



Name : MR SATYAM NAUTIYAL  
Age : 21 Gender : MALE  
Ref. By Dr : SELF

### Laboratory Report.

Sr. Number : 12  
Invoice Date : 02-03-2025 05:08 PM  
Invoice Number : 307  
Registration No.: 307  
Report On : 02-03-2025 05:53 PM



## HAEMATOLOGY

### Complete Blood Cell Count

Test Name	Observed Values	Reference Intervals	Units
WBC	11.80	4--10	10 <sup>3</sup> /UL
LYM%	7.30	20--40	%
MID% /MXD%	4.60	3--10	%
GRANS% / NEUT%	88.10	50--70	%
LYM#	0.8	0.8--7.0	10 <sup>3</sup> /UL
MID# /MXD#	0.5	0.1--1.2	10 <sup>3</sup> /UL
GRANS# / NEUT#	10.5	2--8.0	10 <sup>3</sup> /UL
RBC	5.93	3.5--5.5	10 <sup>12</sup> /L
HGB	16.2	11--17	g/dL
HCT	51.9	36--48	%
MCV	87.6	73--87	fL
MCH	27.3	26--32	pg
MCHC	31.2	32--36	g/dL
RDW-SD	35.9	37--45	fL
RDW-CV	14.9	11.5--14.5	%
PLT	210	150--450	10 <sup>3</sup> /UL



TECHNOLOGIST

RADIOLOGIST

PATHOLOGIST

Facilities : X-ray, E.C.G., Blood, Urine, Stool, Semen, Pregnancy Test, Culture Etc.  
Note : The Above Result Are Subject To Variation Due To Technical Limitation Hence  
Correlation With Clinical Finding And Other Investigation Should Be Done.

THIS REPORT IS NOT VALID FOR MEDICO-LEGAL PURPOSE.



Name : MR SATYAM NAUTIYAL  
Age : 21 Gender : MALE  
Ref. By Dr : SELF

### Laboratory Report.

Sr. Number : 12  
Invoice Date : 02-03-2025 05:08 PM  
Invoice Number : 307  
Registration No.: 307  
Report On : 02-03-2025 05:53 PM



## HAEMATOLOGY MALARIA ANTIGEN CARD TEST

Test Name	Observed Values	Reference Intervals	Units
Pf	Negative		
Pv	Negative		

ADVANTAGE MALARIA CARD is an immunoassay based on the "Sandwich principle". Colloidal gold is conjugated to P.f. specific monoclonal anti-HRP-2 antibody and monoclonal anti-pan specific pLDH (plasmodium lactate dehydrogenase) antibody. The test uses monoclonal anti-Pf HRP-2 antibody (test line F) & monoclonal anti-P.v specific pLDH antibody (test line V) immobilized on a nitrocellulose strip. The test sample is added to the device. On addition of assay buffer, the red blood cells get lysed. If the sample contains P.falciparum or P.vivax or both, the colloidal gold conjugate complexes the HRP-2 / P.vivax specific pLDH in the lysed sample. This complex migrates through the nitrocellulose strip by capillary action. When the complex meets the line of the corresponding immobilized antibody, the complex is trapped forming a purplish pink band which confirms a reactive test result. Absence of a coloured band in the test region indicates a negative test result. To serve as a procedural control an additional line of anti-mouse antibody has been immobilized on the strip as control.



TECHNOLOGIST

RADIOLOGIST

PATHOLOGIST

Facilities : X-ray, E.C.G., Blood, Urine, Stool, Semen, Pregnancy Test, Culture Etc.  
Note : The Above Result Are Subject To Variation Due To Technical Limitation Hence  
Correlation With Clinical Finding And Other Investigation Should Be Done.

THIS REPORT IS NOT VALID FOR MEDICO-LEGAL PURPOSE.



Name : MR SATYAM NAUTIYAL  
Age : 21 Gender : MALE  
Ref. By Dr : SELF

### Laboratory Report.

Sr. Number : 12  
Invoice Date : 02-03-2025 05:08 PM  
Invoice Number : 307  
Registration No.: 307  
Report On : 02-03-2025 05:53 PM



### SEROLOGY

Test Name	Observed Values	Reference Intervals	Units
CRP (C-reactive Protein Quant.)	32.06	0--6	mg/L

Method : **Fluorescence Immunoassay Technology**

Sample Type: **Serum / Plasma / Whole Blood**

#### Clinical Information :

The C-Reactive Protein (CRP) is Synthesized by the Liver in Response to Interleukin-6 and Well Known as One of the Classical Acute-Phase Reactants and as a Marker of Inflammation. CRP is the First Acute-Phase Protein to be Described and is An Exquisitely Sensitive Systemic Marker of Inflammation and Tissue Damage. The Acute-Phase Response Comprises the Nonspecific Physiological and Biochemical Responses of Endothermic animals to most forms of tissue damage, infection, inflammation, and malignant neoplasia. The serum CRP level may rise from a normal level of <5 mg/L to 500 mg/L During the Body's General, Non-Specific Response to Infectious and Other Acute Inflammatory Events. For Some Time, the Measurement of CRP Concentration has Been used as a Clinical Tool for Monitoring Autoimmune Diseases and Infectious Processes, Such as Rheumatoid Arthritis.



TECHNOLOGIST

RADIOLOGIST

PATHOLOGIST

Facilities : X-ray, E.C.G., Blood, Urine, Stool, Semen, Pregnancy Test, Culture Etc.  
Note : The Above Result Are Subject To Variation Due To Technical Limitation Hence  
Correlation With Clinical Finding And Other Investigation Should Be Done.

THIS REPORT IS NOT VALID FOR MEDICO-LEGAL PURPOSE.





Name : MR SATYAM NAUTIYAL  
Age : 21 Gender : MALE  
Ref. By Dr : SELF

### Laboratory Report.

Sr. Number : 12  
Invoice Date : 02-03-2025 05:08 PM  
Invoice Number : 307  
Registration No.: 307  
Report On : 02-03-2025 05:53 PM



## SEROLOGY

### WIDAL (Slide Method)

Test Name	Observed Values					Reference Intervals	Units
Titer Report	1/20	1/40	1/80	1/160	1/320		
S.Typhi. O.	+	+	-	-	-		
S.Typhi. H.	+	-	-	-	-		
S.Para. Typhi. AH.	-	-	-	-	-		
S.Para. Typhi. BH.	-	-	-	-	-		

### WIDAL TEST INTERPETATION

A titre of 1/120 for "O" is diagnostic of enteric fever to be confirmed by rising titre subsequently.

A titre of 1/120 for "O" with any one of the "H" is diagnostic of the type of enteric fever

A titre of 1/60 "O" could be normal titre but to rule out enteric fever test is to be repeated after 5 to 7 days

Only "H" titre even of 1/120 or above is not diagnostic of enteric fever.

NOTE (Salmonella blood culture and sensitivity)

To rule out any uncertainties arising out of widal test a salmonella blood culture (diagnosed in 24 hrs) and also sensitivity (diagnosed in 48 hours) in multi drug resistant typhoid can be done Which is highly sensitive and specific even in first week of infection )



TECHNOLOGIST

RADIOLOGIST

PATHOLOGIST

Facilities : X-ray, E.C.G., Blood, Urine, Stool, Semen, Pregnancy Test, Culture Etc.  
Note : The Above Result Are Subject To Variation Due To Technical Limitation Hence  
Correlation With Clinical Finding And Other Investigation Should Be Done.

THIS RERORT IS NOT VALID FOR MEDICO-LEGAL PURPOSE.



Name : MR SATYAM NAUTIYAL  
Age : 21 Gender : MALE  
Ref. By Dr : SELF

### Laboratory Report.

Sr. Number : 12  
Invoice Date : 02-03-2025 05:08 PM  
Invoice Number : 307  
Registration No.: 307  
Report On : 02-03-2025 05:53 PM



## IMMUNOLOGY

### DENGUE NSI IgG, IgM

Test Name	Observed Values	Reference Intervals	Units
IgG	Negative		
IgM	Negative		
NSI	Negative		

Detected samples should be confirmed by a supplemental assay enzyme-linked immunosorbent assay (elisa) test

#### INTRODUCTION

Dengue virus is a flavivirus found largely in areas of the tropic and sub-tropics. There are four distinct but antigenically related serotypes of dengue viruses, and transmission is by mosquito, principally Aedes aegypti Aedes albopictus.

The mosquito-borne dengue viruses (serotype 1-4) cause dengue fever, a severe flu-like illness.

IgM antibodies are not detectable until 5 -10 days in case of primary dengue infection and until 4-5 days of secondary infection after the onset of illness.

IgG appears after 14 days and persist for the life in case of primary infection and rise within 1-2 days after the onset of symptoms in secondary infection

Primary dengue virus infection is characterized by elevations in specific NS1 antigen levels 0 to 9 days after the onset of symptoms; this generally persist upto 15 days.

#### INTENDED USE

Dengue Day 1 Test is a rapid solid phase immuno-chromatographic test for the qualitative detection of Dengue NS1 antigen and differential detection of IgM and IgG antibodies to dengue virus in human serum/ plasma.

#### LIMITATION OF TEST

This rapid test device will indicate the presence of Dengue antibodies only & should not used as the sole criteria for the diagnosis of Dengue .Serological cross reactivity across the flavivirus group [Dengue, 1, 2, 3, 4, St Louis encephalitis West Nile virus, Japanese encephalitis & yellow fever virus] is common.

Positive result should be confirmed By Other methods [Elisa, P C R , cell Culture].

If result is negative & clinical symptoms persist , additional testing using other clinical methods is essential .A negative result does not at any time preclude the possibility of Dengue infection .

**Advise :-** Close follow up & if suspicion persist repeat after 2 days., & confirm by Elisa

<<< END OF REPORT >>>



TECHNOLOGIST

RADIOLOGIST

PATHOLOGIST

Facilities : X-ray, E.C.G., Blood, Urine, Stool, Semen, Pregnancy Test, Culture Etc.  
Note : The Above Result Are Subject To Variation Due To Technical Limitation Hence  
Correlation With Clinical Finding And Other Investigation Should Be Done.

THIS REPORT IS NOT VALID FOR MEDICO-LEGAL PURPOSE.