



Department of Health and Human Services
Maine Center for Disease Control and Prevention
286 Water Street
11 State House Station
Augusta, Maine 04333-0011
Tel: (207) 287-8016; Fax (207) 287-9058
TTY Users: Dial 711 (Maine Relay)

Maine Health Alert Network (HAN) System

PUBLIC HEALTH ADVISORY

To: Health Care Providers
From: Dr. Isaac Benowitz, State Epidemiologist
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Please review this information about an increased risk of Oropouche virus (OROV) infection in the U.S. associated with an increase in global Oropouche activity and key actions clinicians can take to recognize Oropouche, order appropriate testing, ensure timely reporting to public health authorities, and promote prevention measures.

Oropouche virus is an emerging arthropod-borne virus spread by biting midges and mosquitoes. Initial symptoms are non-specific and may resemble chikungunya, dengue, and Zika. Symptoms usually start 3–10 days after a bite from an infected midge or mosquito and resolve after a few days, but 70% experience recurrent symptoms days to weeks after initial illness.

Clinicians should consider OROV infection in symptomatic patients with recent travel to any area with OROV activity. Because dengue co-circulates in many areas with OROV activity, dengue infection should be ruled out before testing for OROV. **Testing for OROV, for symptomatic patients meeting the [suspect case definition](#), is available only at U.S. CDC and must be approved by Maine CDC.** If approved, samples can be sent to the Maine Health and Environmental Testing Laboratory (HETL). Clinicians can refer to U.S. CDC [specimen requirements](#) to determine the proper specimens to submit for testing. A HETL [Requisition Form](#) and [Human Arbovirus Submission Form](#) are required with all submissions.

U.S. CDC: Increased Oropouche Virus Activity and Associated Risk to Travelers

Summary

The U.S. Centers for Disease Control and Prevention (U.S. CDC) is issuing this Health Alert Network (HAN) Health Advisory to notify clinicians and public health authorities of an increase in Oropouche virus disease in the Americas region, originating from endemic areas in the Amazon basin and new areas in South America and the Caribbean. Between January 1 and August 1, 2024, more than 8,000 cases of Oropouche virus disease were reported, including two deaths and five cases of vertical transmission associated with fetal death or congenital abnormalities. Countries reporting cases include Brazil, Bolivia, Peru, Colombia, and Cuba. In the United States and Europe in 2024, travel-associated cases have been identified in travelers returning from Cuba and Brazil. As testing and surveillance for Oropouche virus disease increase in the Americas, reports of cases from additional countries are expected. This Health Advisory advises on evaluating and testing travelers who have been in impacted areas with signs and symptoms consistent with Oropouche virus infection. It also raises awareness of the possible risk of vertical transmission (e.g., from gestational parent to fetus during pregnancy) and associated adverse effects on pregnancy and highlights prevention measures to mitigate additional spread of the virus and potential importation into unaffected areas, including the United States.

Background

[Oropouche virus](#) belongs to the Simbu serogroup of the genus *Orthobunyavirus* in the *Peribunyaviridae* family. The virus was first detected in 1955 in Trinidad and Tobago and is endemic in the Amazon basin. Previous outbreaks have been described in Bolivia, Brazil, Colombia, Ecuador, French Guiana, Panama, and Peru. One child was infected in Haiti in 2014. The current 2024 outbreak is occurring in endemic areas and new areas outside the Amazon basin; countries reporting locally acquired (autochthonous) cases include Brazil, Bolivia, Peru, Colombia, and Cuba. Although travel-associated cases have been identified in the United States (n=11), no evidence of local transmission currently exists within the United States or its territories.

Sylvatic (enzootic) transmission of Oropouche virus occurs in forested areas between mosquitoes and non-human vertebrate hosts (e.g., sloths, non-human primates, domestic and wild birds, and rodents). Humans can become infected while visiting forested areas and are likely responsible for introducing the virus into urban environments. Humans contribute to the transmission cycle in urban environments since infected humans develop sufficient viremia to serve as amplifying hosts. Biting midges (*Culicoides paraensis*) and possibly certain mosquitoes (*Culex quinquefasciatus*) are responsible for transmitting the virus from an infected person to an uninfected person in urban areas.

Approximately 60% of people infected with Oropouche virus become [symptomatic](#). The incubation period is typically 3–10 days. Initial clinical presentation is similar to diseases caused by [dengue](#), [Zika](#), and [chikungunya](#) viruses, with acute onset of fever, chills, headache, myalgia, and arthralgia. Other symptoms can include retroorbital (eye) pain, photophobia (light sensitivity), nausea, vomiting, diarrhea, fatigue, maculopapular rash, conjunctival injection, and abdominal pain. Clinical laboratory findings can include lymphopenia and leukopenia, elevated C-reactive protein (CRP), and slightly elevated liver enzymes. Initial symptoms typically resolve after a few days, but a high proportion (about 70%) experience recurrent symptoms days to weeks after resolution of their initial illness. Although illness is typically mild, it is estimated less than 5% of patients can develop hemorrhagic manifestations (e.g., epistaxis, gingival bleeding, melena, menorrhagia, petechiae) or neuroinvasive disease (e.g., meningitis, meningoencephalitis). Neuroinvasive disease symptoms may include intense occipital pain, dizziness, confusion, lethargy, photophobia, nausea, vomiting, nuchal rigidity, and nystagmus. Clinical laboratory findings for patients with neuroinvasive disease include pleocytosis and elevated protein in cerebrospinal fluid (CSF).

Although people exposed to biting midges or mosquitoes infected with the virus are most at risk for developing disease, the risk factors for more severe Oropouche virus disease are not well-defined. People at risk for more severe disease likely include those at risk for severe disease with other viral infections transmitted by vectors (e.g., people aged 65 years or older, or those with underlying medical conditions, such as immune suppression, hypertension, diabetes, or cardiovascular disease). Earlier this year, Brazil reported two deaths in otherwise healthy non-pregnant women, and five cases in pregnant people with evidence of vertical transmission of the

virus to the fetus associated with fetal death or congenital abnormalities, including microcephaly. This was the first report of deaths and Oropouche virus vertical transmission and associated adverse birth outcomes.

[Laboratory diagnosis](#) is generally accomplished by testing serum. Cerebrospinal fluid can also be tested in patients with signs and symptoms of neuroinvasive disease. Diagnostic testing is available at some public health laboratories (e.g., Wadsworth Center, NYS Department of Health) and at U.S. CDC. U.S. CDC and other public health laboratories are currently working to validate additional diagnostic assays. Contact Maine CDC at 1-800-821-5821 for more information and to facilitate testing. For current testing and case reporting guidance, [visit U.S. CDC's website](#). In many countries, [outbreaks of dengue](#) are occurring in areas with reported Oropouche virus transmission. For patients with suspected Oropouche virus disease, it is important to rule out dengue virus infection because proper clinical management of dengue can improve health outcomes. Other diagnostic considerations include chikungunya, Zika, leptospirosis, malaria, or infections caused by various other bacterial or viral pathogens (e.g., rickettsia, group A streptococcus, rubella, measles, parvovirus, enteroviruses, adenovirus, Mayaro virus).

No specific antiviral [treatments](#) or vaccines are available for Oropouche virus disease. Treatment for symptoms can include rest, fluids, and use of analgesics and antipyretics. Acetaminophen is the preferred first-line treatment for fever and pain. Aspirin and other non-steroidal anti-inflammatory drugs (NSAIDs) should not be used to reduce the risk of hemorrhage. Patients who develop more severe symptoms should be hospitalized for close observation and supportive treatment. Pregnant people with laboratory evidence of Oropouche virus infection should be [monitored during pregnancy](#) and live-born infants should be carefully evaluated.

Travelers to areas with Oropouche virus transmission should use prevention measures to avoid biting midge and mosquito exposure during travel and for 3 weeks after travel, or if infected during the first week of illness, to mitigate additional spread of the virus and potential importation into unaffected areas in the United States. Oropouche virus disease is not a nationally notifiable condition.

Recommendations for Health Care Providers

- Consider Oropouche virus infection in a patient who has been in an area with documented or suspected Oropouche virus circulation within 2 weeks of *initial* symptom onset (as patients may experience recurrent symptoms), and the following:
 - Abrupt onset of reported fever, headache, and one or more of the following: myalgia, arthralgia, photophobia, retroorbital/eye pain, or signs and symptoms of neuroinvasive disease (e.g., stiff neck, altered mental status, seizures, limb weakness, or cerebrospinal fluid pleocytosis); AND
 - No respiratory symptoms (e.g., cough, rhinorrhea, shortness of breath); AND
 - Tested negative for other possible diseases, in particular dengue. If strong suspicion of Oropouche virus disease exists based on the patient's clinical features and history of travel to an area with virus circulation, do not wait for negative testing for other infections before contacting Maine CDC.
- Contact Maine CDC at 1-800-821-5821 to report all suspected Oropouche virus disease infections, facilitate diagnostic testing, and help mitigate risk of local transmission.
- [Rule out dengue virus infection](#) in travelers with suspect Oropouche virus infection because these viruses often cocirculate and cause similar clinical presentations during acute illness. Early clinical management of dengue can improve health outcomes.
- Be aware that a high proportion of patients (about 70%) with Oropouche virus disease may experience recurrent symptoms days to weeks after resolution of their initial illness.
- Be aware of the risk of vertical transmission and possible adverse impacts on the fetus, including fetal death or congenital abnormalities. [Monitor pregnancies](#) in people with laboratory evidence of Oropouche virus infection and provide thorough infant evaluations.
- Inform pregnant people of the possible risks to the fetus when considering travel to areas with reported Oropouche virus transmission. Counsel these patients to consider the destination, reason for traveling, and their ability to prevent insect bites.
- Pregnant people are currently recommended to reconsider non-essential travel to areas with an Oropouche virus Level 2 [Travel Health Notice](#). If a pregnant person decides to travel, counsel them to strictly prevent insect bites during travel.

- Manage travelers with suspect Oropouche virus disease with acetaminophen as the preferred first-line treatment for fever and pain. Aspirin and other NSAIDS should not be used to reduce the risk of hemorrhage.
- Be aware that people who may be at higher risk for complications or severe disease include pregnant people, older adults (e.g., aged 65 years or older), and people with underlying medical conditions (e.g., immune suppression, hypertension, diabetes, or cardiovascular disease).
- Direct all travelers going to areas with Oropouche virus transmission to use measures to prevent insect bites during travel and for 3 weeks after travel, or if infected, during the first week of illness to mitigate additional spread of the virus and potential importation into unaffected areas in the United States.

Recommendations for Travelers

- All travelers can protect themselves from Oropouche, dengue, Zika, and other viruses transmitted by insects by [preventing insect bites](#), including using an [Environmental Protection Agency \(EPA\) registered insect repellent](#); wearing long-sleeved shirts and pants; and staying in places with air conditioning or that use window and door screens.
- Pregnant travelers should discuss travel plans, reasons for travel, steps to prevent insect bites, and potential risk with their health care provider.
- Pregnant people considering travel to countries with an Oropouche virus Level 2 [Travel Health Notice](#) should reconsider non-essential travel. If travel is unavoidable, pregnant travelers should strictly follow Oropouche virus [prevention recommendations](#) to prevent insect bites during travel.
- Travelers should be aware that the [most common symptoms of Oropouche virus](#) are fever and headache and that symptoms usually begin 3-10 days after being bitten by an infected midge or mosquito. Most people infected with Oropouche virus feel better within a week, but symptoms often come back.
- Travelers who have been in areas with Oropouche virus transmission should [prevent insect bites](#) for 3 weeks after travel.
- Travelers to areas with Oropouche virus transmission, including South America or the Caribbean, who develop fever, chills, headache, joint pain, or muscle pain during or within 2 weeks after travel, should:
 - Seek medical care and tell their health care provider when and where they traveled.
 - Not take aspirin or other NSAIDS (e.g., ibuprofen) to reduce the risk of bleeding.
 - Continue to [prevent insect bites](#) during the first week of illness to avoid further spread, especially in areas where mosquitoes or biting midges are active.

For More Information

- [About Oropouche | U.S. CDC](#)
- [Travel Health Notices | U.S. CDC](#)
- [Preventing Mosquito Bites | U.S. CDC](#)
- [Find the Repellent that is Right for You | EPA](#)
- [Dengue: Guidelines for Diagnosis, Treatment, Prevention and Control | WHO](#)

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