing work among themselves in any way they see fit.
	C. Report Review and Critique	
0 Minutes		
	1. Report presentation	Each team should appoint a speaker to read its report. The speaker should explain exactly what led the team to its conclusion concerning the category or categories of drugs.
	2. Report critique	Solicit questions and commen from students concerning the report they have heard.
		Inquire whether other teams that evaluated this same "role player" reached a different conclusion about the drug category or categories.
		In turn, present and critique the other teams' reports.
		Note: If necessary, this segment can be conducted simultaneously in two separate classrooms, with half of the teams present in each classroom, to allow all reports to be presented and critiqued within the allotted time.

## ROLE PLAY SCENARIOS

## SUBJECT DRUG CATEGORY

Alpha Drug-Free
Bravo Cannabis
Charlie PCP

Delta Narcotic Analgesic

Echo Narcotic Analgesic and CNS Depressant

Foxtrot Cannabis

Golf CNS Stimulant Hotel PCP and Cannabis

India Inhalant

Juliet Alcohol Only (BAC = 0.07)

Kilo Narcotic Analgesic <u>and Alcohol (BAC = 0.05)</u>
Lima CNS Stimulant <u>and Alcohol (BAC = 0.03)</u>

## **Session XXIX**

Classifying a Suspect (Role Play)



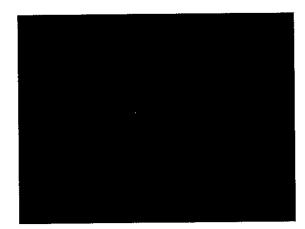
## Classifying a Suspect (Role Play)

Upon successfully completing this session, the participant will be able to:

 Compile a complete, clear and accurate report documenting the conduct and results of a drug evaluation and classification examination

Drug Evaluation & Classification Tenining

XXIX-0



000841	Drug Influence Evaluation	on ·		
Evaluator	DRE No X X / X -   Rolling Log No.			
Recorder/Witness	Crash: ☐ None ☐ Fatal ☐ Injury ☐ Prope	erty		
stee's Name (Last, First, MI)	DOB Sex Race Arre	sting Officer (Name, ID No.)		
ALPHA				
Date Examined/Time/Location	Breath Results: Instrument # 0.00%			
Miranda Warning Given: UYes UNo What By:	t have you eaten today? When? Have you! VOTHING YET TODAY "Ju	been drinking? How much? Time of last drink?  5 T C O FFEE " N / A		
Time now? When did you last sleep? How long? "I HAVEN'T SLEFT IN 2 DAYS"	Are you sick or injured? TYES ON "NO - JUST VERY TIRED	1 *		
Do you take insulin? ☐ Yes No	Do you have any physical defects? □ Yes 💋 N	o Are you under the care of a doctor or dentist?   Yes No		
Are you taking any medication or drugs?   Yes   No	O Attitude PASSIVE, COOPERAT	IVE Coordination SMEWHAT SLOPPY		
	Breath NORMAL	Face FLUSHED		
Speech NORMAL	Eyes:   Reddened Conjunctiva  Normal Bloodshot Watery	Blindness: Tracking:  None OL Eye OR Eye Equal OUnequal		
Collection = reserve	l Size:  #Equal nequal (explain)	Able to follow stimulus:     Yes □ No  Eyelids: □ Normal  Droopy		
Pulse & Time HGN	Left Eye Right Eye Vertical Nystagmus?	□ Yes No One Leg Stand		
Lack of Smooth Pursuit	No No Convergence			
2. 76 / Max. Deviation	No No Right Eye	Left Eye		
3. 76 / Angle of Onset	NONE NONE			
Romberg Balance Walk and Turn T		p Balance		
Арргох. Арргох.	Starts Too 8	I" Nine 2nd Nine Sways While Balancing		
	Stops Walki Misses Heel	ing /		
	Steps Off Li	ne		
	Raises Arms	1* 1* 1		
	Actual Steps	s Taken 9 9 D Puts Foot Down		
Internal Clock  27 Estimated At 30 Sec.  Describe Turn  PROPER	SUT SLOW Cannot Do Test (Ex	F		
○ Right △ Left		Darkness Direct Nasal Area CLEAR		
Draw lines from spot touched	Left Eye 4.5	6.5 3.0 Oral Cavity		
Diaw mes nonspot touched	Right Eye 4.5	6.5 3.0 CLEAR		
4010	Hippus	1 Dilation Reaction To Light  NO NEAR NORMAL		
	AKCHT ARM	LEFT ARM		
2				
	<del>\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac{\frac}}}}}}{\frac{\frac{\frac{\frac{\fin}}{\fint}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}</del>	No Visi		
(5) (6) (7) (AKKS OLE				
Blood Procesure 128 / 84 Terr	90.7			
Muscle Tone: Normal □ Flaccid □ Rigid	np 98,7 . Attach	Photos Of Fresh Puncture Marks		
at Medicine or Drug Have You Been Using?	How Much? Time of Use? Where \	Were The Drugs Used? (Location)		
1. JOTHING - I JUST NEED SOME	SLEEP" N/A	N/A		
Date Time of The Control		art Time . Time Completed		
Member Signature (Include Rank)	ID No. Reviewe	·		
Opinion of Evaluator: □ Rule Out □ Medical	□ Alcohol □ Stimulant □ PCP □ Depressant □ Hallucinogen □ Narco	□ Inhalant tic Analgesic □ Cannabis		

00072		
	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>
LOG NO.	DRE:	ARRESTEE: Alpha
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / DNS 6. MEDICAL PROBLEMS 7. PSYCHOI I 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS
1. LOCATION:		
2. WITNESS:		
3. BREATH TEST:		
4. NOTIFICATION /	INTERVIEW of ARRESTING OFFICER	:
5. INITIAL OBSERVA	ATIONS: Subject appeared to be very tired.	·
6. MEDICAL PROBL	EMS:	
7. PSYCHOPHYSICA	L TESTS:	
8. CLINICAL INDICA	ATORS:	
9. SIGNS of INGEST	ION:	
10. STATEMENTS:		
11. OPINION of EVA	LUATOR:	
12. TOXICOLOGICA	L SAMPLE:	
13. MISCELLANEOU	JS:	/
		/

000843 Drug Influence Evaluation						
Evaluator Evaluator	DRE No XXIX		ling Log No.			
Recorder/Witness	Crash: □ Fatal	□ None □ Injury	O Prope	erty		
stee's Name (Last, First, MI)	DOB	Sex Ra	ce Arre	sting Officer (Nam	e, ID No.)	
Date Examined/Time/Location		Breath Instrum	Results: 0.00	Refused 70 4	Chemical Test  • Refuse	□Urine □Blood d
	have you eaten toda ANDWICH"			been drinking? ICTHING みゃ	How much?	Time of last drink?
Time now? When did you last sleep? How long? "LAST NIGHT - 8 HRS"	Are you sick or i	injured?	□ Yes 💅 N	lo	Are you diabetic or □ Yes 📝 No	epileptic?
Do you take insulin? ☐ Yes   No	Do you have any	y physical defects?	□Yes Z/N	lo .	Are you under the a doctor or dentist	? □Yes No
Are you taking any medication or drugs?   Yes   No	Attitude CAREFRE Breath	E, COOPE	RATIVE	-	Coordination Face	ORMAL
Speech	Eves: □ Rede	dened Conjunctiva		Blindness:	<u> </u>	Tracking:
Normal Corrective Lens: □ None Pupil	□ Normal ØB V Size: ØEqual	loodshot □ Wate ERY BLGOOS	ту нот	Able to follow sting	□ R Eye mulus: 🗗 Yes □ No	ÆEqual □ Unequal     Eyelids:
□ Glasses □ Contacts, if so □ Hard □ Soft □ Un	equal (explain)				——————————————————————————————————————	<b>≠</b> Normal □ Droopy
Pulse & Time HGN	<del></del>	ht Eye Vertica	Nystagmus?	□Yes ØNo	One Leg Sta	ues observed
1. 120 Lack of Smooth Pursuit	<del>                                     </del>	Conver	gence ght Eye	Left Eye	(A)	് ര
2. 112 / Max. Deviation		40	ر 22)		୍ । ବ୍ୟ	(1)
3 Angle of Onset		ONE	Cannot Kee			COUNTED VERY
Romberg Balance Approx. Approx.  Approx. Approx.  Approx. Approx.  Approx. Approx.  Approx. Approx.  Approx. Approx.  Approx. Approx.  Approx. Approx.  Starts Too Soon  I'' Nine 2 rd Nine  Stops Walking  Misses Heel-Toe  Steps Off Line  Raises Arms  Actual Steps Taken  Puts Foot Down						
Internal Clock . Describe Turn  / O Estimated At 30 Sec. PRO	PER	Cam	not Do Test (Ex	xplain)	Type of Foo	twear
○ Right △ Left					rect Nasal Area	LEAR
Draw lines from spot touched	Van	Left Eye Right Eye	<i>5</i> ,0	7.0 4.	— Çıaı Çavışı	JARRATIVE)
4 3110	K) [	Hippus □ Yes Ø No	Rebound Yes		tion To Light VEAR No	RMAL
2		AIGHT A	AM		LEFT ARM	
4	3	=			W. T.	
ANO VISIBLE						
Blood Pressure 168 / 100 Temp 98.6 .						
Muscle Tone: ⊿ Normal □ Flaccid □ Rigid Comments:					sh Puncture Mar	ks
at Medicine or Drug Have You Been Using? No THING - No HARD STUFF AT	How Much? ALL"	Time of Use? "DIDN'T US	ε" " <i>N</i>		DOAY OFF	
Date/Time of Arrest Tim	e DRE Notified			art Time	Time Compl	eted
Member Signature (Include Rank)		ID No.	Reviewe	ed By:		
Opinion of Evaluator:   Rule Out  Medical	□ Alcohol □ Depressant	□ Stimulant □ Hallucinogen	□ PCP □ Narco	tic Analgesic	□ Inhalant □ Cannabis	

	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>			
LOG NO.	DRE:	ARRESTEE: Bravo			
1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.					
1. LOCATION:					
2. WITNESS:					
3. BREATH TEST:					
4. NOTIFICATION / I	INTERVIEW of ARRESTING OFFICER	:			
5. INITIAL OBSERVA	ATIONS: Check of subject's mouth reveale	d small bits of debris (dark green/brown vegetable			
matter) between lower fr	ront teeth.				
6. MEDICAL PROBL	EMS:	•			
7. PSYCHOPHYSICA	L TESTS:				
8. CLINICAL INDICA	ATORS:				
9. SIGNS of INGESTI	ION:				
10. STATEMENTS:					
11. OPINION of EVAL	LUATOR:				
12. TOXICOLOGICA	L SAMPLE:				
13. MISCELLANEOU	US:				

000845	Drug Influence l	Evaluation		
Evaluator	DRE No XXIX·3	Rolling Log No.		
Recorder/Witness	Crash: ☐ None ☐ Fatal ☐ Injury	Property		
stee's Name (Last, First, MI)		Race Arresting Officer	Name, ID No.)	
Date Examined/Time/Location		th Results: 0.00% Refused ument # 0.00%	Chemical Test □ Refuse	□ Urine □ Blood d
Miranda Warning Given: □ Yes □ No What By: □ Ke No □ What	have you eaten today? When? Today?(PAuse) N			Time of last drink?
Time now? When did you last sleep? How long? "THIS MCRNING - 4 HLS"	Are you sick or injured? "I'M VERY Ho		Are you diabetic or Yes ØNo	r epileptic?
Do you take insulin?   Yes No	Do you have any physical defec	ts? Yes No	Are you under the a doctor or dentist	
Are you taking any medication or drugs? □ Yes 📝 No	Attitude DETACHED SLOW Breath NORMA		Coordination Face	
Speech SLOW AND DELIBERATE	Eyes:   Reddened Conjunct  Normal   Bloodshot   Wa	iva Blindness:	Eye CREye	Tracking:  Equal © Unequal
	Size: Æ Equal equal (explain)	Able to folio	w stimulus: AYes □ No	Eyelids:
Pulse & Time HGN  1. 104 Lack of Smooth Pursuit	YES YES Conv	cal Nystagmus? 🗗 Yes 🗆 No ergence Right Eye Left Eye	One Leg Sta	and (18)
2. 108 / Max. Deviation	YES YES	(0)	0	
3. 108 / Angle of Onset  Romberg Balance Walk and Turn T	IMMCDIATE IMMEDIATE	Cannot Keep Balance	HAO TO G	THEN SIMPLY O "ONE THOWSARD"
TEST STOPPED AFTER 90 SECONDS  Internal Clock  Describe Turn	FOLD TO START FTER TURN  S	Misses Heel-Toe Steps Off Line Raises Arms Actual Steps Taken annot Do Test (Explain)	W	Uses Arms to Balance Hopping Puts Foot Down Otwear
1 90	LEAVE FRONT N LINE Pupil Size	Room Light   Darkness	Direct Nasal Area	CLEAR
○ Rìght △ Left	Left Eye	4.0 6.5	3.5 Oral Cavity	CLEAR
Draw lines from spot touched	Right Eye	4.0 6.5	3.5	LEAR
436	Hippus O		Reaction To Light NEAR NEAR	1AL
2	Argu Argu	AT ARM	LEFT ARM	
4	3	No.	V _{ISIBLE}	
3	6	MA	RKS	
Blood Pressure 170 98 Tem	p /00.6 .			
Muscle Tone: □ Normal □ Flaccid Rigid Comments: A KMS V € KY LIGID  at Medicine or Drug Have You Been Using?	How Much? Time of Use?	Attach Photos Of	Fresh Puncture Man	rks
JSING (PAUSE) NOTHING	G" N/A	N/A		
Batto Hind of Taraba	ne DRE Notified	Eval Start Time  Reviewed By:	Time Compl	leted
Member Signature (Include Rank)			□ Inhalant	
Opinion of Evaluator: □ Rule Out □ Medical	☐ Alcohol ☐ Stimulant ☐ Depressant ☐ Hallucinon			

	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>				
LOG NO.	DRE:	ARRESTEE: Charlie				
1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.						
1. LOCATION:						
2. WITNESS:						
3. BREATH TEST:						
4. NOTIFICATION / I	NTERVIEW of ARRESTING OFFICER	:				
5. INITIAL OBSERVA	ATIONS: Subject delayed for several secon	ds prior to responding to most questions. Subject				
stated it was very hot sev	veral times during the examination. Subject	perspired heavily during the examination.				
6. MEDICAL PROBL	EMS:					
7. PSYCHOPHYSICA	L TESTS:					
8. CLINICAL INDICA	ATORS:					
9. SIGNS of INGESTI	ON:					
10. STATEMENTS:	10. STATEMENTS:					
11. OPINION of EVAL	11. OPINION of EVALUATOR:					
12. TOXICOLOGICA	L SAMPLE:					
13. MISCELLANEOU	S:					

000847		luence Eval			
Evaluator	DRE No	Rolling I	og No.		
Recorder/Witness	Crash:	□ None □ Injury	□ Property		
DELTA,	□ Fatal		• •	Maria III Na	<u></u>
stee's Name (Last, First, MI)	DOB	Sex Race	Arresting Officer (	Name, ID No.)	
	1	Breath Resu	lts:0.0  Refused	Chemical Test	□ Urine □ Blood
Date Examined/Time/Location		Instrument #		□ Refused	1
Miranda Warning Given: □ Yes □ No Wha	t have you eaten today	y? When? I	lave you been drinking	? How much ?	Time of last drink?
By:	I DIONT	EAT TODAY	NOTHINE - NO	) B002E	NA
Time now? When did you last sleep? How long?  I DONT Rember	Are you sick or in	njured?	les Mo	Are you diabetic or □ Yes ≅⁄No	epileptic?
Do you take insulin?	Do you have any	physical defects?	res p≪No	Are you under the a doctor or dentist	
Are you taking any medication or drugs?   Yes	o Attitude	ACHED, SIE	DY PASSINE	Coordination	oy - RußbelletteD
I'm ccean	Breath		11, 111530=	Face NORM	2/11
		UOQMA — lened Conjunctiva	Blindness:	100RN	Tracking:
Speech LOW, MUMBIED	Normal □ Bl	oodshot © Watery	b⊈Mone □ I	Eye OR Eye	Equal □ Unequal
Corrective Lens: U None Pup	il Size: Cqual nequal (explain)		Able to follow	w stimulus 🞾 Yes □ No	Eyelids:  Normal Oroopy
Pulse & Time HGN	Left Eye Rigl	nt Eye Vertical Nys	tagmus? □ Yes 🖒 No	One Leg Sta	ınd
Lack of Smooth Pursuit	NON				
1. 52 / Lack of Smooth Pursuit	NO N	Right E	ye Left Eye	( )	l U
3. 521 Angle of Onset	NONE NO	INE		🗳	1
Romberg Balance Walk and Turn	Test VERY RUB	BER-LECTED C	annot Keep Balance 🗸		
Approx. Approx.		"" ""		e 2 nd Nine	Sways While Balancing
Change Walliam					
CREWAL,			eps Off Line		JSES AIMS to Dalance
50/49 C	erenete		ises Arms		Hopping
CILLULAR CILLOS	5		tual Steps Taken	<del></del>	Puts Foot Down
1 METAIL 90 SECOND 1	DID WOT LEA	LI& Cannot D	o Test (Explain)	Type of Foo	otwear
90 -	<b>.</b>	Α.	/n		
MONI FOOT	ONLINE, STALL	Pupil Size Room		Direct Nasal Area	
○ Right △ Left HEF	UJAAQ	T - A Tr			lear
Draw lines from spot touched	Repealency		,0 2,5	20 Oral Cavity	
N A S		Right Eye Z		2.0 C	LEAR
I U Ollo	K	Hippus □ Yes \$\times \text{YNo}	Rebound Dilation □ Yes 🔥 No	Reaction To Light  VERY 5/0	
	17 /	RIGHT ARM		LEFT ARM	FIJE SLABS
2	1 717			(***	
					FOUR
		/	179	and the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of the same of th	LED DOIS
(3) USED WEONL- HA	ND ON XE	(_		. <u></u>	OOZ.MI-
LAST TWO TRIALS		<u> </u>			- Provin
Blood Pressure / 08 / 60 Te	mp <u>97.0</u> .	2			Š
Muscle Tone:   Normal Maria Rigid			Attach Photos Of	Fresh Puncture Ma	rks
Comments: ARMS VERY LOOSEF LACID  at Medicine or Drug Have You Been Using?	o				
nat Medicine or Drug Have You Been Using?	How Much?	Time of Use?	Where Were The Dru		-(2.4.1
Pleases Honest I'm clear		NO, HONES!	FleASE) d	Time Comp	1CA/U
D 11/4/ 11/11/ 05/11/11	ime DRE Notified			, inc comp	
Member Signature (Include Rank)		ID No.	Reviewed By:		
Opinion of Evaluator:	□ Alcohol □ Depressant	□ Stimulant □ Hallucinogen	□ PCP □ Narcotic Analgesic	□ Inhalant : □ Cannabis	

	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>					
LOG NO.	DRE:	ARRESTEE: Delta					
I. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.							
1. LOCATION:							
2. WITNESS:	2. WITNESS:						
3. BREATH TEST:							
4. NOTIFICATION / I	INTERVIEW of ARRESTING OFFICER	:					
5. INITIAL OBSERVA	ATIONS: Subject's eyelids drooped constar	ntly. Subject's head repeatedly nodded forward. At					
times, subject appeared t	to be asleep, but always responded to question	ons. Subject rubbed the face and licked the lips					
repeatedly.							
6. MEDICAL PROBL	EMS:						
7. PSYCHOPHYSICA	L TESTS:						
8. CLINICAL INDICA	ATORS:						
9. SIGNS of INGEST	ION:	-					
10. STATEMENTS:	10. STATEMENTS:						
11. OPINION of EVALUATOR:							
12. TOXICOLOGICA	L SAMPLE:						
13. MISCELLANEOU	JS:						

000849		Drug In	fluence			n		—т	
Evaluator		DRE No		Rolling	Log No.				
Recorder/Witness		Crash: □ Fatal	□ None □ Injury		🗆 Prop	erty			
stee's Name (Last, First, MI)		DOB	Sex	Race	Arre	sting Officer (N	ame, ID No.)		
ECHO			I B	reath Resi	ilts:0.00	□ Refused	Chemical Test	t	□ Urine □ Blood
Date Examined/Time/Location			In	strument	# 123	4	o F	Refused	
Miranda Warning Given: □ Yes		ave you eaten tod	ay? Wher	n?		been drinking?	How much	1?	Time of last drink?
By:	"No	THING				T WATER	Are you diab	ti s an i	N/A
Time now? When did you last sle		Are you sick or	·		Yes D		"No Yes	□ No A	loce"
Do you take insulin?"Not ANY	□ Yes □ No	Do you have an	y physical de	efects?	Yes ₽Y	lo lo	Are you und a doctor or d	er the ca lentist?	are of □ Yes P√No
MORE - I USED TO	Voc. E No.	Attitudo	<u></u>		<u>^</u>		Coordination	n 377 61	GERING GALAT NMAINTAINING BA
Are you taking any medication or drugs? The Yes No Attitude 14351VE,			GENERA WILL OB:	414 (	DOPE.	RATIVE	13		V 1
"NOT ANY MORE"		Breath No.2/	HALLICO Idened Conju	BREA	THING	Blindness:	NOR	a.MA	Tracking:
LOW, MUMBLED, SLUT	10-A AT TOUR	Eyes: □ Red , □ Normal □ E	laenea Conju Bloodshot 🏼 🗗	Watery		None □L	Eye □REye		G Equal □ Unequal
Corrective Lens:	լուարու	1126 13quax	<u></u>			Able to follow	stimulus: Tres C	i No	Eyelids:
□ Glasses □ Contacts, if so □ Har	d Soft Une	qual (explain)					<del></del> -		B Rollina
Pulse & Time HGN		Left Eye Ri	ght Eye V	ertical Ny	stagmus?	□ Yes 🗷 No	One I	Leg Stan	ıd
1. 44 / Lack of	of Smooth Pursuit	YES	YES 0	onvergenc		7 6 El		XV.	3)
2 48 / Max.	Deviation	YES	YES	Right H	Sye	Left Eye		_ ~	4
3 48 / Angle	of Onset	400	400				\\	7	EST STE PED
Romberg Balance	Walk and Turn Te	st - Subtei	T NEARLY	FELL	Cannot Ke	ep Balance_/	<u> </u>	_50/	CANAL FOR CALL FOR
Romberg Balance Approx. Approx.  Walk and Turn Test Starts Too Soon  I'' Nine 2nd Nine  Sways While Balancing									
				<b>R</b>	lisses Hee			□ <b>U</b> :	ses Arms to Balance
1 11 Manuales		1 1		_ / s	teps Off L	ine 🗸	<del>/                                    </del>	о н	opping
HOAD SWIMPED FORDAAD DUGAS TEST					aises Arm	L	M	n Pı	uts Foot Down
Luisoldin Land	/	1/4/	)re stop Test	<i></i>	ctual Ste	18:31	Sport		
Internal Clock  7D Estimated At 30 Sec.	Describe Turn			1		xplain) Su/Sie	-·   ·	of Foot	wear
		N/A	Pupil Size		Light	WO STAMPLE Darkness	Direct Nasz	al Area	
HEAD NODDED FORWARD	t ∆ Left							<u>Ci</u>	EAR
SHOTELT NEVER USED TO LEFT HAND. Draw lines to	re From spot touched ,	. 1	Left Eye	7.	<i>D</i>			Cavity	
N =	-	<b>1</b>	Right Eye		,O		20		EAR_
$\square$ $\square$ $\square$		K)	Hippus	□ Yes ⊡-No	Rebour	D#60		BLE-	REACTION
0				AIGHT ARM		2-INSCHS	LEFT ARM	15 SW	PHOTO A
2	$\leq$ /	<b>/X</b>		E/2	- 4	4144)	484	MAY ?	
10						- DA			2 AGO DOTO CLING FAULD
		<del>√</del> •∕•		/			<i>w</i>		
	- The state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the								$\sum$
				$ \in $					
Blood Pressure 104	58 Tem	<u>97.2.</u>							-
Muscle Tone: □ Normal □ Flacci Comments:	1				Attack	Photos Of F	resh Punctur	e Mar	ks
hat Medicine or Drug Have You	LooSE Been Using?	How Much?	Time of U				s Used? (Locatio		
I STOPPED TWO YE	ARS AGOU		I DIA	st Use"			T USED AN		atad
Date/Time of Arrest	Tim	e DRE Notified				tart Time	Time	Comple	neu
Member Signature (Include Rank)	· · · · · · · · · · · · · · · · · · ·		ID No.		<u> </u>	ved By:			
Opinion of Ethinant-	Rule Out Medical	□ Alcohol □ Depressant	□ Stimul □ Hallud		□ PCP □ Nar	cotic Analgesic	□ Inha □ Can		

	DRUG INFLUENCE EVALUATION	Page_2_of_2
LOG NO.	DRE:	ARRESTEE: Echo
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / ONS 6. MEDICAL PROBLEMS 7. PSYCHON 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS
1. LOCATION:		
2. WITNESS:		
3. BREATH TEST:		
4. NOTIFICATION /	INTERVIEW of ARRESTING OFFICER	d:
5. INITIAL OBSERVA	ATIONS: Subject appeared very drowsy. T	he eyelids drooped constantly, and the head nodded
forward frequently.		•
6. MEDICAL PROBL	EMS:	
7. PSYCHOPHYSICA	al tests:	
8. CLINICAL INDICA	ATORS:	
9. SIGNS of INGEST	ION:	
10. STATEMENTS:		
11. OPINION of EVA	LUATOR:	·
12. TOXICOLOGICA	L SAMPLE:	
13. MISCELLANEOU	JS:	(

000851	Drug Influence l		
Evaluator		Rolling Log No.	
Recorder/Witness	Crash: □ None □ Fatal □ Injury	□ Property	XXIX -6
stee's Name (Last, First, MI)	DOB Sex I	Race Arresting Officer (Nar	ne, ID No.)
FOXTROT		th Results:   Refused	Chemical Test □ Urine □ Blood
Date Examined/Time/Location		ument# 0.00	□ Refused
Miranda Warning Given: ☐ Yes ☐ No What	have you eaten today? When?	Have you been drinking?	How much? Time of last drink?
By:	OTHING" N/A	1 "NOTAING"	
Time now? When did you last sleep? How long?  "LAST NIGHT - ZHR	Are you sick or injured?	□ Yes X No	Are you diabetic or epileptic?
Do you take insulin? ☐ Yes 🕱 No	Do you have any physical defec	ts? □Yes ANo	Are you under the care of a doctor or dentist? □ Yes No
Are you taking any medication or drugs?   Yes	Attitude COOPERA	TIVE	Coordination FAIR
	Breath WORMAL		Face NORINAL
Speech NORMAL	Eyes: Reddened Conjunct	iva Blindness: tery None □ L Ey	e □ R Eye Tracking: Equal □ Unequal
	Size; Equal equal (explain)	Able to follow st	imulus: MYes□No Eyelids: Normal □ Droopy
Pulse & Time HGN	Left Eye Right Eye Verti	cal Nystagmus? □ Yes 🙀 No	One Leg Stand TREMORS VISABLE IN
1. // (6 / Lack of Smooth Pursuit		ergence Right Eye Left Eye	THE ELEVATED LEG
2. 124 / Max. Deviation	NO NO		35/30 (1) 3/30
3. 124 / Angle of Onset	NONE NONE		HAD TO BE REMINDED TO
Romberg Balance Walk and Turn T	est	Cannot Keep Balance Starts Too Soon	
Approx. Approx.		Stops Walking	Sways While Balancing
		Misses Heel-Toe	□ □ Uses Arms to Balance
		Steps Off Line	□ □ Hopping
	STOP STOP	Raises Arms	D D Puts Foot Down
EYELID TREMORS HAD TO BE	REMINDED TO COUNT	Actual Steps Taken ALOUD	8
Internal Clock Describe Turn	SWITEL, NO	annot Do Test (Explain)	Type of Footwear
Estimated At 30 Sec. ABRVBT	TEPS	NA	No. 1 Avec
O Right $\triangle$ Left $TRE$	MORS. Pupil Size	Room Light Darkness D	rirect Nasal Area
	PENEL Left Eye	5.0 6.0 4	.O Oral Cavity
1	SEVERAL Right Eye	5.0 6.0 4	O CLEAR
1 (1) (1)	Hippus C Y	1 44.4	action To Light
	PAGE PAGE	MRA TI	LEFT ARM
2			
4	<u>/3</u>		
5 STOPPED JUST SHORT OF	NOSE NOCHING		
NOSE ON FACU TRY	· · · · · · · · · · · · · · · · · · ·		具具
	<u>. ها. ه ۲</u> م	NO VISIBLE	MARKS
Muscle Tone: Normal   Flaccid   Rigid   Comments:		Attach Photos Of Fre	esh Puncture Marks
at Medicine or Drug Have You Been Using?	How Much? Time of Use?	Where Were The Drugs	Jsed? (Location)
"NONE"	N/A N/A	N/A	I fix a familiar d
Date/Time of Arrest Time	ne DRE Notified	Eval Start Time	Time Completed
Member Signature (Include Rank)	ID No.	Reviewed By;	
Opinion of Evaluator:   Rule Out  Medical	☐ Alcohol ☐ Stimulant ☐ Depressant ☐ Hallucino		□ Inhalant □ Cannabis

	DRUG INFLUENCE EVALUATION	Page <u>2_</u> of_2						
LOG NO.	DRE:	ARRESTEE: Foxtrot						
5. INITIAL OBSERVATION	1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.							
1. LOCATION:	1. LOCATION:							
2. WITNESS:								
3. BREATH TEST:								
4. NOTIFICATION / )	INTERVIEW of ARRESTING OFFICER	:						
5. INITIAL OBSERVA	ATIONS: Subject's eyelids exhibited notice	able tremors during Rhomberg Balance test and						
Finger-To-Nose.								
6. MEDICAL PROBL	EMS:							
7. PSYCHOPHYSICA	L TESTS:							
8. CLINICAL INDICA	ATORS:							
9. SIGNS of INGEST	ION:							
10. STATEMENTS:	10. STATEMENTS:							
11. OPINION of EVAL	LUATOR:							
12. TOXICOLOGICA	L SAMPLE:							
13. MISCELLANEOU	IS:							

000853	Drug In	fluence	Evaluati	ion		
Evaluator	DRE No	Rolling Log No.				
Recorder/Witness	Crash: □ Fatal	□ None □ Injury		operty		XX1X-7
stee's Name (Last, First, MI)	DOB	Sex	Race A	rresting Officer (N	ame, ID No.)	
GOLF					<del></del>	
Date Examined/Time/Location		Ins	eath Results: strument # 호궁 삭	□ Refused	Chemical Test  Refuse	□ Urine □ Blood   d
	have you eaten tod	ay? When	? Have yo	ou been drinking?		Time of last drink?
By: "So"	ME COOKIES	, ,0 %E #5	AGO" "	i od i'acd	350SE"	NO
Time now? When did you last sleep? How long?	Are you sick or	injured?	□ Yes □	□ No	Are you diabetic o	r epileptic? YOU DOING THIS? "
YESTER DAY" "5 HOURS"  Do you take insulin? □ Yes □ No	Do you have an	y physical def	ects? •Yes •	∃ No	Are you under the a doctor or dentist	care of
"Un WHY WAS I STOPPED ?"	"No . No	- 100 - NO	3 - 100"		" NO - NO - NO R doctor or deutist	-120
Are you taking any medication or drugs?   Yes   No	Attitude	บอระโ	AND NE	RUDUS	Coordination FAR BUT	TITTERY
)	Breath		,4,00 100	- V - U - J .	Face	
"00-00 WHAT ? - NO"  Speech VERY RAPID,	Eyes: D Red	ldened Conjun	ctiva	Blindness:	NORMAL	Tracking:
	X Normal □ H	Bloodshot □ V	Watery	None □LI		Equal O Unequal
	Size: X Equal equal (explain)			Able to follow	stimulus: XYes □ No	Eyelids: Normal □ Droopy EYES WDE 00EN
Pulse & Time HGN	Left Eye Ri	ght Eye Ver	rtical Nystagmus	s? □Yes XNo	One Leg St.	· · ·
1. 100 / Lack of Smooth Pursuit	NO I	\o_0 C01	nvergence	Left Fre	NUMBERS	S I O
2. 96 / Max. Deviation	NO 1	<i>30</i>	Right Eye	Left Eye	<u>)</u>	)   Y @
3. 96 / Angle of Onset	NONE N	IONE				2 /30
Pombour Polance Walk and Turn T	est			Keep Balance oo Soon 🗸	24/3	ට /} <del>ව</del> ්
1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Approx. Approx. It Nine 2rd Nine 1/ Sways While Balancing					
Stops Walking						
			Misses H	i i	<b>─</b> ───────────────────────────────────	Oses Arms to Datance
			Raises A			Hopping
			Actual St	teps Taken		Puts Foot Down
SLIGHT CIRCULAR SWAY HAD TO BE BE	MINDED TO CE	<u>αλ4 τζυνα</u>	Cannot Do Test		Type of Fo	otwear
Internal Clock  Describe Turn  12 Estimated At 30 Sec.	-				2,50 2210	
PROPER 5	UT VERY RU	Pupil Size	Room Light		Direct Nasal Area	1
SUBJECT KEPT OPENING EYES AND Right A Left					REDIVES	S IN NOSTRILS
ASKING, AM + Draw lines from spot touched	<b>\</b>	Left Eye	7.5	8.5	Oral Cavit	<b>-</b>
DOING THIS	Vinny	Right Eye	7.5		1.0	(1976-
M.OWO,	$\square$			und Dilation R	eaction To Light	
111 11	N/	<u> </u>	No □ Yes	9 MT40	JERY SLOW	<u> </u>
		^	MRA THESE		LEFT ARM	
2		<b>[</b>		1	~	
	$\sqrt{3}$	No		<b>F</b>	A STATE OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PARTY OF THE PAR	
	\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	MARKS				
(3)	$\overline{\times}_{6}$		,			$\sim$
	Blood Pressure 170 / 100 Temp 99.7.					
Muscle Tone: ▼Normal □ Flaccid □ Rigid Comments:			Attac	ch Photos Of F	resh Puncture Ma	rks
1at Medicine or Drug Have You Been Using?	How Much?	Time of Use	l		s Used? (Location)	
WHAT? NONE! AMI BENG AR	RESTED?"	N/A	1	)/A	Time Comp	lated
Date/Time of Arrest Time	ne DRE Notified			Start Time	Time Comp	1016ft
Member Signature (Include Rank)		ID No.		ewed By:		
Opinion of Evaluator:   Rule Out  Medical	□ Alcohol □ Depressant	□ Stimula □ Hallucir		P arcotic Analgesic	□ Inhalant □ Cannabis	

	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>						
LOG NO.	DRE:	ARRESTEE: Golf						
5. INITIAL OBSERVATION	1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS: 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.							
1. LOCATION:								
2. WITNESS:	2. WITNESS:							
3. BREATH TEST:								
4. NOTIFICATION / I	NTERVIEW of ARRESTING OFFICER	•						
5. INITIAL OBSERVA	ATIONS: Subject appeared very nervous an	d jittery. Kept "stumbling" verbally over words.						
Repeatedly asked "am I	being arrested?"							
6. MEDICAL PROBL	EMS:							
7. PSYCHOPHYSICA	L TESTS:							
8. CLINICAL INDICA	ATORS:							
9. SIGNS of INGEST	ON:							
10. STATEMENTS:								
11. OPINION of EVAL	LUATOR:	•						
12. TOXICOLOGICA	L SAMPLE:							
13. MISCELLANEOU	S:	(						

000855			nfluence	e Eval	uatio	n		
Evaluator	,	DRE No		Rolling	Log No.			
Recorder/Witness		Crash: Fatal	□ None □ Injury		🗆 Prope	erty		XX1X-8
stee's Name (Last, First,	MI)	DOB	Sex	Race	Arre	sting Officer (N	lame, ID No.)	**
HOTEL								
Date Examined/Time/Location	n		I	reath Resunstrument	#	□ Refused ⑤ , ○○	Chemical Test  Refuse	□ Urine □ Blood d
Miranda Warning Given:	Yes □ No What	have you eaten t			Have you	been drinking?	How much?	Time of last drink?
Ву:		DON'T RE	EMEMBER	٤.,	" 200-	THING,		11/12
1	last sleep? How long?	Are you sick	or injured? OT REME!		Yes □ N	lo	Are you diabetic o	r epileptic?
Do you take insulin?	☐ Yes X No	Do you have	any physical de			lo	Are you under the	care of
Do Jou tano mount.			NT REMI				a doctor or dentist	? □Yes XNo
Are you taking any medication	n or drugs? □ Yes □ No	Attitude					Coordination POOR STAG	- CED 11 16
		DAZEL Breath	s Score	0.70	KESP	20 12	Face	
"I DON'T REM	EMBER "	NORM	<u>AL</u>	-		DU de co	FLUSI	NED Tracking:
Speech			Reddened Conju KBloodshot			Blindness:	Eye OR Eye	Equal D Unequal
SLOW DELIBER		VERY T	BLOODSHO	or .		Able to follow	stimulus: X Yes □ No	Eyelids:
Corrective Lens: □ No □ Glasses □ Contacts, if so		Size: KEqual equal (explain)				Able to lollow	Stimurus, A 100 0 110	Normal Droopy
Pulse & Time	HGN	Left Eye	Right Eye V	ertical Nys	stagmus?	X(Yes □ No	One Leg St	and
1.104/	Lack of Smooth Pursuit	YES	YES C	onvergence		T - 01 T3		7 18/
2. 128/	Max. Deviation	YES	YES .	Right E	ye	Left Eye	15 _{30_} [6	) (
		<del></del>		<u> </u>	2)		0	(1)
3. \ <u>J.</u> (0) /	Angle of Onset	I STAKBHAII		C	annot Ke	ep Balance	SEVERSE	TREMORS IN LEGS
Romberg Balance Approx. Approx.	Walk and Turn T Sub IECT SIMPO AFTER TURNING -	NSIES LONCHED A LOCK & NOWN	INC STEPS		tarts Too	Soon	L R	
Stone Walking Walking								
4" 4" 4" 4			ļ		isses Hee	<u>v</u>	ML 10 10	Uses Arms to Balance
	END			J St	teps Off L	ine 🗸	STEPS D	Hopping
				€D'R	aises Arm	<u> </u>		
		STOP		A	ctual Step		8 4 4	Puts Foot Down
SWAY CIRCULAR	Describe Turn	c'en ( c si) st		Cannot I	o Test (E	xplain)	Type of Fo	otwear
Internal Clock  (C) Estimated At 30	اددیجومی	STEPS TOU	UMBD THE	N/		•	ļ	
<del>-</del>	RIGHT		Pupil Size	l	Light	Darkness	Direct Nasal Area	a
SUBJECT HAD TO BE REMINDED TO	⊃Right △ Left		Pupii Size	1,00011	Light	Darwicoo	CLEAR	
TOUCH EU	VICER TO NOSE. We lines from spot touched	_	Left Eye	4.	5	7.0	4.0 Oral Cavit	ty Bits OF
\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	EYELID TREMORS	\( \( \lambda_{\text{m}} \)	Right Eye			7.0	GREEN	CHIBEL CREAMERS
N		7	Hippus	□ Yes			Reaction To Light	C Deresta / a
(X -	7(1)	KI		XN₀	□ Yes		NEAR NORM	1AL_
				RIGHT ARM			LEFT ARM	
(2)		<u>/1</u> \	,			,		
	= /	/3				><. <del>2</del> y		
(4)		^	, No	<del>_</del>			San in	
(5)		$\sqrt{6}$	VISIBL MARK			_		
SUBJECT KEPT STE	10 TSOHE BUILDING	F THE NOSE	1					
· · · · · · · · · · · · · · · · · · ·	VERY RIGID ARM	100.4	1	-	/			
Blood Pressure 172  Muscle Tone: □ Normal		пр <u>(ОО, Т</u>				D1 / CCT	5 . 1. m 3.5	lva
Comments'					Attach	Photos Of I	Fresh Puncture Ma	arks
hat Medicine or Drug Ha	HANDS RIGID ve You Been Using?	How Much?	Time of U	se?	Where	Were The Drug	gs Used? (Location)	
" NOTHING"		N/A	N/		N/	A		
Date/Time of Arrest	Ti	me DRE Notified			1 .	tart Time	Time Comp	pleted
	Ponk		ID No.		Review	red By:	L	
Member Signature (Include				1	□ PCP	-	□ Inhalant	
Opinion of Evaluator:	□ Rule Out □ Medical	□ Alcohol □ Depressant	□ Stimu □ Hallu		□ PCP □ Narc	otic Analgesic	© Cannabis	

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	DRUG INFLUENCE EVALUATION	Page <u>2</u> of <u>2</u>				
LOG NO.	DRE:	ARRESTEE: Hotel				
1. LOCATION 2. WITNESS 3. BREATH TEST 4. NOTIFICATION / INTERVIEW ARRESTING OFCR. 5. INITIAL OBSERVATIONS 6. MEDICAL PROBLEMS 7. PSYCHOPHYSICAL 8. CLINICAL INDICATORS 9. SIGNS OF INGESTION 10. SUSPECTS STATEMENTS 11. OPINION 12. TOXICOLOGY SAMPLE 13. MISC.						
1. LOCATION:						
2. WITNESS:						
3. BREATH TEST:	WA-7					
4. NOTIFICATION / I	NTERVIEW of ARRESTING OFFICER	:				
5. INITIAL OBSERVA	ATIONS: Subject appeared very stiff, rigid.	Subject delayed for several seconds before				
responding to most ques	tions. Subject's eyes were extremely red. S	Subject exhibited a blank stare throughout the				
evaluation.						
6. MEDICAL PROBL	EMS:					
7. PSYCHOPHYSICA	L TESTS:					
8. CLINICAL INDICA	ATORS:					
9. SIGNS of INGESTI	ION:					
10. STATEMENTS:						
11. OPINION of EVALUATOR:						
12. TOXICOLOGICA	L SAMPLE:					
13. MISCELLANEOU	IS:					

000857		fluence Eva	luation		<del></del>
Evaluator	DRE No		Log No.		
Recorder/Witness	Crash: □ Fatal	□ None □ Injury	□ Property		YX1X-9
stee's Name (Last, First, MI)	DOB	Sex Race	Arresting Officer (		
Date Examined/Time/Location		Breath Res Instrument		Chemical Test ☐ Refuse	□ Urine □ Blood d
Miranda Warning Given: ☐ Yes ☐ No What By:	have you eaten tod		Have you been drinking"	? How much?	Time of last drink?
Time now? When did you last sleep? How long?	Are you sick or	injured?	Yes □ No	Are you diabetic o □ Yes No	
Do you take insulin?	Do you have an	ny physical defects?	Yes A No	Are you under the a doctor or dentist	
Are you taking any medication or drugs? □ Yes 🛛 N	O Attitude COOPERAT Breath NORWAL		いしていきもり	Coordination STUMBLINE Face FLUSHED	STAGGERING
Speech LOW, MUMBLED SOME TIMES SLULRED	Eyes: □ Rec	Idened Conjunctiva Bloodshot 🗆 Watery	Blindness:	Eye □ R Eye	Tracking: ¡KŒqual □ Unequal
Corrective Lens:   None   Pupi	l Size: X Equal equal (explain)		Able to follow	v stimulus: ¥Yes □ No	Eyelids: ☑ Normal □ Droopy
Pulse & Time HGN	Left Eye Ri	ght Eye Vertical Ny	stagmus? 🗆 Yes 💢 No	One Leg St	and (13)
1. Clc / Lack of Smooth Pursuit 2. Cl2 / Max. Deviation	<del>                                     </del>	Convergence Right		A	)   0
3. 94 / Angle of Onset	<del></del>	30"		254	35/30
Romberg Balance CIRCULAR Walk and Turn T		DEED S	Cannot Keep Balance V Starts Too Soon 1 ² Nin stops Walking V Jisses Heel-Toe	_   ✓   · · · · ·	Sways While Balancing Uses Arms to Balance
STAGGERED TO THE LEFT	فصفت	man s	Steps Off Line Raises Arms Actual Steps Taken	\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	Hopping Puts Foot Down
Internal Clock  45 Retinated At 30 Sec.  Two CE  PERMINSED  Describe Turn  STAGERS	D TWO STE	Cannot	Do Test (Explain)	Type of Fo	otwear
O Right \( \triangle \text{ Left} \)	E EIGH	Pupil Size Room	n Light Darkness	Direct Nasal Area	
Draw lines from spot touched	M	Right Eye 4.	<del></del>	3.5 Oral Cavit	•
るのうの	n	Hippus	Rebound Dilation	3.5 CLEA Reaction To Light NEAR NORM	
	19	RIGHT ARM		LEFT ARM	
	<u>73</u> \	NO VISIBLE /			
5	<u> </u>	MARKS (			
Blood Pressure 148 / 88 Ter Muscle Tone: Normal Flaccid Rigid Comments:	np <u>98.8</u> .		Attach Photos Of	Fresh Puncture Ma	rks
nat Medicine or Drug Have You Been Using?	How Much?	Time of Use?	Where Were The Dru	gs Used? (Location)	
Date/Time of Arrest Ti	me DRE Notified		Eval Start Time	Time Comp	leted
Member Signature (Include Rank)		ID No.	Reviewed By:	er T. 1. 3.	
Opinion of Evaluator: □ Rule Out □ Medical	□ Alcohol □ Depressant	⊡ Stimulant □ Hallucinogen	□ PCP □ Narcotic Analgesic	□ Inhalant □ Cannabis	

	DRUG INFLUENCE EVALUATION	Page_2_ of _2
LOG NO.	DRE:	ARRESTEE: India
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / DNS 6. MEDICAL PROBLEMS 7. PSYCHOI 1 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS
1. LOCATION:		
2. WITNESS:		
3. BREATH TEST:		
4. NOTIFICATION / I	INTERVIEW of ARRESTING OFFICER	:
5. INITIAL OBSERVA	ATIONS: Subject appeared dazed and confi	ised. Near the end of the evaluation, subject
complained of nausea.		
6. MEDICAL PROBL	EMS:	
7. PSYCHOPHYSICA	L TESTS:	
8. CLINICAL INDICA	ATORS:	
9. SIGNS of INGEST	ION:	
10. STATEMENTS:	•	
11. OPINION of EVA	LUATOR:	
12. TOXICOLOGICA	L SAMPLE:	
13. MISCELLANEOU	JS:	

000859		Drug I	nfluence l	<u>Evaluati</u>	on		
Evaluator		DRE No	]	Rolling Log No.			
Recorder/Witness		Crash:	□ None □ Injury	□ Proj			XX1X-10
stee's Name (Last, First	, MI)	DOB	Sex	Race Arr	esting Officer	(Name, ID No.)	<u>-</u>
JULIET By Francisco Company			Brea	th Results:	□ Refused	Chemical Test	□ Urine □ Blood
Date Examined/Time/Location	)U		Insta	numant#	0.07	□ Refu	
Miranda Warning Given:		have you eaten t	today? When?	Have you	ı been drinking	=	Time of last drink?
Ву:	"501		L" "AT BREAK			EER"	A60"
Time now? When did you "LAST WILL	last sleep? How long?	Are you sick		□ Yes 💆		Are you diabetic	0
Do you take insulin?	□ Yes 🔍 No	Do you have	any physical defec	ts? □Yes 💢	No		he care of ist? □ Yes 'KNo
Are you taking any medicati	on or drugs? □ Yes 🛚 🕱 No	Cocheile Breath	TIVE BUT			Coordination SCOPPY U	
Speed		CDOR OF Eves: DI	Reddened Conjunct	iva	Blindness:	NORTHA	Tracking:
Speech Low, MVMBLED		□ Normal	XBloodshot □W: BLOODSHOT			L Eye DR Eye	≱ Equal □ Unequal
Corrective Lens: ON Glasses Contacts, if so		Size: SEqual			Able to follo	ow stimulus: A Yes □ No	Eyelids:  O Normal A Droopy SUGHTLY DROOPY
Pulse & Time	HGN	Left Eye	Right Eye Vert	ical Nystagmus?	□ Yes □ No	One Leg	Stand HAD TO BE CECHILDAED TO CECHIT OUT
1. <u>87/</u>	Lack of Smooth Pursuit	YES .		vergence Right Eye	Left Eye		A LOUD
2. <u>80 /</u>	Max. Deviation	YES	00	•	$\langle \langle \rangle$	ര	UIU®
3. <u>%O</u> /	Angle of Onset	45°	45°			· \	5/30 27/30
Romberg Balance	Walk and Turn Te	est		Cannot Ko Starts Too		l l R	
Approx. Approx.			ace p	▲ Stops Wall			Sways While Balancing
```\\`\`\\\\\\\\\\\\\\\\\\\\\\\\\\\			1	Misses He		\$ W 5	Uses Arms to Balance
				Steps Off I	Line \( \frac{1}{} \)		Hopping
			متست	Raises Arr	m ,		Puts Foot Down
CIRCULAR SWAY				Actual Ste	9	9	
Internal Clock	Describe Turn			annot Do Test (I	Explain)	Type of	Footwear
37>Estimated At 30	Sec. PROPER	BJT SLE		N/A Room Light	Darkness	Direct Nasal A	rea
	○ Right △ Left		Pupil Size	voom ragut	Darkness	Direct Nasai A	
Dra	w lines from spot touched ,	<b>\</b>	Left Eye	4.5	6.0	3.5 Oral Ca	· · ·
	-	1	Right Eye	4.5	6.0	3.5 CLE	AR_
M 3	3110	KI	Hippus D	Yes Rebou No □ Yes	nd Dilation No	Reaction To Light NEAR NOR	MAL
2			,86	HT ARM		LEFT ARM	
4	、三人	73	, No		> <u>&gt;</u>		
	\ /1	6	VISIBLE				
(5)	" End to the second second second	7 - 1	MARKS			-	
100	/ 84 Tem	<u>98.7</u>		5			~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~
Blood Pressure 128  Muscle Tone: Normal Comments:		p to. C		Attacl	h Photos Of	Fresh Puncture N	Marks
hat Medicine or Drug Ha	wa You Been Heine?	How Much?	Time of Use?	Where	Were The Dr	ugs Used? (Location)	
hat Medicine or Drug Ha	to Tou Deen Come:		NA	2	/A		
Date/Time of Arrest	Tin	ne DRE Notified		Eval S	Start Time	Time Co.	mpleted
Member Signature (Includ	e Rank)		ID No.	Revie	wed By:		
Opinion of Evaluator:	□ Ruie Out	□ Alcohol	l □ Stimulan			□Inhalan	
	□ Medical	□ Depressant	□ Hallucino	ogen □ Nar	cotic Analgesi	c □ Cannab	ıs

	DRUG INFLUENCE EVALUATION	Page_2_of_2
LOG NO.	DRE:	ARRESTEE: Juliet
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / DNS 6. MEDICAL PROBLEMS 7. PSYCHOI I 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS
1. LOCATION:		
2. WITNESS:		
3. BREATH TEST:		
4. NOTIFICATION /	INTERVIEW of ARRESTING OFFICER	:
5. INITIAL OBSERVA	ATIONS: Subject appeared drowsy and gav	e off a moderate odor of alcoholic beverage.
6. MEDICAL PROBL	EMS:	
7. PSYCHOPHYSICA	L TESTS:	
8. CLINICAL INDICA	ATORS:	
9. SIGNS of INGÉST	ION:	
10. STATEMENTS:		
11. OPINION of EVA	LUATOR:	
12. TOXICOLOGICA	L SAMPLE:	
13. MISCELLANEOU	JS:	

000861		<u>iluence</u>	Evaluation	on		
Evaluator	DRE No		Rolling Log No.			
Recorder/Witness	Crash: □ Fatal	□ None □ Injury	□ Prop	perty		XX1X-11
stee's Name (Last, First, MI)	DOB	Sex	Race Arr	esting Officer (Na	me, ID No.)	
, KILO			<u> </u>			
Date Examined/Time/Location		Ins	eath Results: strument # 234	©.05	Chemical Test □ Refuse	□ Urine □ Blood d
	have you eaten tod コアトルじつ"		? Have you	been drinking? と BととP		Time of last drink?
Time now? When did you last sleep? How long?	Are you sick or	injured?	□Yes X	No	Are you diabetic or □ Yes XNo	epileptic?
Do you take insulin?	Do you have an	y physical def	ects? □Yes X	No	Are you under the a doctor or dentist	care of ? □ Yes XNo
Are you taking any medication or drugs?   Yes No	Breath		BUT DROI	WSY AGE	Coordination STUMBUNC Face NORMAL HEAD NOODS	STAGGERING COLOR NG, DEDWISY
Speech State, Low, FRASPY	Eyes: Red	ldened Conjun	ctiva	Blindness: None □ L E		Tracking: Equal © Unequal
Corrective Lens: D None Pupil	Size: Equal qual (explain)			Able to follow s	timulus: A Yes □ No	Eyelids: Normal ADroopy  VERY DROOPY
Pulse & Time HGN	Left Eye Ri	ght Eye Ve	rtical Nystagmus?	□Yes XNo	One Leg Sta	m in m
1. (3) / Lack of Smooth Pursuit	YES Y	Co.	nvergence Right Eye	Left Eye	STOPPED A	STOPPED
2. 60 / Max. Deviation		<u>)</u>	4-0	( <u>Ç</u>	TEST U	1 1 6 153.
3/ Angle of Onset	45° 4	15°				1
Romberg Balance Walk and Turn Te Approx. Approx.	st र्ं रंं	υP	Cannot Ke Starts Too	eep Balance <u>//</u> Soon	L R	
	STOP		Stops Walk Misses Hee Steps Off I Raises Arm Actual Step	el-Toe J ine J J J	<i>J b d</i> 1	Sways While Balancing Jses Arms to Balance Hopping Puts Foot Down
Internal Clock Describe Turn	-UNSEC SIE	05.40	Cannot Do Test (E	Explain)	Type of Foc	twear
THE RICH	7 MODDED	Pupil Size	N/A Room Light	Darkness D	Direct Nasal Area	
- ( ) (	uaris Frequently	Left Eye	1.5	1.5	.5 Oral Cavity	
NE E	<b>V</b> 2	Right Eye	1.5		.5 CLEAN	2
M Silo	K)		Yes Rebour (No □ Yes	1-/ x t -	action To Light NE OBSERVA	હાદ
2		А	OGHT ARM	SCARD	LEFT ARM SCAT	2
	<del>\</del> 3\	Į	=			SCABS
	6			7.50	Sep. 1	SCARS
(5)						PUDCTURE
Blood Pressure 112 / 66 Tem  Muscle Tone: □ Normal Flaccid □ Rigid  Comments:			Attach	n Photos Of Fr	Pਸਰਾਹ A esh Puncture Mai	DETS AVID
( ) (	How Much?	Time of Use	TIM 1	Were The Drugs	Used? (Location)	1 00
NOTHING IM CLEAR Time Date/Time of Arrest Time	e DRE Notified	CLEAN	<u> </u>	tart Time	Time Compl	eted
Member Signature (Include Rank)		ID No.		ved By:		
Opinion of Evaluator:	□ Alcohol □ Depressant	□ Stimula □ Hallucir		cotic Analgesic	□ Inhalant □ Cannabis	

	DRUG INFLUENCE EVALUATION	Page_2_of_2
LOG NO.	DRE:	ARRESTEE: Kilo
5. INITIAL OBSERVATIO	ESS 3. BREATH TEST 4. NOTIFICATION / DNS 6. MEDICAL PROBLEMS 7. PSYCHOL I 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS
1. LOCATION:		
2. WITNESS:		
3. BREATH TEST:		
4. NOTIFICATION / I	INTERVIEW of ARRESTING OFFICER	:
5. INITIAL OBSERVA	ATIONS: Subject gave off a moderate odor	of an alcoholic beverage. Subject scratched the arms
and face frequently, and	licked the lips repeatedly.	
6. MEDICAL PROBL	EMS:	
7. PSYCHOPHYSICA	L TESTS:	
8. CLINICAL INDICA	ATORS:	·
9. SIGNS of INGEST	ION:	
10. STATEMENTS:		
11. OPINION of EVAL	LUATOR:	
12. TOXICOLOGICA	L SAMPLE:	
13. MISCELLANEOU	īs:	

000863		ıfluence ]	Evaluatio	n	1	1
Evaluator	DRE No		Rolling Log No.			
Recorder/Witness	Crash:	□ None □ Injury	D Prop	•		XXIX-15
stee's Name (Last, First, MI)	DOB	Sex	Race Arre	sting Officer	Name, ID No.)	
1 LIMA				- D . 1	CU	□ Urine □ Blood
Date Examined/Time/Location			rument#	□ Refused	Chemical Test ☐ Refuse	d
	t have you eaten to ME TOAST" "	day? When?	Have you	been drinking はらったし		Time of last drink?
Time now? When did you last sleep? How long?	Are you sick o	r injured?	□ Yes □ N	Vo	Are you diabetic o	
"YESTEROAY" 5 HOULS"  Do you take insulin?	Do you have a	ny physical defec	ts? □Yes XI	yo .	Are you under the a doctor or dentist	? □Yes □No
Are you taking any medication or drugs?   Yes	No Attitude	5 ANXIO	<u>.</u>		"OF COURSE Coordination JUTERY (	
"OF COVESE NOT"	Breath	AN ALCO		VEDA6.8	Face NUKMAL	
Speech No.	Eves: DRe	eddened Conjunct Bloodshot   We	iva	Blindness:	LEye □REye	Tracking: ★Equal □ Unequal
Corrective Lens:	il Size: Æqual nequal (explain)			Able to follo	w stimulus: ▼Yes □ No	Eyelids:  XNormal □ Droopy
Pulse & Time HGN	Left Eye F	light Eye Vert	ical Nystagmus?	□ Yes □ No	3	
1. 100 / Lack of Smooth Pursui	t YES		vergence	× 4.11	(3)	4)
2. 104 / Max. Deviation	130	200	Right Eye	Left Eye		)   (4
3. 104-/ Angle of Onset	NONE !	none		ep Balance	457	444
VERY SUCHT CHACKED SWAY  Internal Clock  Describe Turn	OPPED COUNT	DE OUT LOAD DE COURT LANDED	Stops Walk Misses Hee Steps Off L Raises Arm Actual Step annot Do Test (E	ing l-Toe ine ss V/		Sways While Balancing Uses Arms to Balance Hopping Puts Foot Down いなみ ないにメート otwear
O Right \( \triangle \text{Left} \)	NEL-NO SMAL	Pupil Size	Room Light	Darkness	Direct Nasal Area	REDIVERS IN NOSTRIUS
Draw lines from spot touched	<b>N</b> 1	Left Eye	7.5	8.5	7.0 Oral Cavit	
1/		Right Eye	7.5	8.5	7.0 CEAR	
U.Ollo	K	Hippus 🗅		nd Dilation	Reaction To Light VERY SLOUD	
2	PA	***	URA THE		LEFT ARM	
4)	3	<b>₽</b> 0				
5 AFTER 3M TRIAL SUBJECT	OPENED SO	MYBRE				
EYES, AND ASKED, AM I WOOD	ARRESTI	_	<b>E</b> ,_			
Blood Pressure 170 / 100 To  Muscle Tone: Normal   Flaccid   Rigid  Comments:	mp 99.6.	_	Attach	Photos Of	Fresh Puncture Ma	rks
nat Medicine or Drug Have You Been Using?	How Much?	Time of Use?	Where		ags Used? (Location)	
Date/Time of Arrest	ST?	N/A	1 '	tart Time	Time Comp	leted
Member Signature (Include Rank)	ing Die Hounes	ID No.		ved By:		
Opinion of Evaluator: ☐ Rule Out ☐ Medical	□ Alcohol □ Depressant	□ Stimulan □ Hallucino		otic Analgesic	□ Inhalant □ Cannabis	-

	DRUG INFLUENCE EVALUATION	Page_2_ of _2_
LOG NO.	DRE:	ARRESTEE: Lima
5. INITIAL OBSERVATION	ESS 3. BREATH TEST 4. NOTIFICATION / ONS 6. MEDICAL PROBLEMS 7. PSYCHO N 10. SUSPECTS STATEMENTS 11. OPINIC	PHYSICAL 8. CLINICAL INDICATORS
1. LOCATION:		
2. WITNESS:		
3. BREATH TEST:		
4. NOTIFICATION / 1	INTERVIEW of ARRESTING OFFICER	R:
5. INITIAL OBSERVA	ATIONS: Subject gave off a slight odor of	alcoholic beverage. Subject kept asking, "am I being
arrested?" Subject ofter	n repeated statements and frequently "trippe	d" verbally over words.
6. MEDICAL PROBL	EMS:	
7. PSYCHOPHYSICA	AL TESTS:	
8. CLINICAL INDICA	ATORS:	
9. SIGNS of INGEST	ION:	
10. STATEMENTS:		
11. OPINION of EVA	LUATOR:	
12. TOXICOLOGICA	L SAMPLE:	
13. MISCELLANEOU	JS:	

### GUIDELINES FOR ROLE PLAYERS

As a "role player", you have the important task of helping students practice the administration and interpretation of drug evaluation and classification examinations. You will also be expected to coach the students as they are practicing. To help insure that you do the best possible job, please follow these guidelines carefully.

- 1. Study the exemplar for your assigned role play carefully and thoroughly.

  Become familiar with all of the information it contains. You do not have to memorize the exemplar. Instead, you will carry the exemplar with you, and you will refer to it as the students administer their tests to you. But you must be familiar with the exemplar to make sure that you give the students all of the information they need to classify "your" drug category or categories.
- 2. <u>Do not</u> attempt to "act" impaired. Let the information on the exemplar speak for itself.
- 3. Start by informing the students of your role play "name" (Alpha, Bravo, etc.). State your actual age. Instruct students to record your actual sex and race, and the actual date and time.
- 4. <u>Inform</u> the students of the BAC for your role.
- 5. For the Preliminary Examination:
  - a. Answer each question exactly as indicated on your exemplar.
  - b. Instruct students to record your answers exactly as you give them.
  - c. Allow students to conduct the preliminary examinations of your <u>eyes</u>.

    <u>Coach</u> them as necessary during the preliminary eye checks to make sure they conduct the checks properly. When they have finished, tell them to record the information given on your exemplar.
  - d. Allow students to conduct the first check of your <u>pulse</u>. Coach them as necessary during to make sure that they check pulse properly. When they have finished, tell them to record the information given on your exemplar.
- 6. For the Eye Examinations:
  - a. Allow the students to conduct the full tests of Horizontal Gaze
    Nystagmus, Vertical Gaze Nystagmus and Lack of Convergence.

    Coach them as necessary to make sure they conduct the tests properly.

b. As they complete each test, instruct them to record the information given on your exemplar.

## 7. For the Psychophysical Tests:

- a. <u>Do not</u> actually perform the Romberg test. Instead, allow the students to give you the Balance test instructions, then comment on their performance in giving the instructions. Tell them to record the Romberg test information given on your exemplar.
- b. <u>Do not</u> actually perform the Walk and Turn test. Instead, place your feet in the heel-to-toe stance for the "instructions stage" and allow the students to give you the Walk and Turn instructions. When the instructions are completed, comment on the students' performance in giving the instructions. Then, tell them to record the Walk and Turn information given on your exemplar.
- c. <u>Do not</u> actually perform the One Leg Stand test. Instead, allow the students to give you the One Leg Stand instructions (for <u>one</u> leg), then comment on their performance in giving the instructions. Tell them to record the One Leg Stand information given on your exemplar.
- d. You <u>will</u> have to perform the Finger-to-Nose test, since students give instructions throughout that test. <u>Try</u> to place your finger tips on the points indicated in the diagram on your exemplar. When the test is completed, <u>show</u> the diagram to the students and instruct them to replicate it on their record form.

# 8. For the Vital Signs Examinations:

- a. Allow the students to conduct the full checks of blood pressure, temperature and pulse. <u>Coach</u> the students as necessary to make sure they conduct the tests properly.
- b. As they complete each test, instruct them to record the information given on your exemplar.

### 9. For the Dark Room Examinations:

- a. Allow the students to conduct the full checks of pupil size, pupil reaction to light, nasal area and oral cavity. <u>Coach</u> them as necessary to make sure they conduct the checks properly.
- b. As they complete each check, instruct them to record the information given on your exemplar.

- 10. Examinations for Muscle Tone and Injection Sites:
  - a. Allow the students to conduct these examinations, and coach them as appropriate. Allow students to conduct the third check of your <u>pulse</u>.

    <u>Coach</u> them as necessary to make sure that they check pulse properly. When they have finished, tell them to record the pulse measurement shown on your exemplar.
  - b. Instruct them to record the information given on your exemplar.
- 11. Give the students the information (if any) contained on the reverse side of your exemplar. Do not make any other statements.
- 12. When you finish working with one team of students, move on to the next team.

Two Hours and Thirty Minutes

# SESSION XXX

TRANSITION TO THE CERTIFICATION PHASE OF TRAINING

# SESSION XXX TRANSITION TO THE CERTIFICATION PHASE OF TRAINING

During this session, the participant will:

- o Demonstrate their mastery of the knowledge the course was intended to help them develop.
- o Summarize the key topics covered.
- o Offer comments and suggestions for improving the course.
- o Receive assignments for Certification Training.

Content Segments		<u>Lear</u>	Learning Activities		
A.	Summary	O	Participant led Presentations		
В.	Post-Test	0	Participants' Anonymous		
C.	Critique	_	Critique of Course		
D.	Certification Training Assignments	0	Knowledge Examination		
	and Schedule	0	Instructor led Presentation		
E.	Closing Remarks				

Aides	Lesson Plan	Instructor Notes
	TRANSITION TO THE CERTIFICATION PHASE OF TRAINING	Total Lesson Time: Approximately 150 Minutes Session title on wallchart.
XXX-O (Objectives)		Briefly review the objectives, content and activities of this session.
15 Minutes	A. Summary .	
13 Minutes	1. The seven categories of drugs.	Ask students to name the seven categories. Make sure all categories are named.
	<ul> <li>a. CNS Depressants</li> <li>b. CNS Stimulants</li> <li>c. Hallucinogens</li> <li>d. PCP</li> <li>e. Narcotic Analgesics</li> <li>f. Inhalants</li> <li>g. Cannabis</li> </ul>	
	2. The drug evaluation and classification procedure.  a. Breath Alcohol Test b. Interview of Arresting     Officer c. Preliminary Examination d. Examinations of Eyes e. Divided Attention Tests f. Vital Signs Examinations g. Dark Room Examinations h. Check for Muscle Rigidity i. Inspection for Injection     Sites j. Statements and	Ask students to name the components of the procedure. Make sure all components are named. Ask students to discuss the kinds of evidence/information gleaned from each component.

Toxicological Examination

Observations k. Opinions of Evaluator

1.

Aides	Lesson Plan	Instructor Notes
	3. Major signs and symptoms.	Instruct students to turn to the symptomatology chart in their manuals.
		Briefly summarize and review the major signs and symptoms associated with each drug category.
		Solicit students' questions concerning the major content topics of the course.
		Inform the students that the final exam in a "closed book" test. Instruct them to put all books and notes away.
	B. Post-Test	
100 Minutes		
	1. Knowledge examination.	Distribute post-test knowledge examinations.
		Allow students approximately 80 minutes to complete the knowledge examination.
		Collect the completed knowledge examination.
		Grade the knowledge exams.
15 Minutes	C. Critique	Distribute the anonymous critique forms.
10 1/1111111005		Collect the completed forms.
10 Minutes	D. Certification Training Assignments and Schedule	
. V MALLEMUCO		

		-		
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_		n	Н.	

### Lesson Plan

### **Instructor Notes**

- 1. Remind students that during certification all evaluations must be supervised by instructors to count towards minimum certification requirements.
- 2. In situations where an instructor is not available to observe a student evaluation, the student should check the local policy governing this. These evaluations do NOT count toward certification requirements. It is important to remember that the ultimate goal is to remove the impaired driver from the highway.
- 3. Instructors should take the time to explain the state's requirements for certification and the final knowledge examination.

Hand out sheets to each student outlining his or her schedule of certification training.

Explain logistics and administrative procedures for certification training.

Note: The minimum national standards for certification are at the back of the instructor manual. (State requirements may be more stringent than the national standards.)

Remind students that they must bring their Certification. Progress Logs to each night of Certification Training. Also remind them to bring their "Rolling Logs" (last five pages of the Student's Manual) to all Certification Training nights.

Solicit students' questions concerning certification training.

Brief closing remarks will be offered by appropriate representatives of the department and faculty.



10 Minutes

E. Closing Remarks





# **Session XXX**

Transition to the Certification Phase of Training





# Transition to the Certification Phase of Training

During this session, the participant will:

- Demonstrate his or her mastery of the knowledge the course was intended to help him or her develop
- · Summarize the key topics covered
- · Offer comments and suggestions for improving the course
- · Receive his or her assignments for Certification Training

Drug Evaluation & Classification Training

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# Congratulations!

Drug Evaluation & Classification Training

# PROFICIENCY EXAMINATION CHECKLIST (For Use During Certification Training)

Dat	te _	Examiner
[.	Pre	eliminary Examination
	1.	Did the student ask all preliminary examination questions?
		yesno
	(If	No: What questions were deleted?
	2.	Did the student properly estimate pupil size?
		yesno
	3.	Did the student properly assess the eyes' tracking ability?
		yesno
	4.	Did the student properly measure pulse rate?
		yesno
I.	Еу	re Examinations
	1.	Did the student properly administer the Horizontal Gaze Nystagmus test?
		yesno
	(If	no, explain deficiencies
•	2.	Did the student properly administer the Vertical Gaze Nystagmus test?
		yesno
	(If	no, explain deficiencies

3.	Did the student properly administer the test for Lack of Convergence?
	yesno
(If	no, explain deficiencies
I.	Psychophysical Tests
1.	Did the student properly administer the Romberg Balance test?
	yesno
1. (If 1) 3. (If 1) 4.	no, explain deficiencies
***	
2	Did the student properly administer the Walk and Turn test?
Д.	
(If y	yesno no, explain deficiencies
\ <u></u>	
3.	Did the student properly administer the One Leg Stand test?
	yesno
(If r	no, explain deficiencies
4.	Did the student properly administer the Finger To Nose test?
	yesno
(If r	o, explain deficiencies

V.	<u>Vital Signs Examinations</u>				
	1.	Did the student properly measure blood pressure?			
		yesno			
	(If	no, explain deficiencies			
	2.	Did the student properly measure temperature?			
		yesno			
	(If	no, explain deficiencies			
	3.	Did the student properly measure pulse?			
		yesno			
	(If	no, explain deficiencies			
IV.		Dark Room Examinations			
	1.	Did the student properly control the pen light for the two checks of pupil size?			
		yesno			
	(If	no, explain deficiencies			
	2.	Did the student accurately estimate pupil size?			
		yesno			
	3.	Did the student properly check the nasal area?			
		yesno			

	4.	Did the student properly check the oral cavity?	
		yesno	
VI.		Examinations of Muscle Tone	
	1.	Did the student adequately inspect for muscle tone?	
		yesno	-
	(If	no, explain deficiencies	
v.	Exa	aminations of Injection Sites and Third Pulse	
	1.	Did the student adequately inspect for injection sites?	
		yesno	
	(If i	no, explain deficiencies	·····
	2.	Did the student properly measure pulse?	
		yesno	
	(If 1	no, explain deficiencies	
VII		Evaluator's Opinion of Student's Proficiency	
	(Off	fer appropriate, specific comments concerning the student's progress)	
			· · · · · · · · · · · · · · · · · · ·
			·····

Course Location	Date

## DRE SCHOOL STUDENT'S CRITIQUE FORM

### 1. Rating The Various Segments Of The School

On a scale from 1 (="low") to 5 (="high"), please indicate how important each major topic or activity of this school was for you personally. Drugs In Society and In Vehicle Operation Development and Effectiveness of the DEC Program Overview of the Drug Recognition Expert Procedures Physician's Desk Reference Eye Examinations: Explanation and Demonstrations by Instructors Eye Examinations: Hands-on Practice by Students Vital Signs: Explanations and Demonstrations by Instructors Vital Signs: Hands-on Practice by Students Physiology and Drugs The Alcohol Workshop The "Practice: Test Interpretation" Sessions The Sessions on the Individual Drug Categories Overview of Signs and Symptoms **Drug Combinations** Resume Preparation and Maintenance Preparing the Narrative Report Case Preparation and Testimony The Mid-Course Review Session The Role Play Session (Instructors "simulating" drug impaired subjects).

The Quizzes

2. If		For Improving The School had to cut four hours out of this	school, what topics or sessions				
	ould you reduce						
	you could add fo ditional time bo	our hours to the School, how wo	uld you recommend that the				
3.	Specific Fea	tures Of The School					
		appropriate word to indicate you ing statements.	r agreement or disagreement with				
1.	The DRE Scho	ool is at least one day too long.					
	Agree	Disagree	Not Sure				
2.	We spent too 1	We spent too much time in hands-on practice.					
	Agree	Disagree	Not Sure				
3. Now that I've had the DRE School, I believe the needed.			nat the <u>PRE</u> -School really wasn't				
	Agree	Disagree	Not Sure				
4. Some of the instructors didn't seem to be as well prepared as they sbeen.			ell prepared as they should have				
	Agree	Disagree	Not Sure				
5.	I do <u>not</u> feel co accurately.	onfident about my ability to estin	nate nystagmus onset angle				
	Agree	Disagree	Not Sure				
3.	This School was much harder than I thought it would be.						
	Agree	Disagree	Not Sure				

We should have spent more time in hands-on practice.			
Agree	Disagree	Not Sure	
3. The instructors seemed to know their material, but some of them didn't ge across very well.			
Agree	Disagree	Not Sure	
We spent too much tim	e on the details of each drug catego	ory.	
Agree	Disagree	Not Sure	
I am <u>not</u> confident that	I can measure blood pressure accu		
Agree	Disagree	Not Sure	
I would have to say tha	at the final examination was hard,	but fair.	
Agree	Disagree	Not Sure	
2. Some of the instructors "threw the bull" a bit too much.			
Agree	Disagree	Not Sure	
3. Now that I've had the DRE School, I am more convinced than ever that the <u>PRE</u> -School is very important.			
Agree	Disagree	Not Sure	
4. I am still very confused about drug combinations and their effects.			
Agree	Disagree	Not Sure	
I am not confident that	I can estimate pupil size accuratel	y.	
Agree	Disagree	Not Sure	
I would have to say that be.	at this School wasn't quite as hard a	as I thought it would	
Agree	Disagree	Not Sure	
	The instructors seemed across very well.  Agree We spent too much time Agree I am not confident that Agree I would have to say that Agree Some of the instructors Agree Now that I've had the Inpre-School is very impressed agree I am still very confused Agree I am not confident that Agree I am not confident that Agree I would have to say that be.	The instructors seemed to know their material, but some across very well.  Agree Disagree  We spent too much time on the details of each drug category. Agree Disagree  I am not confident that I can measure blood pressure accurately. Agree Disagree  I would have to say that the final examination was hard, and agree Disagree  Some of the instructors "threw the bull" a bit too much. Agree Disagree  Now that I've had the DRE School, I am more convinced to the pressure accurately. Agree Disagree  I am still very confused about drug combinations and their Agree Disagree  I am not confident that I can estimate pupil size accurately agree Disagree  I would have to say that this School wasn't quite as hard a be.	

17.	There were to	oo many quizzes in this Scho	ool.	
	Agree	Disagree	Not Sure	
18.	18. The final examination was much harder than it should have been.			
	Agree	Disagree	Not Sure	
19.	9. We did <u>not</u> receive enough information about the effects, signs and symptoms of the various drug categories.			
	Agree	Disagree	Not Sure	
20.	I am confiden	at that I will succeed in the G	Certification Stage of my training.	
	Agree	Disagree	Not Sure	
21.	The DRE Sch	ool is at least one day too sh	ort.	
	Agree	Disagree	Not Sure	
4.	Rating of In	structors		
	a scale from 1 essment of eac		, please indicate your overall	
Ins	tructor		Rating	
Ins	tructor		Rating	
Ins	tructor		Rating	
Īns	tructor		Rating	
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# The International Standards of the Drug Evaluation and Classification Program



### A Product of

The DEC Standards Revision Subcommittee of the Technical Advisory Panel of the IACP Highway Safety Committee

Revised June 2, 1999

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#### EXECUTIVE SUMMARY

Since 1984, the National Highway Traffic Safety Administration (NHTSA) has supported the Drug Evaluation and Classification Program. The program which was initially developed by the Los Angeles, California, Police Department, was validated through both laboratory and field studies conducted by Johns Hopkins University. In 1987, the Highway Safety Committee of the International Association of Chiefs of Police (IACP) was requested by NHTSA to participate in the development and national expansion of the program. As the program grew, it became apparent that in order to ensure continued success, nationally accepted standards needed to be established. These standards, which establish criteria for the selection, training and certification of drug recognition experts, helped to ensure the continued high level of performance of the Drug Evaluation and Classification Program. In 1988, NHTSA asked the IACP and its Highway Safety Committee to develop this system of nationally accepted standards.

In March of 1989, the IACP and NHTSA sponsored a meeting at the Transportation Safety Institute in Oklahoma City, Oklahoma. Persons invited to this meeting included experienced drug recognition experts, instructors, curriculum specialists, toxicologists, prosecutors and training administrators. The participants met in working groups to reach consensus concerning the many issues relating to the Drug Evaluation and Classification Program and to develop recommended minimum standards to the Highway Safety Committee. The standards were drafted and presented to the committee for review at its mid-year meeting in June 1989. In addition, the committee agreed to name a Drug Evaluation and Classification Technical Advisory Panel to assist and advise the committee concerning technical aspects relating to the operation of the program.

The Highway Safety Committee, by resolution, adopted the *Interim National Standards of the Drug Evaluation and Classification Program*. The standards were subsequently approved by the voting membership of the IACP. The standards were adopted on an interim basis pending the outcome of an evaluation of the effectiveness of the program to be performed by NHTSA. In October 1992, the standards were officially approved and adopted. Revisions and updates are periodically made to the standards.

Presented in this document are standards specifying the requirements for certification and recertification of DREs and DRE instructors; standards for decertification and reinstatement; and standards for agency participation. Also, for those agencies participating in the program, a set of administrative guidelines is provided.

These standards, when adopted by other countries, will be administered pursuant to their political structure.

#### DEFINITIONS

Associate Instructor: Persons not certified as DREs but who possess knowledge, expertise or credentials deemed valuable to the program may be designated as associate instructors for the Drug Evaluation and Classification Program.

Blood Alcohol Concentration (BAC): A person's blood alcohol concentration indicates the grams of alcohol per 100 milliliters of blood. For example, a BAC of 0.10% means that there is one-tenth of a gram of alcohol in 100 milliliters of the person's blood.

Candidate DRE: An individual in the process of achieving certification as a drug recognition expert. To achieve certification, a person must successfully complete a training program consisting of

An IACP/NHTSA-approved SFST training course of instruction A two-day IACP/NHTSA-approved DRE preschool A seven-day IACP/NHTSA-approved DRE school On-the-job field certification

Candidate DRE Instructor: An individual in the process of achieving certification as a DRE instructor. To achieve certification, a DRE must successfully complete the IACP/NHTSA-approved DRE instructor training, conduct a minimum of two hours of DRE training, and witness two drug evaluations.

Course Manager: An individual who ensures that each training event follows the standardized curriculum and evaluates the training event to identify ways to improve it. The course manager represents the National Highway Traffic Safety Administration and the International Association of Chiefs of Police and resolves issues with the content and/or delivery of the training.

DRE Coordinator: The appropriate DRE coordinator will be one of the following:

Agency Coordinator: The person designated within each department or agency responsible for maintaining program records, ensuring maintenance of program standards and conducting training and certification sessions within the agency. Responsibility for this function may rest with one individual, in the case of a small or closely coordinated effort, or may be decentralized among several people throughout the agency. If there is no designated agency coordinator, the appropriate DRE coordinator shall be the state coordinator.

State Coordinator: In each of the states in which the Drug Evaluation and Classification Program has been implemented under the auspices of the National Highway Traffic Safety Administration, an individual has been designated to act as the statewide coordinator for the DEC Program. The duties of the position generally include but are not limited to

1. Acting as an information clearinghouse and central communication point for the program within the state.

2. Assisting in coordinating training and other support activities for all agencies participating in the program within the state.

3. Coordinating the assignment of instructors in response to requests for service from federal and other sources.

The Governor's Office of Highway Safety shall be responsible for designating the state coordinator. If there is no designated state coordinator, the appropriate DRE coordinator shall be the TAP regional coordinator, who shall assume the duties and responsibilities as described above.

TAP Regional Coordinator: One DRE from each of the four regions, as established by the Division of State and Provincial Police, is appointed by the IACP Highway Safety Committee Chair to serve on the Technical Advisory Panel.

**DRE Instructor:** Individuals who, having been trained and certified as drug recognition experts, receive further training and experience instructing within the Drug Evaluation and Classification Program. Certified instructors will usually be certified DREs with experience in performing drug evaluations and in providing testimony in court in the area of drug recognition. Certified instructors are responsible for observing, evaluating and verifying the performance of candidate DREs.

Criminal Justice Agency: For purposes of these standards, a criminal justice agency is any organization, funded by public monies, that is involved in the apprehension, prosecution, adjudication of public miscreants; or in the incarceration, detention, supervision or control of said miscreants following apprehension, prosecution and/or adjudication.

**Drug:** For purposes of the Drug Evaluation and Classification Program, a drug is any substance that, when taken into the human body, can impair the ability to operate a vehicle safely. Note that this is not necessarily a strict medical definition.

**Drug Evaluation:** A process of systematically examining a person suspected of being under the influence of a drug, for the purpose of ascertaining what category of drugs (or combination of categories) is causing the person's impairment. A trained DRE can identify, with a high degree of reliability, the distinguishing signs and symptoms of seven broad categories of drugs.

Drug Evaluation and Classification Technical Advisory Panel: This group was formed to assist the Highway Safety Committee of the International Association of Chiefs of Police on specific matters relating to the Drug Evaluation and Classification Program. These matters include the revision of the approved training curriculum, review and approval of proposed alternative training programs, and other matters relating to the technical aspects of the DEC Program.

Drug Recognition Expert (DRE): An individual who has successfully completed all phases of training requirements for certification established by the International Association of Chiefs of Police and the National Highway Traffic Safety Administration.

Highway Safety Committee: A standing committee of the IACP that addresses highway safety issues.

Horizontal Gaze Nystagmus (HGN): A loss of the normal control of the eyes observed as an involuntary jerking occurring when a person attempts to follow as stimulus with the eyes and/or looks to the left or right side.

Impairment: One of the several terms used to describe the degradation of mental and/or motor abilities necessary for safely operating a motor vehicle.

Implied Consent: Every state has enacted a version of an Implied Consent law, which serves to encourage persons arrested for DWI to submit to a chemical test to determine blood alcohol content. Many states also allow for the testing of blood, breath or urine for the presence of drugs and/or alcohol. The concept of implied consent is that the state views the suspect as already having agreed to take the test, as a condition of operating a vehicle in the state. The typical wording of an implied consent law is as follows: "Any person who operates a motor vehicle upon the public highways of this state shall be deemed to have given consent to a chemical test or tests for the purpose of determining the alcohol (or drug) content of his or her blood, when arrested for any act alleged to have been committed while the person was operating a vehicle while under the influence of alcohol (or any drug)."

The law further provides that, if the arrestee nevertheless refuses to submit to the chemical test, he or she will not be forced to submit, but the driver's license will be suspended or revoked.

IACP Staff: With grant assistance from the National Highway Traffic Safety Administration, the Division of State and Provincial Police of the IACP has agreed to develop standards and assist in managing the certification process for the Drug Evaluation and Classification Program. As part of this agreement, the IACP will perform necessary staff and coordination functions for the program. The staff of the Division of State and Provincial Police is responsible for maintaining records for the program and will coordinate certification and recertification processes.

Instructor Trainer: An experienced DRE instructor who conducts instructor training courses and who must be knowledgeable of and have audited all phases of the Drug Evaluation and Classification training program and must be fully conversant with the student and instructor manuals.

Intoxication: One of the several terms used to describe the degradation of mental and/or motor skills and other faculties due to ingestion of alcohol or other drugs.

NHTSA: The National Highway Traffic Safety Administration, within the United States Department of Transportation that exercises primary responsibility for coordinating federal efforts to ensure the safe design and operation of motor vehicles.

Standardized Field Sobriety Tests: The Standardized Field Sobriety Tests include three tests that were developed and validated through a series of controlled experiments supported by research grants from NHTSA. The three tests include Horizontal Gaze Nystagmus (HGN); Walk and Turn (WAT); and One Leg Stand (OLS).

The HGN test is described elsewhere in these definitions.

Walk and Turn and One Leg Stand are divided attention tests. As such, they require the suspect to concentrate on more than one thing at a time.

The training course developed by IACP and NHTSA, "DWI Detection and Standardized Field Sobriety Testing," is a program designed to train traffic enforcement officers to administer the sobriety tests. The training includes two approved alcohol workshops. During these workshops, students practice administering the test battery. In order to complete the course satisfactorily, students must pass a written examination and demonstrate proficiency in administering the field sobriety test battery.

# STANDARDS FOR THE DRUG EVALUATION AND CLASSIFICATION PROGRAM

## I. Standards for Certification as a Drug Recognition Expert

The standards in this section specify the criteria that must be met prior to an individual's being certified as a drug recognition expert (DRE). These criteria outline the knowledge and skills required to be considered for training, as well as the knowledge and proficiencies required for final certification.

The currently approved curriculum involves a three-phase training process. Prior to beginning the training program, students are required to be trained in and demonstrate proficiency in the use of the IACP/NHTSA-approved standardized field sobriety tests, including the horizontal gaze nystagmus test. Phase I of the drug recognition training consists of a two-day (16-hour) preschool. During this preschool, students are taught the definition of the term "drug" as it is used in the Drug Evaluation and Classification Program, and become familiar with the techniques of the drug evaluation. Students also begin to learn the techniques and procedures for evaluating persons suspected of drug impairment.

Phase II of training is a seven-day (56-hour) classroom program during which students receive detailed instruction in the techniques of the drug evaluation examination as well as in physiology, the effects of drugs and legal considerations. Upon completion of this phase of training, the student must pass a comprehensive written examination before proceeding to Phase III of training, the field certification.

The field certification portion of training follows completion of the classroom training and is conducted at periodic intervals for the next sixty to ninety days. During this portion of the training, students, under the direction of certified instructors, evaluate subjects suspected of being impaired by drugs other than alcohol. After participating in and documenting the results of at least twelve drug evaluations and completing a comprehensive examination, the student is certified as a drug recognition expert.

1.1 In order to be considered for certification as a drug recognition expert, a person shall be in the employ and under the direct control of a public criminal justice agency or institution involved in providing training services to officers of law enforcement agencies.

Commentary: At the discretion of the agency head or administrator, and with the consent of the training body, other persons may audit or observe any or all portions of the DRE training. Persons attending the course as auditors or observers shall not be eligible for certification.

Persons pursuing certification as drug recognition experts for the purpose of instructing in the Drug Evaluation and Classification Program must meet all requirements for certification and recertification in order to maintain their standing as DREs or DRE instructors.

1.2 The candidate DRE must have experience in preparing comprehensive investigative reports and in providing detailed court testimony.

Commentary: The technical nature of the drug evaluation process and the need to provide detailed and accurate documentation of findings and conclusions requires proficiency in preparing reports. Candidate DREs should have demonstrated the ability to investigate, document and prepare detailed reports of incidents such as major traffic crashes or criminal violations. In addition, DREs must be able to provide court testimony concerning their methods and results, as well as their training and qualifications.

1.3 All DRE candidates must attend and complete the IACP/NHTSA-approved course of instruction in Standardized Field Sobriety Testing, or an equivalent curriculum approved by the IACP Highway Safety Committee and Technical Advisory Panel. They shall demonstrate proficiency in the use of Standardized Field Sobriety Tests, to the satisfaction of a DRE instructor, by the conclusion of the IACP/NHTSA DRE Pre-school or a school meeting Standard 1.2 above.

Commentary: The drug evaluation process requires that the contribution of alcohol to observed impairment be determined. The National Highway Traffic Safety Administration has developed, and the IACP has adopted, the Standardized Field Sobriety Test procedure in conjunction with immediate breath testing, as a means of identifying the alcohol-impaired driver. If the effects of alcohol are determined not to be the sole cause of impairment, the officer can begin the evaluation process to determine what other causes may be responsible.

In order to conform to the IACP/NHTSA model curriculum, SFST training must contain the specified number of hours and include at least two approved alcohol workshops. In addition, the training must instruct students in the administration of the horizontal gaze nystagmus, walk and turn, and one leg stand tests.

Each agency should ensure that candidates submitted for DRE training have had adequate time prior to beginning the training program to develop and to demonstrate proficiency in the use of SFST/HGN or allow for refresher training in these techniques as necessary.

1.4 All DRE candidates must attend and complete the IACP/NHTSA DRE Preschool or an IACP-recognized equivalent prior to progressing to Phase II, the DRE School.

- 1.5 Prior to attending phase II of the DRE training, the candidate shall have met the learning objectives for phase I of the training program, the IACP/NHTSA-approved DRE preschool. The candidate shall be able to
  - 1. Define the term "drug" as it is used in the DEC Program;
  - 2. Name the seven drug categories identified in the DRE training program;
  - 3. Measure vital signs, including blood pressure, pulse and body temperature;
  - 4. Show familiarity with the 12-step drug recognition evaluation process;
  - 5. Demonstrate proficiency in the administration of the Standardized Field Sobriety Tests, including Horizontal Gaze Nystagmus;
  - 6. Show familiarity with the administration of the eye examinations, including pupil size, vertical nystagmus and lack of convergence.

These learning objectives are generally met through completion of Phase I, the DRE preschool. However, agencies have the latitude to determine the best means of ensuring that candidate DREs meet the prerequisites. The agency must verify, through the application process to the instructor responsible for delivering the training, that a candidate meets all requirements. Each candidate DRE will be required to demonstrate the knowledge and skills outlined.

Administrative guidelines and suggested application forms containing the necessary information will be provided by IACP staff to agencies and training institutions.

1.6 The candidate DRE shall complete an approved classroom training course which shall, at minimum, achieve the learning objectives as stated in the IACP-approved training curriculum.

Commentary: The National Highway Traffic Safety Administration and the International Association of Chiefs of Police have developed a classroom training course that, when completed, qualifies the student to proceed to the field certification portion of the training program. Because of differences in the type and level of training for officers in the detection of the impaired subject, agencies should determine the most effective means of providing classroom training in drug recognition. However, in order to maintain the credibility and integrity of the certification program, agencies that use a training program other than that currently approved by the IACP, must have the alternative curriculum approved by the IACP Technical Advisory Panel as meeting learning objectives. In addition, the Technical Advisory Panel will be responsible for providing periodic updates and modifications to the IACP training curriculum.

1.7 All candidate DREs shall attend and complete all classroom portions of an approved DRE curriculum prior to progressing to Phase III (the field certification phase) of the training. This shall include satisfactorily completing all assignments and required examinations. Students shall not be permitted to "test out" of portions of the training, nor shall they be permitted to attend only those classes that they have not previously completed.

Commentary: Class sessions missed should be made up prior to the final exam.

1.8 In order to complete satisfactorily the classroom portion of the training and proceed to field certification, candidate DREs must complete an IACP-approved final examination with a score of not less than eighty percent (80%). Candidates scoring less than 80% on the final examination may be retested one time, under the supervision of a certified DRE instructor. The retest shall be completed not less than fifteen nor more than thirty days following the completion of the classroom training.

Commentary: Upon satisfactory completion of the examination, the candidate may then proceed to field certification. The examination used to retest the candidate shall be an IACP-approved examination and shall not have been administered to the candidate previously. If the candidate does not achieve a passing score on reexamination, the candidate must retake the classroom portion of the training and pass the knowledge examination before proceeding further in the certification process.

1.9 Upon completion of the field certification phase of training, the candidate must demonstrate the ability to conduct a complete drug evaluation in an approved sequence and appropriately document and interpret the results. The candidate must also be able to document the findings of the evaluation and demonstrate proficiency in interviewing techniques.

Commentary: One of the primary factors in the success of the Drug Evaluation and Classification Program has been the emphasis upon a standardized approach to the drug recognition process. The training stresses the importance of a systematic, structured approach to performing the drug evaluation. This includes completing all portions of the evaluation in the appropriate sequence. Upon conclusion of an evaluation the DRE reviews the results of all tests, examinations and observations; documents the findings; and draws a conclusion based on the totality of the evidence.

1.10 To be considered for certification as a drug recognition expert, the candidate must satisfactorily complete a minimum of twelve (12) drug evaluations, during which the candidate must encounter and identify subjects under the influence of at least three of the drug categories as described in the DRE training program. All three drug categories must be supported by toxicology.

Of the evaluations required for certification, the candidate shall administer at least six evaluations. The candidate may observe the remaining evaluations. Certification training evaluations will be conducted in accordance with the current procedures and guidelines established in the DECP training curricula.

All evaluations, either administered or observed, and documented for certification purposes, shall be observed and supervised by at least one certified DRE instructor.

Commentary: Ideally, a drug evaluation will be performed by no more than two persons: the evaluator and one observer. At no time should more than four persons participate in an evaluation, as the results of the evaluation may be influenced by the distraction caused by a large number of persons observing the process.

1.11 Prior to completing the certification phase of training, the candidate DRE must demonstrate the ability to draw correct conclusions consistent with observed physiological signs and symptoms. In addition, the conclusions must be supported by the findings of a forensic toxicology laboratory. No candidate DRE shall be certified as a drug recognition expert unless blood, urine, or other appropriate biological samples are obtained and submitted from at least nine (9) subjects whom the candidate DRE has examined for certification purposes. These may include subjects for whom the candidate DRE served as the examination recorder or observer as well as those subjects directly evaluated by the candidate DRE. Further, the candidate DRE cannot be certified unless the opinion concerning the drug category or categories affecting the subject is supported by forensic toxicological analysis seventy-five percent (75%) of the time, or in at least seven (7) of the nine (9) samples submitted for certification purposes. For purposes of this standard, a candidate DRE's opinion is supported if the toxicological analysis discloses the presence of at least one drug category named by the candidate DRE. In the event that the candidate DRE has concluded that three or more categories of drugs are involved, at least two categories must be supported by toxicology results.

Commentary: Successful and uniform application of this standard places important forensic toxicological requirements on the program. First, the blood or urine specimen must be obtained as soon as possible after the arrest so that the contents of the sample refer to the subject's status at the time of the offense. Second, the sample must be properly sealed, stored, transported to the forensic toxicology laboratory and analyzed in a timely fashion to maintain the integrity of the specimen. Third, the drug recognition examination should be conducted as soon as possible after the offense so that the results of the evaluation accurately refer to the subject's status at the time of the offense. Fourth, the laboratory should use its full powers of analysis and detection to attempt to identify each category named by a candidate DRE; in some cases this may require the laboratory to modify its routine screening and confirmation procedures. Finally, the laboratory must complete its report on the samples as soon as possible and provide a copy of the report to the arresting officer, DRE or candidate DRE submitting the sample. It is the submitting officer's responsibility to provide a report to each DRE or candidate DRE who participated in the evaluation.

Although the candidate DRE must complete a minimum of twelve (12) drug evaluations (standard 1.10), standard 1.11 requires only 75 percent of those to include a biological sample. This allows for those cases in which a biological sample is unavailable, such as when a subject refuses or cannot provide one. In those cases when an evaluation is not supported by forensic toxicology, a certified DRE instructor should ensure that the candidate DRE's opinion was based on observable signs and symptoms consistent with the opinion.

1.12 Prior to concluding field certification training, the candidate shall satisfactorily complete an approved "Certification Knowledge Examination." The examination shall be administered and the results reviewed by at least one certified instructor. The examination shall only be administered after the candidate has completed not less than three drug evaluations.

Commentary: The "Certification Knowledge Examination" consists of a comprehensive written examination followed by a detailed interview with the reviewing instructor(s). As stated previously, certification is based on the evaluation by the instructor(s) of the skills and abilities of the candidate rather than on the completion of a specified set of tasks. The purpose of the examination and interview is to aid the instructor(s) in evaluating the candidate's qualifications, performance and general abilities.

The examination should be administered when, in the judgment of the reviewing instructor(s), the candidate has demonstrated proficiency in conducting, evaluating and documenting results of the drug evaluation process.

1.13 The candidate DRE shall complete the field certification phase of training within six months following completion of the classroom training, unless the time limit is extended by the appropriate DRE coordinator.

Commentary: Under normal circumstances, a candidate not completing field certification within the prescribed time period will be dropped from the program. However, a reevaluation of the candidate's qualifications and the reasons for non-completion may be conducted by the appropriate DRE coordinator to determine whether or not circumstances exist that indicate that the candidate should continue in the program.

1.14 By the time the candidate DRE has completed field certification training, the candidate shall have prepared a résumé which shall reflect the candidate's training and experience in drug recognition. The résumé shall include a complete log of all evaluations in which the candidate has participated.

Commentary: In order to be accepted as a credible witness, the drug recognition expert must be able to document and articulate a body of information concerning training, qualifications and experience in the field of drug evaluation and classification. Toward this end, candidates are instructed in the importance and proper preparation of a professional résumé.

1.15 When the candidate DRE has satisfactorily completed all requirements of the classroom and field certification portions of training, at least two certified DRE instructors who have observed the candidate during the field certification process will verify that the candidate meets all requirements for certification as a drug recognition expert.

Commentary: The certification process relies in large part on the judgment of the instructor(s) as to the abilities and performance of the candidate. Experience has shown that in many cases, particularly those in which a candidate's qualifications may be in question, the opinion of a second instructor as to readiness for certification is of value. In addition, the use of a second instructor to evaluate the candidate may overcome any bias, either for or against a candidate. For these reasons, each candidate must be evaluated by at least two instructors prior to becoming certified as a DRE.

1.16 Following completion of certification requirements, copies of all documents, including test results, evaluation logs and drug evaluation reports shall be forwarded to the agency DRE coordinator who shall forward all documents to the state coordinator. The state DRE coordinator shall forward the names and copies of certification progress logs of the DREs they have certified as having successfully completed all phases of the DRE training program. The IACP will then credential each applicant and will register him as a certified drug recognition expert.

Commentary: The IACP staff shall maintain current listings of persons certified as drug recognition experts. Upon notification that a person has met all requirements, staff shall complete and forward to the state coordinator a certificate indicating that he meets all requirements of the Drug Evaluation and Classification Program as a drug recognition expert. The state coordinator shall forward these documents to the agency which, in turn, will present them to the DRE.

### II. Standards for Certification as Drug Recognition Expert Instructor

Because of the highly technical nature of the functions performed by the drug recognition expert, only persons experienced in the techniques of drug evaluation should instruct in the Drug Evaluation and Classification Program. In general, these instructors will be certified drug recognition experts with experience in performing drug evaluations and in providing testimony in court in the area of drug recognition. However, persons who possess specialized skills or credentials may be utilized to teach certain parts of the training course as associate instructors. Dedicated, qualified instructors are critical to the continued success of the Drug Evaluation and Classification Program.

Certified instructors are responsible for observing, evaluating and verifying the performance of candidate DREs throughout the training and certification process. In addition, certified instructors must provide periodic update training to DREs already certified.

Also addressed in this section are standards for the use of instructor trainers in the program. These individuals are responsible for the training of DRE instructors.

2.1 Only persons certified as drug recognition experts may be certified as DRE instructors.

Commentary: Persons not certified as DREs but who possess knowledge, expertise or credentials deemed valuable to the program may be designated as associate instructors for the Drug Evaluation and Classification Program. Persons who might be considered for such designation may include medical professionals, attorneys and others who possess knowledge in a designated field of expertise. Associate instructors must be familiar with the Drug Evaluation and Classification Program and fully conversant with the lesson plans for their assigned blocks of instruction. Classes taught by associate instructors shall be taught in cooperation with certified DRE instructors to ensure consistency.

Each associate instructor should provide to the state coordinator a biographical sketch to be included in the file of approved instructional staff. The biographical sketch shall include those segments of the training curricula that the associate instructor is qualified to teach.

2.2 A DRE desiring to become an instructor in the Drug Evaluation and Classification Program shall make written application to the agency coordinator. The agency coordinator will ensure that the candidate meets all requirements to become an instructor and will refer the application to the state coordinator.

Commentary: The agency head shall verify to the training provider that a candidate instructor meets all prerequisites to enter DRE instructor training. Prerequisites may also include any state, local or agency requirements specified for instructors within the jurisdiction. The state coordinator shall provide to requesting agencies the administrative guide and sample application forms for candidate instructors.

2.3 The candidate shall satisfactorily complete the IACP/NHTSA-approved Drug Evaluation and Classification Instructor Training Program, or an approved equivalent, which shall include both knowledge and practical examination of candidate instructors.

Commentary: This requirement does not preclude states or local jurisdictions from placing additional requirements on persons wishing to teach in the local law enforcement community.

2.4 Upon satisfactory completion of the IACP-approved classroom portion of training or completion of an equivalent program, the student shall be designated as a candidate instructor for purposes of completing instructor certification. To complete instructor certification, the candidate instructor must teach for a minimum of two hours in the classroom portion of an approved drug recognition training program; and supervise the administration of not less than two drug evaluations performed by candidate DREs during certification training.

The candidate instructor's progress shall be monitored and evaluated by at least one certified DRE instructor.

Commentary: The National Highway Traffic Safety Administration and the IACP have developed a training curriculum for instructors in the Drug Evaluation and Classification Program. The learning objectives for this program emphasize specific techniques for teaching the specialized information contained in the drug recognition training program.

The Technical Advisory Panel shall be responsible for reviewing and evaluating alternative training programs submitted by agencies. Those programs meeting or exceeding the approved learning objectives for instructor training shall be deemed "equivalent." This does not preclude agencies or states from adopting more stringent standards.

2.5 Upon satisfactory completion of instructor training, copies of all documentation, including instructor progress logs, examination scores and instructor evaluations, shall be forwarded to the appropriate DRE coordinator. The agency DRE coordinator will forward these documents to the state coordinator who shall certify that they have successfully completed all phases of DRE instructor training. The IACP will then credential each applicant and will register him as a certified DRE instructor.

Commentary: The IACP staff will maintain a current register of persons certified as instructors in the Drug Evaluation and Classification Program. Upon notification that a person has met all requirements, the staff shall complete and forward to the state coordinator a certificate indicating that he/she meets all requirements as a DRE instructor. The state coordinator shall forward these documents to the agency who, in turn, will present them to the DRE instructor.

The administrative guidelines shall provide sample forms for necessary progress logs and certification documents.

**2.6** To ensure the proper conduct and delivery of the approved curriculum, all training sessions conducted as part of the Drug Evaluation and Classification Program shall be coordinated by a certified DRE instructor who has previously instructed. All classes taught by associate or candidate instructors shall be supervised directly by a certified DRE instructor.

Commentary: To ensure that all training classes are conducted in accordance with applicable standards, it is recommended that the instructor coordinating the training program have a minimum of one-year experience as a drug recognition expert instructor.

- 2.7 An instructor trainer shall have demonstrated proficiency as an instructor.
- 2.8 An instructor trainer must be knowledgeable of and have audited all phases of the Drug Evaluation and Classification training program and must be fully conversant with the student and instructor manuals.

Commentary: An instructor trainer must present evidence of the satisfactory completion of the NHTSA/IACP Instructor's Development Course or equivalent. Instructor trainers must be familiar with the Drug Evaluation and Classification Program and fully conversant with the lesson plans for their assigned blocks of instruction. To ensure consistency, classes taught by instructor trainers shall be taught in cooperation with certified DRE instructors.

Each instructor trainer shall provide to the appropriate DRE coordinator a biographical sketch to be included in the file of approved instructional staff. The biographical sketch shall include those segments of the training curricula that the instructor trainer is qualified to teach.

The state coordinator should maintain a record of persons qualified as instructor trainers in the Drug Evaluation and Classification Program.

- 2.9 The course manager shall perform four duties: planning and preparation, onscene course management, data collection, and reporting. These responsibilities involve the following:
  - 1. Assigning instructors, and verifying in advance that the training is conducted in the standardized manner and that it is properly evaluated;
  - 2. Ensuring at the training site that all necessary conditions exist to maximize the students' ability to learn;
  - 3. Collecting certain data following every training event and forwarding it to the host state coordinator; and
  - 4. Preparing a comprehensive report following every training event.

#### III. Standards for Recertification

Recertification is necessary to ensure that DREs and DRE instructors maintain proficiency. Just as the standards in the previous sections have outlined the criteria for original certification, the standards outlined in this section are required to ensure that professional integrity is maintained throughout the recertification process.

3.1 The following records concerning certification and recertification shall be maintained:

Individual DRE/

Copies of all drug evaluations

DRE Instructor

Evaluation logs

Resume

Certification and recertification progress logs

Certificates

Agency DRE Coordinator

Copies of evaluation logs

Certification progress logs

Copies of certificates

Instructor ratings and summaries of student critiques

Records of classes taught by each instructor

State DRE Coordinator and/or IACP Staff

Copies of evaluation logs (optional)

Certification progress logs

File of certified DREs and instructors

Recertification information

Commentary: Guidelines for the retention of pertinent records concerning the program operation help to ensure integrity of the program and provide valuable information for purposes of statistics and court verification of training. Other records as deemed appropriate by local agencies or certification commissions may be required of the individual DRE or the appropriate DRE coordinator.

- 3.2 DREs shall be required to renew their certificates of continuing proficiency every two years. A one-year grace period following the lapse of certification may be allowed for those not meeting recertification standards. During the grace period, the DRE may be rectified without having to repeat the original certification process.
- 3.3 The state coordinator shall be notified of those DREs in need of recertification at least six months prior to the expiration of the certificates. The state DRE coordinator shall forward to the IACP staff required documentation indicating the completion of recertification requirements. The staff will issue new cards when requirements are met.

Commentary: In the absence of a state coordinator, the TAP regional coordinator will perform these functions.

3.4 A DRE shall demonstrate continuing proficiency by

Performing a minimum of four (4) acceptable evaluations since the date of last certification, all of which shall be reviewed and approved by a certified DRE instructor and one (1) of which shall be witnessed by a certified DRE instructor. These evaluations may be performed on subjects suspected of drug and/or alcohol impairment or during classroom simulations; and Completing a minimum of eight hours of recertification training since the date of the DRE's most recent certification, which may alternatively be presented in two sessions of no less than four hours, and which shall be consistent with any IACP standards for such training; and Presenting an updated resume and rolling log to the appropriate coordinator or his/her designee for review.

Commentary: All coordinators are responsible for maintaining the integrity of the program, and the appropriate coordinator, consistent with this responsibility, is encouraged to withhold recertification for, or refer for remediation, any DRE whose rolling log indicates an unacceptable level of accurate evaluations, as indicated by toxicology results.

- 3.5 When a DRE has completed all requirements for recertification, a certified DRE instructor shall verify to the appropriate DRE coordinator that minimum recertification requirements have been met.
- 3.6 A certified instructor shall maintain instructor certification so long as DRE certification is maintained.

Commentary: An instructor may be decertified for cause, such as for conducting substandard instructional programs, and still maintain certification as a DRE.

# IV. Standards for Decertification of Drug Recognition Experts and Instructors

The standards in this section outline the circumstances and procedures for decertifying individual DREs or DRE instructors. In order to ensure that standards of performance are maintained, a means is needed for removing from the roles of the program those persons unable to meet the criteria of competence and professionalism. The responsibility for maintaining program standards lies with the agency and the appropriate DRE coordinator. It shall be incumbent upon all DRE coordinators to ensure that certified DREs meet approved standards for conduct and qualifications.

4.1 Decertification shall occur when a DRE or DRE instructor fails to meet minimum standards and requirements for certification or recertification, or demonstrates evidence of poor performance, inconsistent findings, or other substantiated acts on the part of the DRE that reflect discredit upon the Drug Evaluation and Classification Program.

Commentary: All DREs are responsible for maintaining and forwarding to the appropriate DRE coordinator information regarding required training or experience. If such information is not provided in a timely manner, certification will lapse.

Local agencies and licensing/certification bodies may, at their discretion, establish certification and decertification criteria to conform to local laws or rules. Nothing in these standards should be construed to overrule local authority in establishing standards no less stringent for the performance of officers in this area or to prevent an agency from following internal disciplinary or administrative personnel procedures.

- 4.1.1 Before decertification is finalized, a DRE or DRE instructor will be given written notice by the initiating DRE coordinator of the reasons for decertification. The subject of the action shall have the opportunity for a written or an oral response to the initiating DRE coordinator.
- 4.2 Requests for voluntary description will be honored when submitted by a DRE or DRE instructor to the section IACP staff and with approval of the agency appropriate DRE coordinator.
- 4.3 Cases involving poor performance or inconsistent findings shall be referred to the agency appropriate DRE coordinator for investigation, recommendation and action.
- 4.4 Certification of a DRE shall not terminate as long as the DRE meets the requirements of Standards 1.1 and 4.1.

4.5 The state DRE coordinator, upon the recommendation of the agency DRE coordinator or based on substantiated independent knowledge shall initiate the decertification process against a DRE or DRE instructor. The state coordinator shall inform the IACP staff of all decertification actions. In instances where these complaints have not been resolved by the appropriate coordinator, these complaints will be referred to the state's Governor's Office of Highway Safety for resolution.

# V. Standards for Reinstatement of a Decertified Drug Recognition Expert

The standards in this section outline the procedures for reinstating a previously decertified DRE and/or DRE instructor.

- **5.1** An individual can be reinstated as a DRE when the following conditions are met:
  - (1) The applicant must pass the 100-item exam (same as that given at the end of the DRE school, or the make-up exam) as witnessed by a certified DRE instructor, with a score of at least 80%.
  - (2) The applicant must complete four (4) hands-on drug evaluations within a one-year period from the date of request to be reinstated.
  - (3) The applicant's eligibility and reinstatement as a DRE is reviewed and approved by the DRE's agency and the agency, state, and TAP regional DRE coordinators, where applicable.
- **5.2** An individual can be reinstated as a DRE instructor when the following conditions are met:
  - (1) The applicant meets conditions 5.1 and is reinstated as a DRE.
  - (2) The applicant's eligibility and reinstatement as a DRE instructor is reviewed and approved by the DRE's agency and the agency, state, and TAP regional DRE coordinators, where applicable.

Commentary: In many instances, a DRE certification lapses through no fault of the DRE due to transfers, promotions, etc., and recertification requirements have not been met. In many cases a DRE may want to reapply DRE skills with a new assignment. IACP suggests that a written request for reinstatement to the program come from the applicant to the appropriate coordinator, through the proper agency channels. A form is provided by the IACP to DEC state and TAP regional coordinators for the purpose of reinstatement. All coordinators are cautioned to conduct a thorough check on the cause of the applicant's decertification and reason for application for reinstatement.

### VI. Standards for Agency Participation

Since 1986, the National Highway Traffic Safety Administration has endeavored to expand the Drug Evaluation and Classification Program. In an effort to contain costs, ensure the most efficient use of resources and maintain a high probability of program success, NHTSA developed site selection criteria to be used in assessing potential suitability of sites. Factors such as demographics, favorable legislation, agency operations and system support for the program are considered in evaluating potential sites for the implementation of the Drug Evaluation and Classification Program.

It is recognized that law enforcement agencies, in considering the implementation of new traffic enforcement programs, must be aware of both short- and long-term costs that are involved. In order for the program to achieve maximum results, the Drug Evaluation and Classification Program requires that agencies commit considerable resources long term to the detection and apprehension of the drug-impaired driver.

**6.1** A DEC Program site should be a state, a political subdivision of a state, or a group of subdivisions.

Commentary: Experience has shown that a DEC Program will take firm root only if the resources to support the program are concentrated in a relatively small geographical area, such as a major city or county. Given that these new sites will begin operations with a small cadre of DREs, a community-focused DEC Program will allow the DREs to respond quickly to the location(s) where drug-impaired drivers might be taken for processing. By concentrating its forces, the program can ensure that a qualified DRE is available at any time or place needed. The concentrated focus of a community-based program allows the DREs ample opportunity to conduct evaluations and maintain skills at peak proficiency.

**6.2** A proposed program site should be able to produce enough drug-impaired driving arrests to (1) justify the expense of training the DREs, and (2) provide enough evaluation opportunities for DREs to maintain proficiency.

Commentary: Studies indicate that up to 40 percent of the persons arrested for impaired driving are actually under the influence of drugs, either alone or in combination with alcohol. Thus, a site should produce an adequate number of DUI arrests annually per DRE to provide ample drug evaluation opportunities.

**6.3** Prior to implementation of a DEC Program, a site should be located in a state with an implied consent law that

Explicitly allows the chemical test sample to be analyzed to determine the presence and/or concentration of drugs other than alcohol:

Explicitly indicates that the "consent" applies to multiple tests, i.e., that the person is "deemed to have given consent to a test or tests of blood, breath or urine"; and

Empowers the arresting officer and/or the law enforcement agency to select the types of chemical tests to be taken, rather than giving the suspect the option of choosing the tests.

In the absence of an implied consent law, a site must certify that the above three criteria are met and apply to the Technical Advisory Panel for consideration for acceptance to the program.

Commentary: It is pointless to evaluate drivers for drug-induced impairment unless those found to be so impaired can be prosecuted successfully. The requirements for multiple chemical tests are essential because both a breath test and blood or urine tests are integral components of the drug recognition process. In addition to implied consent legislation, the effectiveness of DEC programs is greatly enhanced by legislation that

Allows the fact of a suspect's refusal to submit to the chemical test to be introduced as evidence in court; and

Makes it an offense to drive under the influence of any drug.

**6.4** At least eighty percent (80%) of a participating agency's traffic law enforcement officers must be fully trained and proficient in the use of the IACP/NHTSA-approved standardized field sobriety tests, including the horizontal gaze nystagmus test.

Commentary: It is recommended that the agency's SFST training program is consistent with the IACP/NHTSA model curriculum. In particular, the training must contain the specified number of hours and include at least two approved alcohol workshops.

- 6.5 Participating agencies must maintain accurate and timely records of
  - Alcohol and drug-related arrests and convictions;
  - Alcohol and drug offense processing time;
  - All toxicological examinations; and
  - All drug recognition evaluations to include documenting and collecting of basic data which includes, but is not limited to, the name and age of arrestee, date of arrest, sex, the DRE opinion, and the name of evaluator.

Commentary: In order to evaluate critically the effectiveness of the Drug Evaluation and Classification Program, it is necessary that, at a minimum, the above records be maintained. In addition to evaluation purposes, the records may prove beneficial in establishing program validity for court purposes. The IACP and NHTSA has endorsed a data collection software program which DECP states are encouraged to use.

**6.6** Participating agencies should have the capability to establish centralized booking or processing of all DUI arrestees.

Commentary: The ideal situation is one in which all persons arrested for DUI are taken to a single location for processing. One or two DREs could then be stationed at that location to ensure prompt access to all suspects apprehended for drug-impaired driving. However, it is feasible for a jurisdiction to have a few centralized processing facilities as long as there are enough DREs to staff them adequately and enough DUI arrests to ensure that the DREs conduct frequent evaluations.

**6.7** Each location where DRE evaluations are conducted must have adequate facilities, including

A room sufficiently large to permit unobstructed administration of the Standardized Field Sobriety Tests;

A separate room that can be completely darkened for the eye examination;

Storage space for test data forms, reference documents, blood pressure kits, etc;

Access to breath testing equipment producing on-the-spot results; and

Facilities and materials for collecting blood and/or urine samples.

Commentary: Because of the unique requirements of the DEC Program, it is sometimes more economical for several agencies within a site to share DUI processing facilities. Other desirable characteristics for a DUI processing facility include

Adequate holding cells for arrestees;

Separate interrogation and report writing areas that provide privacy from the general prisoner population; and

Testing facilities that are out of main traffic patterns and allow the drug evaluation process to be performed without interruption or distraction.

**6.8** Participating agencies must have access to laboratories that are capable of identifying the presence of the most commonly abused drugs when these drugs are present in sufficient concentrations to produce impairment.

Commentary: Ideally, the laboratories will also be able to identify the concentration of these drugs. In any case, the accuracy of the chemical analysis should be consistent with state-of-the-art drug testing. In other words, screening tests are not sufficient; a jurisdiction should be able to produce a confirmatory analysis. Although either blood or urine samples are acceptable, it is best if the jurisdiction has the ability to test both.

**6.9** All agencies and states interested in participating in a Drug Evaluation and Classification Program must have the following endorsements:

The state governor's representative for highway safety;

The chief elected official of each political subdivision to be included in the site;

The commanding officer of each participating law enforcement agency;

The administrative judge of each court that tries people arrested for DUI within the jurisdiction;

The chief prosecuting attorney for each court in the jurisdiction; and

Representatives of any other agencies that would be involved in covering the costs of developing and sustaining the DEC Program.

# DRUG EVALUATION AND CLASSIFICATION PROGRAM ADMINISTRATIVE GUIDELINES INTERNATIONAL ASSOCIATION OF CHIEFS OF POLICE

With grant assistance from the National Highway Traffic Safety Administration (NHTSA), the International Association of Chiefs of Police has developed certification standards and administers the Drug Evaluation and Classification Program. Under these administrative guidelines, it will be the responsibility of the individual and all coordinators to ensure that specific requirements of the standards are met. The staff at the IACP will be responsible for maintaining records, issuing certificates of completion, coordinating certain training-related events and maintaining and updating training materials as required.

The following procedures have been developed by the staff of the International Association of Chiefs of Police for use by agencies participating in the Drug Evaluation and Classification Program and wishing to certify drug recognition experts and instructors in their employ.

Obtaining certification as a drug recognition expert or DRE instructor ensures that an individual meets minimum requirements for training and experience as established by the IACP and the IACP Technical Advisory Panel. The Drug Evaluation and Classification Administrative Guidelines accompany the International Standards of the Drug Evaluation and Classification Program.

For the certification process to operate efficiently, it is recommended that coordinators at the agency, and state, and regional levels be identified. The responsibilities of the coordinators may include reviewing the qualifications of the candidate DREs, supplying required documentation that minimum standards have been met, and maintaining individual and program records. The coordination functions may be performed by one person or may be divided among several persons, as operational needs demand.

# 1. Notification of Candidate Drug Recognition Experts

When an individual has completed all agency application requirements for admission for training as a drug recognition expert, the agency shall provide the following information to the appropriate coordinator:

- 1. Candidate's name
- 2. Mailing address
- 3. Sponsoring agency
- 4. Social security number
- 5. Verification that candidate has satisfactorily completed a NHTSA/IACP-approved course in Standardized Field Sobriety Testing

In addition, the appropriate DRE coordinator shall provide the above information to the agency or individual responsible for providing training to ensure that all students meet prerequisites prior to the beginning of the training phase:

State program coordinators shall forward to the IACP staff the above information on all candidate DREs at the following address:

International Association of Chiefs of Police Division of State and Provincial Police 515 North Washington Street Alexandria, VA 22314

# 2. Obtaining Certification as a Drug Recognition Expert

All candidates for certification under the International Drug Evaluation and Certification Program must demonstrate completion of all requirements specified in Section I of the International Standards of the Drug Evaluation and Classification Program. Each candidate's progress toward meeting certification requirements shall be documented on the "Certification Progress Log," which shall be supplied to all appropriate DRE coordinators by the IACP staff. Each candidate shall be responsible for maintaining a certification progress log.

Completion of each step in the certification process shall be verified by the signature of at least one certified DRE instructor. Final recommendation for certification must be verified by the signatures of two certified instructors. Upon completion of all certification requirements, copies of the certification progress log shall be forwarded to the agency DRE coordinator and to the state coordinator. The state coordinator shall verify all information on the certification *progress* log and ensure that all entries are correct. The state coordinator shall forward to the IACP staff a copy of each candidate's completed certification progress log.

Upon receipt of the completed certification progress log, the IACP staff shall ensure that all necessary information is complete. Upon verifying that the information is complete, the IACP staff shall forward to the DRE state or TAP regional coordinator a certificate of completion and an identification card signifying that the candidate has met or exceeded all requirements for certification as a drug recognition expert. In the event that proper documentation is not provided, notification will be sent to the state coordinator indicating the specific reasons(s) for non-qualification.

The IACP staff shall maintain records of all certified DREs. Each record will contain the following information:

- 1. Name
- 2. Social Security Number
- 3. Department/agency
- 4. Mailing address

- 5. Telephone number
- 6. Dates of all events specified on the progress log
- 7. Name(s) of instructors verifying completion of training events
- 8. Date certificate is awarded
- 9. Date certification expires

### 3. Obtaining Certification as DRE Instructor

Candidates for certification as DRE instructors must demonstrate that they meet all requirements specified in Section II of the *International Standards of the Drug Evaluation and Classification Program*. The candidate instructor's progress toward completing certification requirements shall be documented on the form, "DRE Instructor's Certification Progress Log," which shall be supplied by IACP staff to all appropriate DRE coordinators. The individual candidate DRE instructor shall be responsible for maintaining the log.

Completion of each step in the instructor certification phase shall be verified by at least one certified DRE instructor. Upon completion of all certification requirements, copies of the DRE instructor's certification progress log shall be forwarded to the agency DRE coordinator and to the state DRE coordinator. The state DRE coordinator, after verifying that all information on the logs is complete and accurate, shall forward copies of all completed instructors' certification progress logs to the IACP staff.

Upon receipt of the instructor certification progress log, the IACP staff shall verify that all information on the log is complete. Upon verification, the IACP staff shall forward to the state coordinator a certificate of completion signifying that the candidate meets or exceeds all requirements of the Drug Evaluation and Classification Program as a DRE instructor. The IACP staff shall send notification to the state coordinators that the instructor has been certified. In the event that the instructor does not meet all requirements for certification, notification will be sent to the state coordinators indicating the specific reason(s) for non-qualification.

The IACP staff will maintain records of all certified DRE instructors. Each record will contain the following information:

- 1. Name
- 2. Social Security Number
- 3. Department/agency
- 4. Mailing address
- 5. Telephone number
- 6. Dates of all training events specified in the progress log
- 7. Name(s) of instructors verifying completion of training events
- 8. Date certificate was awarded
- 9. All pertinent information relating to the instructor's experience and credentials

Drug recognition expert instructors shall maintain certification as long as DRE certification is maintained. State coordinators will maintain a list of persons designated as associate instructors or as instructor trainers for the Drug Evaluation and Classification Program. In order that the list for instructors and associate instructors may be kept current and, therefore, of use to the participants, agencies hosting DRE training events (pre-schools, DRE training, instructor schools) should provide the state coordinator a list of all instructors and their instruction assignments.

# 4. Procedures for Recertification of Drug Recognition Experts and DRE Instructors

As specified in Section III of the *International Standards of the Drug Evaluation* and Classification Program, all drug recognition experts must be recertified every two years following original certification. DRE instructors shall maintain their instructor certification as long as DRE certification remains in effect. All applicable recertification standards for DREs shall apply to DRE instructors.

The following process will be utilized to ensure timely notification and compliance with recertification requirements:

- 1. Eighteen (18) months following the date of original certification, the IACP will send a renewal advisory notice to state DRE coordinators.
- 2. The DRE shall forward to his state coordinator evidence of completion of all recertification requirements as well as a recertification form signed by his agency coordinator. The state coordinator, after signing the recertification form, will forward a copy to IACP staff.
- 3. Upon notification that a person has met all requirements under section III of the *International Standards of the Drug Evaluation and Classification Program*, IACP staff shall issue a card recertifying the DRE for a period of two years.

In the event that information verifying completion of recertification requirements is not received by the IACP staff prior to the expiration of certification, the IACP staff will notify the state coordinators that certification has expired. Following expiration of certification, the DRE may renew certification without penalty for a period of one year by providing proof of completion of recertification requirements. A decertified DRE wishing to be reinstated following the expiration of the one-year grace period must complete all training and certification requirements enumerated in Section V of the International Standards of the Drug Evaluation and Classification Program.

#### 5. Decertification of Drug Recognition Experts

Decertification of a drug recognition expert may take place if one or more of the following conditions exist:

- 1. The requirements as enumerated in Section III of the *International Standards of the Drug Evaluation and Classification Program* are not met by the individual DRE, allowing certification to lapse.
- 2. A DRE voluntarily requests decertification.
- 3. There is evidence of poor performance, inconsistent findings, or other acts on the part of the DRE that reflect discredit upon the Drug Evaluation and Classification Program.

In the case of a lapse of certification, the procedures in Section 4 of the Administrative Procedures shall be followed.

A DRE wishing to be decertified shall submit a written request through the appropriate agency and state coordinators to the IACP staff. Upon receipt of approval of the request by the state DRE coordinator, IACP staff shall remove the name of the individual from the list of certified DREs.

Agency DRE coordinators shall monitor the performance of DREs within their agencies and shall investigate complaints arising from their activities in the drug evaluation area. When, in the opinion of the agency coordinator, and with the approval of the agency head or his designee, a DRE's actions warrant decertification, the agency shall notify the state coordinator that the DRE is no longer certified within that agency.

Nothing in this procedure should be construed as to prevent an agency from following internal disciplinary or administrative personnel procedures. The IACP staff will maintain records of all decertified DREs and the reason(s) for decertification.

# 6. Approval of Drug Recognition Training Curricula

The National Highway Traffic Safety Administration (NHTSA) and the International Association of Chiefs of Police (IACP) have developed a course of instruction to train police officers in the techniques of drug recognition. This course of training has been adopted by the IACP as the minimum training requirement for certification for DREs and DRE instructors. NHTSA and IACP are responsible for revising and updating the DRE training curricula.

The course of instruction adopted by the IACP requires a total of seventy-two hours of classroom instruction followed by field certification during which a candidate must participate in a minimum of twelve drug evaluations. In the course of the required drug evaluations, a candidate must encounter and correctly identify subjects under the influence of at least three different categories of drugs. The complete requirements for certification as a DRE are enumerated in Section I of the International Standards of the Drug Evaluation and Classification Program.

In recent years, several training programs have been developed by police agencies and commercial training institutions with the aim of training individuals to detect persons impaired by drugs. A number of agencies currently utilize portions of the NHTSA/IACP approved program or variations of it in teaching officers the techniques of detecting the drug-impaired driver.

Section I of the International Standards of the Drug Evaluation and Classification Program requires that a candidate for certification complete "...an approved classroom training course which shall, at minimum, achieve the learning objectives as stated in the IACP approved training curriculum." The Highway Safety Committee of the IACP is charged with overseeing the operation and development of the Drug Evaluation and Classification Program. In order to maintain the high standards of the program, the committee has established the Technical Advisory Panel. Responsibilities of this panel, appointed by the IACP Highway Safety Committee, include the review of proposed alternative training programs to determine whether or not course content and learning objectives are consistent with approved standards.

Organizations wishing to submit proposed training curricula for review and approval as equivalent programs for the purpose of certifying individuals as drug recognition experts shall submit lesson plans, visual aids and any other required materials to the IACP staff. The IACP staff will submit the proposed course to the Technical Advisory Panel for evaluation. Courses that meet applicable standards and learning objectives shall be termed as equivalent courses. Completion of said courses shall qualify the candidate for certification as a DRE.