



Dengue Case Investigation

- Dengue-like illness
- Dengue
- Severe Dengue

NBS Patient ID: _____

PLEASE PRINT LEGIBLY

Patient Information

Last Name: _____ First Name _____
 Date of Birth: ____/____/____ Sex: Male Female Unknown
 Street Address: _____ City, State, Zip: _____
 Patient Phone: _____ County of Residence: _____
 Race: Asian American Indian/Alaskan Native
 Black or African American Native Hawaiian/Pacific Islander
 White Unknown Other: _____
 Ethnicity: Hispanic Not Hispanic Unknown

Clinical Information

Physician: _____ Address _____
 City, State, Zip: _____ Phone: _____ Fax: _____
 Was the patient hospitalized for this illness? Yes No Unknown
 If yes, provide name of hospital: _____
 Dates of hospitalization: Admission ____/____/____ Discharge ____/____/____
 Date of Illness Onset: ____/____/____
 Is the patient deceased? Yes No Unknown
 If yes, provide date of death: _____ (submit documentation if due to arbovirus)

Clinical Evidence

Dengue-like illness (reported by patient or healthcare provider):

Fever Yes No Unknown

Dengue (fever PLUS one or more of the following):

- Headache Yes No Unknown
- Retro-orbital pain Yes No Unknown
- Nausea/Vomiting Yes No Unknown
- Abdominal pain Yes No Unknown
- Myalgia Yes No Unknown
- Joint/bone pain Yes No Unknown
- Rash Yes No Unknown
- Leukopenia (total white blood cell count <5,000mm³) Yes No Unknown
- Extravascular fluid accumulation Yes No Unknown
- Positive tourniquet test Yes No Unknown
- Petechiae Yes No Unknown
- Purpura/Ecchymosis Yes No Unknown
- Mucosal bleeding Yes No Unknown
- Liver enlargement > 2 cm Yes No Unknown
- Increasing hematocrit with thrombocytopenia Yes No Unknown

Severe Dengue (Dengue PLUS one or more of the following):

- Severe plasma leakage with respiratory distress Yes No Unknown
- Severe bleeding (i.e. melena, menorrhagia) Yes No Unknown
- Severe organ involvement Yes No Unknown
- Elevated liver transaminases (ALT or AST ≥ 1,000 U/L) Yes No Unknown
- Impaired consciousness Yes No Unknown

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Patient Name: _____

Epidemiology

Did the patient donate or receive blood, blood products, or organ/tissue in the last 30 days?

Yes No Unknown

If yes: Type of product: Blood Blood products Organ/tissue

Donation date(s): ____/____/____; ____/____/____; ____/____/____

Transfusion/transplant date(s): ____/____/____; ____/____/____; ____/____/____

Blood Collection Agency/Medical Facility: _____

Does this patient have a recent vaccination against a flavivirus (e.g. Yellow fever or Japanese encephalitis)?

Yes No Unknown

Was the patient pregnant during illness?

Yes No Unknown N/A

Was the patient breastfeeding within 2 weeks of onset?

Yes No Unknown N/A

Occupation: _____

(give exact job, type of business or industry, work shift and % of time spent outside while at work)

In the 30 days prior to onset, how many hours did the patient spend outdoors each day?

<2 2-4 5-8 >8 Unknown

When outdoors, what percentage of the time did the patient use mosquito repellent?

Always 75% 50% 25% Never Unknown

Did the patient travel outside of their residence County within 15 days of illness onset? Yes No Unknown

If yes, provide dates of travel and locations: _____

Is case thought to be imported?

Yes No Unknown

If yes, from where: _____

Is this a dengue-endemic area?

Yes No Unknown

Is there evidence of ongoing transmission with other flaviviruses?

Yes No Unknown

Does the patient know anyone else experiencing a similar illness?

Yes No Unknown

If yes, provide names and contact information on page 3.

Transmission Mode: Vector-borne In-Utero (transplacental) Perinatal Blood-borne

Indeterminate Other (explain): _____

Was the patient viremic while in Texas (during 7 days after onset)?

Yes No Unknown

If yes, provide dates and locations where the patient may have been bitten by mosquitoes on page 3.

Laboratory Findings

| Test (IgM, IgG, PCR, or PRNT) | Date Collected | Lab | Source | Result | Interpretation |
|-------------------------------|----------------|-----|--------|--------|---|
| | | | | | <input type="checkbox"/> Positive <input type="checkbox"/> Negative |
| | | | | | <input type="checkbox"/> Positive <input type="checkbox"/> Negative |
| | | | | | <input type="checkbox"/> Positive <input type="checkbox"/> Negative |
| | | | | | <input type="checkbox"/> Positive <input type="checkbox"/> Negative |

Comments or Other Pertinent Epidemiological Data *(Use page 3 if necessary):*

Date First Reported: ____/____/____ Investigation: Started ____/____/____ Completed ____/____/____

Reporting Facility: _____

Name of Investigator: _____ *(Please print clearly)*

Agency: _____ *(Please do not abbreviate)*

Phone: _____ E-Mail: _____

