

PR.

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

Phone: 0522-4062223, 9305548277, 8400888844 9415577933, 9336154100, Tollfree No.: 8688360360

E-mail: charak1984@gmail.com

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

Visit No Patient Name : Ms.RENU SEHTA : CHA250039148

Age/Gender Registration ON : 54 Y/F : 04/Mar/2025 04:36PM Lab No Sample Collected ON : 10136443 : 04/Mar/2025 04:38PM Referred By : Dr.VISHAL SINGH NEGI Sample Received ON : 04/Mar/2025 04:39PM

Refer Lab/Hosp Report Generated ON : CGHS (DEBIT) : 04/Mar/2025 06:01PM

HCV,HBSAg,HIV,PT/PC/INR,RANDOM,CALCIUM,KIDNEY FUNCTION TEST - I,LFT,ECG,CBC+ESR Doctor Advice :



Test Name	Result	Unit	Bio. Ref. Range	Method	
CBC+ESR (COMPLETE BLOOD COUNT)					
Erythrocyte Sedimentation Rate ESR	30.00		0 - 20	Westergreen	







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Test Name	Result	Unit	Bio. Ref. Range	Method
SERUM CALCIUM				
CALCIUM	10.5	mg/dl	8.8 - 10.2	dapta / arsenazo III
PT/PC/INR				
PROTHROMBIN TIME	13 Second		13 Second	Clotting Assay
Protrhromin concentration	100 %		100 %	
INR (International Normalized Ratio	1.00		1.0	







**PATHOLOGIST** 



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Test Name	Result	Unit	Bio. Ref. Range	Method
HEPATITIS B SURFACE ANTIGEN (HBsAg)				
Sample Type : SERUM				

HEPATITIS B SURFACE ANTIGEN NON REACTIVE <1 - Non Reactive **CMIA** >1 - Reactive

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

#### 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
- -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	Result	Unit	Bio. Ref. Range	Method
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.

## **HEPATITIS C VIRUS (HCV) ANTIBODIES**

HEPATITIS C VIRUS (HCV) ANTIBODIES NON REACTIVE

Non Reactive

## (TRIO DOT ASSAY)

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

CHARAK





P.R.

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HCV,HBSAg,HIV,PT/PC/INR,RANDOM,CALCIUM,KIDNEY FUNCTION TEST - I,LFT,ECG,CBC+ESR Doctor Advice :

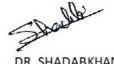
|--|--|--|--|

Test Name	Result	Unit	Bio. Ref. Range	Method
CBC+ESR (COMPLETE BLOOD COUNT)				
Hb	12.8	g/dl	12 - 15	Non Cyanide
R.B.C. COUNT	4.40	mil/cmm	3.8 - 4.8	Electrical
				Impedence
PCV	39.8	%	36 - 45	Pulse hieght
				detection
MCV	90.9	fL	80 - 96	calculated
MCH	29.2	pg	27 - 33	Calculated
MCHC	32.2	g/dL	30 - 36	Calculated
RDW	12	%	11 - 15	RBC histogram
				derivation
RETIC	1.0 %	%	0.5 - 2.5	Microscopy
TOTAL LEUCOCYTES COUNT	7370	/cmm	4000 - 10000	Flocytrometry
DIFFERENTIAL LEUCOCYTE COUNT				
NEUTROPHIL	64	%	40 - 75	Flowcytrometry
LYMPHOCYTE	29	%	20-40	Flowcytrometry
EOSINOPHIL	4	%	1 - 6	Flowcytrometry
MONOCYTE	3	%	2 - 10	Flowcytrometry
BASOPHIL	0	%	00 - 01	Flowcytrometry
PLATELET COUNT	297,000	/cmm	150000 - 450000	Elect Imped
PLATELET COUNT (MANUAL)	297000	/cmm	150000 - 450000	Microscopy.
Mentzer Index	21			
Peripheral Blood Picture	CH			

Red blood cells are normocytic normochromic. Platelets are adequate. No immature cells or parasite seen.









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: 04/Mar/2025 05:24PM

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Test Name	Result	Unit	Bio. Ref. Range	Method		
BLOOD SUGAR RANDOM						
BLOOD SUGAR RANDOM	100.2	mg/dl	70 - 170	Hexokinase		
LIVER FUNCTION TEST						
TOTAL BILIRUBIN	0.62	mg/dl	0.4 - 1.1	Diazonium Ion		
CONJUGATED ( D. Bilirubin)	0.12	mg/dL	0.00-0.30	Diazotization		
UNCONJUGATED (I.D. Bilirubin)	0.50	mg/dL	0.1 - 1.0	Calculated		
ALK PHOS	98.60	U/L	30 - 120	PNPP, AMP Buffer		
SGPT	28.0	U/L	5 - 40	UV without P5P		
SGOT	31.0	U/L	5 - 40	UV without P5P		
KIDNEY FUNCTION TEST - I						
Sample Type : SERUM						
BLOOD UREA	21.70	mg/dl	15 - 45	Urease, UV, Serum		
CREATININE	0.60	mg/dl	0.50 - 1.40	Alkaline picrate- kinetic		
SODIUM Serum	135.0	MEq/L	135 - 155	ISE Direct		
POTASSIUM Serum	3.8	MEq/L	3.5 - 5.5	ISE Direct		

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

CHARAK





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Report Generated ON : 04/Mar/2025 04:56PM

# **ECG-REPORT**

RATE : 80 bpm

\* RHYTHM : Normal

\* P wave : Normal

\* PR interval : Normal

\* QRS Axis : Normal

Duration : Normal

Configuration : Normal

\* ST-T Changes : None

\* QT interval :

\* QTc interval : Sec.

\* Other :

**OPINION:** ECG WITH IN NORMAL LIMITS

(FINDING TO BE CORRELATED CLINICALLY )

[DR. RAJIV RASTOGI, MD, DM]

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

