

# THIS IS AN OFFICIAL NH DHHS HEALTH ALERT

Distributed by the NH Health Alert Network  
[Health.Alert@nh.gov](mailto:Health.Alert@nh.gov)  
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## Arboviral Disease in New Hampshire: Preparation for the 2015 Season

### NH Division of Public Health Services (NH DPHS) recommends:

1. Vigilance is needed during the summer months to consider mosquito-borne diseases, including West Nile Virus (WNV) and Eastern Equine Encephalitis (EEE), in patients with compatible clinical features. Laboratory testing is recommended and may be arranged by calling (603) 271-4496 during business hours or (603) 271-5300 after hours. Forms and human testing information are available at <http://www.dhhs.state.nh.us/dphs/cdcs/arboviral/index.htm>.
2. Remind patients to take preventive measures, including avoiding mosquito bites by use of insect repellents and wearing protective clothing, and environmental reduction of mosquito populations.
3. Equine vaccines are available for both EEE and WNV and should be used to protect horses. Vaccines are not available for human use.
4. Report all arboviral illnesses, confirmed or suspected, to the Division of Public Health Services (DPHS) within 24 hours at 603-271-4496 (after hours 1-800-852-3345, x5300).

### Background

Arboviruses in NH include West Nile virus (WNV) and Eastern Equine Encephalitis (EEE) virus, both transmitted to humans through the bite of an infected mosquito. In 2013, the first human case of locally acquired Jamestown Canyon virus (JCV) was identified in NH. EEE and WNV are maintained in a bird-mosquito cycle with humans considered incidental hosts. JCV is maintained in a deer-mosquito cycle, and reports of human illness are rare. The greatest risk for human acquisition of arboviral diseases is between July and October. Year-round transmission is possible in some geographic locations in the US.

Nationally last season (2014), there were 2,122 human cases of WNV reported in the US, including 85 deaths. Neuroinvasive Disease (meningitis and/or encephalitis) was recorded in 1,283 cases, while 839 cases were diagnosed with milder West Nile fever. There were 8 human cases of EEE reported in the US.

In NH during the 2014 season, there was one WNV-positive mosquito batch. There were no veterinary or human cases of WNV reported. There were 18 EEE positive mosquito batches and three EEE positive animals. Three human cases of EEE were reported.

In 2014, the first local transmission of chikungunya virus (CHIKV) was identified in the US, and the first travel-associated cases were reported in NH. There were 2,481 travel associated cases reported in the US and 11 locally-transmitted cases (all were acquired in Florida). There were 22 travel-associated cases of CHIKV reported in NH. The mosquito species that are capable of transmitting CHIKV have not been identified in NH.

### When to Suspect Arboviral Illness

The incubation period following the bite of an infected mosquito ranges from 3 to 14 days. Most arboviral infections are mild and non-apparent. Mild forms of disease normally present as a febrile illness but sudden onset of symptoms can occur and may include headache, myalgias and arthralgias. Approximately 20% of those infected with WNV develop a mild illness known as West Nile Fever.

The more severe forms of arboviral infection include altered mental status and/or neurological dysfunction (cranial and peripheral neuritis or other neuropathies, including acute flaccid paralysis syndrome). A minority of patients with severe disease develop a diffuse maculopapular or morbilliform rash. Approximately 1 in 150 WNV infections will result in severe neurological disease with encephalitis more common than meningitis. Older patients are at increased risk of developing severe West Nile Virus infections. For EEE, approximately one-third of all people who develop clinical encephalitis will die from the disease. Among those who recover, many suffer from permanent brain damage. Severe disease can be seen in any age group, including children.

The typical laboratory findings are normal or elevated total leukocyte counts, lymphocytopenia and anemia, and hyponatremia in peripheral blood. Examination of cerebrospinal fluid (CSF) shows pleocytosis (usually with a predominance of lymphocytes), elevated protein, and normal glucose levels. For about one-third of WNV patients, magnetic resonance imaging (MRI) shows enhancement of the leptomeninges, the periventricular areas, or both, while MRI of EEE patients often reveal abnormalities of the basal ganglia and thalami.

Treatment is supportive, often involving hospitalization, intravenous fluids, respiratory support, and prevention of secondary infections for patients with severe disease.

### When to Report Suspected Cases of Arboviral Illness

Clinicians, hospitals, and laboratories should report within 24 hours any patient meeting the following criteria:

1. Any patient with encephalitis or meningitis from July through November, who meet criteria a, b and c below without an alternative diagnosis:
  - a. Fever  $\geq$  38.0 C or 100 F, and
  - b. CNS involvement including altered mental status (altered level of consciousness, confusion, agitation, lethargy) and/or other evidence of cortical involvement (e.g., focal neurologic findings, seizures), and
  - c. Abnormal CSF profile suggesting a viral etiology (a negative bacterial stain and culture) showing pleocytosis with predominance of lymphocytes. Elevated protein and normal glucose levels.

### How to Report Suspect Cases of Arboviral Illness

All suspected arboviral cases should first be reported to the New Hampshire Division of Public Health Services by telephone. A completed case report form (attached) must be faxed to the NH Infectious Disease Investigation Section (603-271-0545) *and* a copy submitted with the laboratory specimen(s) to the NH Public Health Laboratories (PHL). DPHS staff members are available 24/7 to help determine if the clinical presentation meets the case criteria and whether further testing would be appropriate. Specimen submission guidelines are attached.

For additional information on arboviral illness and maps of recent activity, please visit the NH DHHS website at <http://www.dhhs.nh.gov/dphs/cdcs/arboviral/results.htm>. For fact sheets on WNV and EEE, go to <http://www.dhhs.nh.gov/dphs/cdcs/arboviral/publications.htm>

### Laboratory Testing for Arboviral Illnesses

Due to a national reagent shortage, some clinical and reference laboratories (LabCorp, ARUP, Mayo) have been forced to stop arbovirus antibody testing for EEE, Western equine encephalitis (WEE), California encephalitis, and Saint Louis encephalitis (SLE).

The NH PHL performs a microsphere immunoassay test developed by Centers for Disease Control and Prevention (CDC), which is not affected by the reagent shortage. The NH PHL's arbovirus panel includes IgM antibody testing for EEE, SLE, and WNV.

Chikungunya virus real time reverse transcriptase polymerase chain reaction (RT-PCR) testing is available at the NH PHL. If the Bureau of Infectious Disease Control is consulted and testing for this condition is appropriate, there will be no charge for this test. Without prior consultation, the cost for the RT-PCR test is \$72.00. Serum specimens should be taken during the acute phase of the disease ( $\leq 7$  days post symptom onset). If the patient makes the first visit to the physician after day 7 of symptoms, the sample is unlikely to yield positive PCR results. Serum collected  $>8$  days post onset will be sent to CDC for antibody testing. Serum should be stored, refrigerated and shipped to the PHL as soon as possible.

### For additional information on WNV and EEE please refer to:

1. The NH DHHS website at: <http://www.dhhs.nh.gov/dphs/cdcs/arboviral/index.htm>
2. The Centers for Disease Control, Division of Vector-Borne Infectious Diseases website at: <http://www.cdc.gov/ncidod/dvbid/westnile/clinicians/>.

If you or other health care providers have questions, please call Bureau of Infectious Disease Control at (603) 271-4496 or 1-800-852-3345, extension 4496 during business hours (8 am to 4:30 pm). Nights or weekends call the New Hampshire Hospital switchboard at 1-800-852-3345 extension 5300 and request the Public Health Professional on-call.

For any questions regarding the contents of this message, please contact NH DHHS, DPHS, Bureau of Infectious Disease Control at 603-271-4496 (after hours 1-800-852-3345 ext.5300).

To change your contact information in the NH Health Alert Network please call 603-271-4596 or email [health.alert@nh.gov](mailto:health.alert@nh.gov).

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Practitioners, Infectious Disease Specialists, Community Health Centers, Hospital CEOs, Hospital Emergency Departments, Nurses, NHHA, Pharmacists, Laboratory Response Network, Manchester Health Department, Nashua Health Department, Public Health Networks, DHHS Outbreak Team, DPHS Investigation Team, DPHS Management Team, Northeast State Epidemiologists, Zoonotic Alert Team, Health Officers, Deputy Health Officers, MRC, NH Schools, EWIDS

From: Elizabeth A. Talbot, MD – Deputy State Epidemiologist  
Originating Agency: NH Department of Health and Human Services, Division of Public Health Services

**Attachments:**

- 1) NH Arboviral Case Report Form
- 2) Laboratory Submission Guidelines for Arboviral Testing

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**New Hampshire Case Report  
Arboviral Infection  
Encephalitis/Meningitis**

**This form must be faxed to the New Hampshire Communicable Disease Control Section (603-271-0545) and a copy submitted with the laboratory specimen(s) to the NH Public Health Laboratories**

**Prior to submission of suspect chikungunya virus specimens for testing, a Public Health Nurse at the Bureau of Infectious Disease Control must be consulted in order to avoid a testing fee.**

Please indicate the nurse contacted for tracking purposes: \_\_\_\_\_

**PATIENT INFORMATION**

Name: \_\_\_\_\_ Date of Birth: \_\_\_\_/\_\_\_\_/\_\_\_\_  Male  Female  
Last First MI mm dd yy

Home Address: \_\_\_\_\_ Homeless  Yes  No  
Street City State Zip

Phone (H) \_\_\_\_\_ (W) \_\_\_\_\_ (Cell) \_\_\_\_\_

RACE  White  Black/African American  Asian  Native Hawaiian/Pacific Islander  American Indian/Alaska Native  Unknown  
 ETHNICITY  Unknown  Hispanic  Non-Hispanic

**CLINICAL INFORMATION**

Current Diagnosis:  Encephalitis  Meningitis  Other \_\_\_\_\_

Hospitalized?  Yes  No If yes, Hospital: \_\_\_\_\_

Date of Admission: \_\_\_\_/\_\_\_\_/\_\_\_\_ Date of Discharge/Transfer: \_\_\_\_/\_\_\_\_/\_\_\_\_

Physician/Provider: \_\_\_\_\_ Phone: \_\_\_\_\_

**SYMPTOMS:** Date of first symptoms \_\_\_\_/\_\_\_\_/\_\_\_\_ Date of first *neurologic* symptoms \_\_\_\_/\_\_\_\_/\_\_\_\_

	YES	NO	UNK		YES	NO	UNK		YES	NO	UNK
Fever $\geq 100^{\circ}\text{F}$	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Disorientation	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Convulsions	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Highest Temp (if known) _____ <sup>o</sup> F				Delirium	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Paralysis/Paresis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Headache	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Lethargy	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Acute Flaccid Paralysis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Stiff Neck	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Stupor	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Cranial Nerve Palsy	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Tremor	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Coma	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Rash	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Vomiting/Nausea	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Muscle Weakness	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Location of Rash			
Diarrhea	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Hyperreflexia	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Hemorrhage	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Confusion	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Muscle Pain	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Joint Pain	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Seizures	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Rigidity	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>				
Other _____											

OUTCOME  Recovered  Residual Symptoms  Died  Unknown If patient died, date of death \_\_\_\_/\_\_\_\_/\_\_\_\_

**LABORATORY INFORMATION/TEST RESULTS (attach laboratory sheets)**

Acute specimens (serum or CSF) must be collected within 3 to 10 days after onset of symptoms. Convalescent specimens should be collected 2-3 weeks after acute sample. If CSF is collected and submitted, please include serum sample.

CSF (specify units) Date \_\_\_\_/\_\_\_\_/\_\_\_\_ Abnormal?  Yes  No  Unknown Glu \_\_\_\_\_ Prot \_\_\_\_\_ RBC \_\_\_\_\_  
 WBC \_\_\_\_\_ Diff. Segs% \_\_\_\_\_ Lymphs% \_\_\_\_\_ Gram stain \_\_\_\_\_ Bacterial Culture \_\_\_\_\_  
 Fungal/Parasitic tests \_\_\_\_\_ Viral test results (Culture/Serology/PCR) \_\_\_\_\_  
 CBC (specify units) Date \_\_\_\_/\_\_\_\_/\_\_\_\_ WBC \_\_\_\_\_ Diff.Segs% \_\_\_\_\_ Lymphs% \_\_\_\_\_  
 MRI Date \_\_\_\_/\_\_\_\_/\_\_\_\_ Result \_\_\_\_\_  
 CT Date \_\_\_\_/\_\_\_\_/\_\_\_\_ Result \_\_\_\_\_  
 EMG Date \_\_\_\_/\_\_\_\_/\_\_\_\_ Result \_\_\_\_\_

**ANTIVIRAL TREATMENT**  Yes  No  Unk If Yes, list below. **Date Started** \_\_\_\_/\_\_\_\_/\_\_\_\_

**RISK FACTOR INFORMATION FOR PRELIMINARY OR CONFIRMED POSITIVE CASES OF ARBOVIRAL ILLNESS**

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_/\_\_\_\_/\_\_\_\_

1. Does the patient's residence have screened windows? Yes No Unknown
2. During the two weeks before onset of illness does the patient recall being bitten by mosquitoes?  
Yes No If yes, dates and places \_\_\_\_\_
3. Is the patient a smoker? Yes No Unknown  
If yes, do they smoke outdoors? Yes No Unknown
4. On average, how much time has the patient spent outdoors each day in the two weeks prior to onset? \_\_\_\_\_  
List any unusually long periods spent outside during the two weeks prior to onset: \_\_\_\_\_
5. Does the patient use any prevention measures to avoid mosquito bites? Yes No Unknown  
If yes, list \_\_\_\_\_  
Does the patient use mosquito repellent when outdoors: Always Sometimes Rarely Never  
Does the repellent contain DEET (N, N-diethyl-meta-toluamide, or N, Ndiethyl-3-methylbenzamide), Picaridin, or Oil of Lemon Eucalyptus? Yes No Unknown
6. During the two weeks before onset did the patient travel outside the county of residence?  
Yes No Unknown If yes, specify when and where: \_\_\_\_\_
7. Has the patient traveled outside of New Hampshire in the two weeks prior to onset? Yes No Unknown  
If yes, specify when and where: \_\_\_\_\_
8. Has the patient traveled outside the U.S. in the two weeks prior to onset? Yes No Unknown  
If yes, specify when and where: \_\_\_\_\_
9. Does the patient have any underlying medical conditions? Yes No Unknown  
If yes, specify: \_\_\_\_\_
10. What is the patient's occupation? \_\_\_\_\_

**BLOOD DONATION/TRANSFUSION/TRANSPLANT HISTORY/PREGNANCY**

11. Has the patient received an organ transplant or blood product transfusion in the month prior to onset?  
Yes No Unknown  
If yes, specify when and where: \_\_\_\_\_
12. Has patient donated blood products or been a living organ donor in the one month prior to onset? Yes No Unknown
13. Is the patient currently pregnant? Yes No Unknown Not applicable  
If yes, weeks pregnant \_\_\_\_\_ due date \_\_\_\_/\_\_\_\_/\_\_\_\_
14. Is the patient breastfeeding or planning to breastfeed? Yes No Unknown

**COMMENTS:** \_\_\_\_\_  
\_\_\_\_\_

**REPORTED BY:** \_\_\_\_\_ **DATE OF REPORT:** \_\_\_\_/\_\_\_\_/\_\_\_\_

Last Name \_\_\_\_\_ First Name \_\_\_\_\_ Title(ICN, Resident, Attending) \_\_\_\_\_

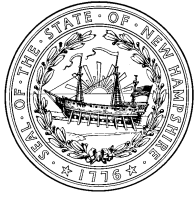
Work address \_\_\_\_\_ City \_\_\_\_\_ State \_\_\_\_\_ Zip Code \_\_\_\_\_

Phone \_\_\_\_\_ Pager \_\_\_\_\_

**FOR DHHS USE:**

Initial Report Taken by: \_\_\_\_\_ Report Completed by: \_\_\_\_\_

Case Status: Confirmed Probable Not a Case Unknown Other State



STATE OF NEW HAMPSHIRE

DEPARTMENT OF HEALTH AND HUMAN SERVICES

29 HAZEN DRIVE, CONCORD, NH 03301-6503
603-271-4612 1-800-852-3345 Ext. 4612
Fax: 603-271-4827 TDD Access: 1-800-735-2964



Nicholas A. Toumpas
Commissioner

Marcella Jordan Bobinsky
Acting Director

NH Public Health Laboratories

How to Collect and Submit Clinical Specimens for Arboviral Testing

All suspect arbovirus cases should be reported to the Communicable Disease Control Section at 1-800-852-3345, ext. 4496 or the Public Health Laboratories at (603) 271-4661 before specimens are submitted.

Diagnostic testing: The arboviral testing panel is a serological test for West Nile virus (WNV), Eastern Equine Encephalitis virus (EEE), St. Louis Encephalitis virus (SLE), and may include Powassan virus (depending on availability of reagents).

- The most efficient diagnostic method measures IgM antibodies in CSF or serum collected within 8 days of illness onset. The PHL uses the Microsphere Immunoassay (MIA) for detection of IgM antibody.
Since the MIA is a preliminary test, Plaque Reduction Neutralization test (PRNT) is required for case confirmation.
The IgM antibody does not cross the blood-brain barrier; IgM antibody in CSF strongly suggests central nervous system infection.
Serologic tests have a lower sensitivity due to cross-reactivity to related flaviviruses (e.g., yellow fever, Japanese encephalitis, dengue) and the persistence of WNV IgM antibodies in serum for 6 months or longer after infection.

Fee Schedule:

Table with 2 columns: TEST, CPT. Rows include Eastern Equine Encephalitis (EEE) virus antibodies, IgM (86652), St. Louis Encephalitis virus antibodies, IgM (86653), and West Nile Virus (WNV) antibodies, IgM (86788).

All specimens submitted to the Public Health Laboratories will be screened for EEE, SLE, and WNV.

The Total Cost Per Screen is \$105.00.

Note: All spinal fluid submissions must be accompanied by a corresponding serum sample. There will be only a single charge for the paired specimens.

Specimens:

Cerebrospinal fluid (CSF): As early as the first few days of illness, IgM antibody can be demonstrated in CSF by MIA.

Since other viruses can cause encephalitis, culture for additional viruses (other arboviruses, enteroviruses, and herpesviruses) may be performed at the discretion of the laboratory.

Submit 2-5 ml in sterile, empty, screw-capped container.

- **Serum:** Acute serum (3ml) should be collected and sent immediately to PHL for testing. Serum will be tested for IgM arboviral antibody. If specimen is IgM positive, then a convalescent specimen will be requested to determine the timing of infection.

**Ideal timing of specimens for serology:**

Specimen	Timing
Acute	3 to 10 days after onset of symptoms
Convalescent	2-3 weeks after acute sample

**All spinal fluid submissions must be accompanied by a corresponding serum sample.**

The following information is critical for accurate interpretation of test results and should be recorded on the accompanying case report form:

- **Date of onset of disease symptoms**
- Date of specimen collection
- Unusual immunological status of patient (e.g. immunosuppression)
- Brief clinical summary including suspected diagnosis (e.g., encephalitis or meningoencephalitis)
- Current address and travel history to flavivirus-endemic areas
- History of prior vaccination against flavivirus disease (e.g., yellow fever, Japanese encephalitis, or Central European encephalitis)
- Disease history (e.g., previous history of viral encephalitis or Dengue fever)

**Procedure for submission of serum or CSF:**

1. Perform lumbar puncture or venipuncture (SST or whole blood tube) by standard aseptic technique.
2. Label the specimen tubes with patient's full name and the date of collection.
3. If possible, centrifuge blood to separate serum.
4. For CSF, tightly seal cap and then wrap parafilm around seal to provide additional protection from leakage during transport.
5. Fill out requisition form completely, being sure to request "Arbovirus IgM"
6. Place CSF inside zip-lock biohazard bag and seal.
7. Place blood tube inside inner metal liner. Be sure there is enough absorbent material to cushion tubes in transit or to absorb liquid in case of leaking or broken tubes. Cap liner tightly.
8. Wrap the requisition form around the OUTSIDE of the inner metal liner.
9. Insert the metal liner into the outer cardboard container, and cap tightly. Make certain that the mailing container is labeled with the name and address of the NH PHL.
10. Mail first class or hand/courier deliver to the PHL. For emergency pickup after hours, contact the PHL at 1-800-852-3345. Refrigerate at 2-8° C if it is not possible to send specimen immediately.

The arboviral collection kit consists of:

- ❖ A labeled cardboard outer mailing container
- ❖ An aluminum inner liner
- ❖ An SST vacutainer blood collection tube
- ❖ A polypropylene tube and parafilm for transport of CSF
- ❖ Absorbent material
- ❖ Requisition form

**To order specimen collection kits, please call 271-4661, or 1-800-852-3345, extension 4661. For further technical information regarding diagnostic testing, please call Denise Bolton, at 271-3684, or 1-800-852-3345, extension 3684.**