

**Erythrocyte Sedimentation Rate ESR** 

PR.

292/05, Tulsidas Marg, Basement Chowk, Lucknow-226 003

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CMO Reg. No. RMEE 2445133 NABL Reg. No. MC-2491 Certificate No. MIS-2023-0218

0 - 15

Visit No Patient Name : Mr.ANUBHAV MISHRA : CHA250040738

24.00

Age/Gender Registration ON : 15 Y/M : 06/Mar/2025 04:44PM Lab No Sample Collected ON : 10138033 : 06/Mar/2025 05:16PM Referred By : Dr.RDSO LUCKNOW Sample Received ON : 06/Mar/2025 05:28PM Refer Lab/Hosp : RDSO LUCKNOW Report Generated ON : 06/Mar/2025 06:38PM

DENGUE PROFILE,MF BY CARD,HCV-RNA QUANTITATIVE (EDTA PLASMA),HIV,HBSAg,WIDAL,LFT,KIDNEY FUNCTION TEST - I, DEC+ESR Doctor Advice :

Westergreen

Test Name	Result	Unit	Bio. Ref. Range	Method	
CBC+ESR (COMPLETE BLOOD COUNT)					

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Test Name	Result I	Unit	Bio. Ref. Range	Method
MF BY CARD				
MICROFILARIA ANTIBODY (MF)	Negative		NEGATIVE	by card
WIDAL				
Sample Type : Serum				
SALMONELLA TYPHI O	1/160			

SALMONELLA TYPHI H NOTE:

1/160 **POSITIVE** 







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Test Name	Result	Unit	Bio. Ref. Range	Method
DENGUE PROFILE				
Dengue ( NS1) Antigen	NON REACTI	VΕ	Non Reactive	(Rapid Card Test)
DENGUE IgG	NON REACTI	VΕ	Non Reactive	(Rapid Card Test)
DENGUE IgM	NON REACTI	VΕ	Non Reactive	(Rapid Card Test)

## **COMMENTS:**

- -Primary dengue virus infection is characterized by elevation of specific IgM levels 3 to 5 days after the onset of symptoms and persists for 30 to 60 days. IgG levels become elevated 10 to 14 days and remain detectable for many years.
- -During secondary infection, IgM levels generally rise more slowly than in primary infection while IgG levels rise rapidly from 1 to 2 days after the onset of symptoms.
- -The test detects all four subtypes, DEN1, DEN2, DEN3 & DEN4 of dengue virus.

## **LIMITATIONS:**

- -This is only a screening test and will only indicate the presence or absence of dengue antibodies in the specimen. All reactive samples should be confirmed by confirmatory tests.
- -The patient clinical history, symptomatology as well as serological data should be considered.
- -False positive results can be obtained due to cross-reaction with EBV, RA, Leptospira, malaria, Hepatitis A, Influenza A & B, Salmonella typhi etc.
- -Immuno-depressive treatments presumably after the immune response to infection, inducing negative results in dengue patients.

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Test Name Result Unit Bio. Ref. Range Method

HEPATITIS B SURFACE ANTIGEN (HBsAg)

Sample Type: Serum

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HEPATITIS B SURFACE ANTIGEN NON REACTIVE

<1 - Non Reactive

CMIA

>1 - Reactive

Note: This is only a Screening test. Confirmation of the res<mark>ult (Non Reactive/Reactive)should be done by performing a PCR based test.</mark>

## **COMMENTS:**

-HBsAg is the first serological marker after infection with Hepatitis B Virus appearing one to ten weeks after exposure and two to eight weeks before the onset of clinical symptoms. HBsAg persists during the acute phase and clears late in the convalescence phase. Failure to clear HBsAg within six months indicates a chronic HBsAg carrier state. HBsAg assays are used to identify the persons infected with HBV and to prevent transmission of the virus by blood and blood products as well as to monitor the status of infected individuals in combination with other hepatitis B serological markers.

-Borderline cases must be confirmed with confirmatory neutralizing assay

## LIMITATIONS:

- -Results should be used in conjunction with patient history and other hepatitis markers for diagnosis of acute and chronic infections.
- -Specimens from patients who have received preparations of mouse monoclonal antibodies for diagnosis or therapy may contain human anti-mouse antibodies (HAMA) which may produce anomalous values when tested with assay kits that employs mouse monoclonal antibodies.
- -Heterophilic antibodies in human serum can react with reagent immunoglobulins, interfering with in vitro immunoassays. Patients routinely exposed to animals or animal serum products can be prone to this interference and anomalous results may be observed.
- -Cross reactivity for specimens from individual with medical conditions (Pregnancy, HIV etc) has been observed -HBsAg mutations may result in a false negative result in some HBsAg assays.
- -If HBsAg results are inconsistent with clinical evidence, additional testing is suggested to confirm the result.



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**Test Name** Bio. Ref. Range Unit Result HIV

**HIV-SEROLOGY** < 1.0: NON REACTIVE NON REACTIVE >1.0: REACTIVE

Done by: Vitros ECI (Sandwich Assay)

Note:-Elisa test is a screening method for HIV.It is known to give false Positive & Negative result.

Hence confirmation: "Western Blot" method is advised.







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Test Name	Result	Unit	Bio. Ref. Range	Method	
CBC+ESR (COMPLETE BLOOD COUNT)					
Hb	13.7	g/dl	12 - 15	Non Cyanide	
R.B.C. COUNT	4.50	mil/cmm	3.8 - 4.8	Electrical	
				Impedence	
PCV	41.7	%	36 - 45	Pulse hieght	
				detection	
MCV	92.1	fL	80 - 96	calculated	
MCH	30.2	pg	27 - 33	Calculated	
MCHC	32.9	g/dL	30 - 36	Calculated	
RDW	13.3	%	11 - 15	RBC histogram	
				derivation	
RETIC	1.2 %	%	0.5 - 2.5	Microscopy	
TOTAL LEUCOCYTES COUNT	6290	/cmm	4000 - 10000	Flocytrometry	
DIFFERENTIAL LEUCOCYTE COUNT					
NEUTROPHIL	75	%	40 - 70	Flowcytrometry	
LYMPHOCYTE	22	%	20-40	Flowcytrometry	
EOSINOPHIL	1	%	1 - 6	Flowcytrometry	
MONOCYTE	2	%	2 - 10	Flowcytrometry	
BASOPHIL	0	%	00 - 01	Flowcytrometry	
PLATELET COUNT	98,000	/cmm	150000 - 450000	Elect Imped	
PLATELET COUNT (MANUAL)	117000	/cmm	150000 - 450000	Microscopy.	
Mentzer Index	20		N 1/		
Peripheral Blood Picture	GH				

Red blood cells are normocytic normochromic. WBC show relative neutrophilia. Platelets are mild reduced with gaint form. No immature cells or parasite seen





DR. SHADABKHAN



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Test Name	Result	Unit	Bio. Ref. Range	Method
LIVER FUNCTION TEST				
TOTAL BILIRUBIN	0.40	mg/dl	0.4 - 1.1	Diazonium Ion
CONJUGATED ( D. Bilirubin)	0.10	mg/dL	0.00-0.30	Diazotization
UNCONJUGATED (I.D. Bilirubin)	0.30	mg/dL	0.1 - 1.0	Calculated
ALK PHOS	199.00	U/L	82 - 331	PNPP, AMP Buffer
SGPT	100.0	U/L	5 - 40	UV without P5P
SGOT	124.0	U/L	5 - 40	UV without P5P
KIDNEY FUNCTION TEST - I				
Sample Type : Serum				
BLOOD UREA	1 <mark>7.7</mark> 0	mg/dl	15 - 45	Urease, UV, Serum
CREATININE	0.60	mg/dl	0.50 - 1.40	Alkaline picrate- kinetic
SODIUM Serum	<mark>135.0</mark>	MEq/L	135 - 155	ISE Direct
POTASSIUM Serum	4.2	MEq/L	3.5 - 5.5	ISE Direct

\*\*\* End Of Report \*\*\*

CHARAK



