



Government of The Punjab
Health Department, Lahore
In collaboration with
DEAG, Services Institute of Medical Sciences, Lahore.

NIC Number: _____

Serial No: _____

Dengue Case Report

PATIENT INFORMATION			
First Name	Middle Name	Last Name	Primary language <input type="checkbox"/> English <input type="checkbox"/> Other <input type="checkbox"/> Urdu <input type="checkbox"/> Punjabi
Father's Name	Husband Name	DOB (dd/mm/yyyy)	Age
House Number		Street	
Locality (like Shadman, Baghbanpura etc.)		City	
Home Telephone		Cellular phone	
E-mail Address		Phone number of relative / attendant	
Work Address			
Locality (like Shadman, Baghbanpura etc.)		City	Education <input type="checkbox"/> Uneducated <input type="checkbox"/> Under matric <input type="checkbox"/> Undergraduate <input type="checkbox"/> Others: _____
Gender <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Other (Please specify): _____			
Pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown		If yes, Est. Delivery Date (dd/mm/yyyy)	
CLINICAL INFORMATION			
Physician Name – Last Name		First Name	Telephone Number

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SIGNS AND SYMPTOMS

Symptomatic? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown			Onset Date (dd/mm/yyyy)				Date First Sought Medical Care (dd/mm/yyyy)				
Signs / Symptoms	Yes	No	Unk	If yes, specify as note		Signs / Symptoms	Yes	No	Unk	if yes, specify as noted	
Fever				Highest temperature (specify °F/°C)		Purpura / Ecchymosis					
Headache						Abdominal pan					
Eye pain						Sweats					
Muscle pain						Epistaxis					
Joint pain						Bleeding gums					
Nausea or vomiting						Hematuria					
Diarrhea						Vaginal bleeding					
Irritability in infants											
Chills						Hypotension				SystolicBP	
Cough										Diastolic BP	
Petechiae						Other symptoms (specify)				Pulse Pressure	

TRAVEL HISTORY

Did patient travel outside of his district in last 10 days?
 Yes No Unknown

Has the patient traveled outside Pakistan during the last 10 days period?
 Yes No Unknown

If yes or either of these questions, specify all locations and dates below.

TRAVEL HISTORY – DETAILS

Location (City, county, country)	Date travel started (dd/mm/yyyy)	Date travel ended (dd/mm/yyyy)

EXPOSURES / RISK FACTORS

Did patient recall any mosquito bites during the incubation period?

Yes No Unknown

Occurrence of confirmed cases of dengue fever in family or neighborhood

Yes No Unknown

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PAST MEDICAL HISTORY

Has the patient been previously diagnosed with dengue?

 Yes No Unknown

If yes, date of Diagnosis (dd/mm/yyyy)

Serotype (if known)

 1 2 3 4**PREVIOUS HOSPITAL VISIT**

Did patient visit emergency room for illness?

 Yes No Unknown

Was the patient hospitalized?

 Yes No Unknown

If yes, how many total hospital nights

Was patient placed in isolation ward?

 Yes No Unknown

if there were any ER or hospital stays related to this illness, specify details below

HOSPITALIZATION – DETAILS

Hospital Name 1	Hospital Address	Admission Date (dd/mm/yyyy)
	Telephone	Discharge / Transfer Date (dd/mm/yyyy)
		Discharge Diagnosis
Hospital Name 2	Hospital Address	Admission Date (dd/mm/yyyy)
	Telephone	Discharge / Transfer Date (dd/mm/yyyy)
		Discharge Diagnosis

LABORATORY INFORMATION**Hematology**

Hematology

 Yes No Unknown

Date Collected (dd/mm/yyyy)

WBC

HCT

Hb

Platelets

Serology

Specimen Type 1

Collection date

 Serum Other(Please specify): _____

Type of test

ELISA-IgM

ELISA-IgG

NS 1

PCR

Other (please specify): _____

Level

Level

Pos

Neg

Pos

Neg

Laboratory Name

Telephone Number

Specimen Type 2

Collection Date

 Serum Other(Please specify): _____

Type of test

ELISA-IgM

ELISA-IgG

NS 1

PCR

Other (please specify): _____

Level

Level

Pos

Neg

Pos

Neg

Laboratory Name

Telephone Number

OUTCOME

Outcome?

 Survived Died Unknown

If survived,

Alive on

_____(dd/mm/yyyy)

Date of death (dd/mm/yyyy)

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NOTES / REMARKS

REPORTING AGENCY

Reporting Officer	Union Council No. (Area of City)	Telephone Number	Date (dd/mm/yyyy)
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First reported by

 Clinician Laboratory Other (please specify): _____
DISEASE CASE CLASSIFICATION

Case classification (see case definition below)

 Confirmed Probable Suspected
CASE DEFINITION**DENGUE (2012)**

Dengue Fever

Dengue Hemorrhagic Fever

Dengue Shock Syndrome

CLINICAL DESCRIPTION**Dengue Fever**

Dengue fever (DF) is most commonly an acute febrile illness defined by the presence of fever and two or more of the following, retro-orbital or ocular pain, headache, rash, myalgia, arthralgia, leukopenia, or hemorrhagic manifestations (e.g., positive tourniquet test, petechiae; purpura/ecchymosis; epistaxis; gum bleeding; blood in vomitus, urine, or stool; or vaginal bleeding) but not meeting the case definition of dengue hemorrhagic fever. Anorexia, nausea, abdominal pain, and persistent vomiting may also occur but are not case-defining criteria for DF.

Dengue Hemorrhagic Fever

Dengue hemorrhagic fever (DHF) is characterized by evidence of **plasma leakage** as shown by hemoconcentration (an increase in hematocrit \geq 20% above average for age or a decrease in hematocrit \geq 20% of baseline following fluid replacement therapy), OR pleural effusion, or ascites or hypoproteinemia. **Plus** one of the following:

- Fever lasting from 2-7 days,
- Evidence of hemorrhagic manifestation or a positive tourniquet test,
- Thrombocytopenia (\leq 100,000 cells per mm³), AND

Dengue Shock Syndrome

Dengue shock syndrome (DSS) has all of criteria for DHF plus circulatory failure as evidenced by:

- Rapid and weak pulse and narrow pulse pressure (< 20mm Hg), OR
- Age-specific hypotension and cold, clammy skin, and restlessness.

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CASE DEFINITION (CONTINUED)**LABORATORY CRITERIA FOR DIAGNOSIS****Confirmatory:**

- Isolation of dengue virus from or demonstration of specific viral antigen or genomic sequences in tissue, blood, cerebrospinal fluid (CSF), or other body fluid by polymerase chain reaction (PCR) test, immunofluorescence or immunohistochemistry, OR
- Seroconversion from negative for dengue virus-specific serum Immunoglobulin M (IgM) antibody in an acute phase (≤ 5 days after symptom onset) specimen to positive for dengue-specific serum IgM antibodies in a convalescent-phase specimen collected ≥ 5 days after symptom onset, OR
- Demonstration of a ≥ 4 -fold rise in reciprocal Immunoglobulin G (IgG) antibody titer or Hemagglutination inhibition titer to dengue virus antigens in paired acute and convalescent serum samples, OR
- Demonstration of a ≥ 4 -fold rise in PRNT (plaque reduction neutralization test) end point titer (as expressed by the reciprocal of the last serum dilution showing a 90% reduction in plaque counts compared to the virus infected control) between dengue viruses and other flaviviruses tested in a convalescent serum sample, OR
- Virus-specific immunoglobulin M (IgM) antibodies demonstrated in CSF.

Presumptive/Probable:

- Dengue-specific IgM antibodies present in serum with a P/N ratio ≥ 2

Exposure:

- Travel to a dengue endemic country or presence at location with ongoing outbreak within previous two weeks of dengue-like illness, OR
- Association in time and place with a confirmed or probable dengue case.

Case Classification:

Suspected: A clinically compatible case of DF, DHF, or DSS that is epidemiologically linked to a confirmed case

Probable: A clinically compatible case of DF, DHF, or DSS with laboratory results indicative of presumptive infection

Confirmed: A clinically compatible case of DF, DHF, or DSS with confirmatory laboratory results

Comment:**Asymptomatic Blood or Tissue Donor**

Dengue virus - specific viral antigen or genomic sequences demonstrated in donated blood or organs during screening and confirmatory testing in the absence of symptoms in the donor.

Dengue viruses are members of the Flaviviridae and have sufficient antigenic similarity to yellow fever virus, Japanese encephalitis virus, and West Nile virus that previous infection or vaccination may raise cross-reactive serum antibodies. After a primary infection with a heterologous flavivirus, subsequent antibody testing by ELISA may produce false positive results for a different flavivirus. PRNT can often resolve cross-reactive serum antibodies in this situation and identify the infecting virus. However, high-titered cross-reactive antibody levels produced from multiple previous flavivirus infections cannot be resolved by PRNT. This demonstrates the complexity inherent in serological diagnosis and differentiation in populations living in regions where more than one flavivirus co-circulates.

Reference testing is available from DEAG, Services Institute of Medical Sciences, Jail Road, Lahore, Pakistan

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