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

Patient Name : Ms. NASHRA Visit No : CHA250044103

Age/Gender : 18 Y/F Registration ON : 11/Mar/2025 06: 22PM Lab No : 10141398 Sample Collected ON : 11/Mar/2025 06: 23PM Referred By : Dr.UZMA MUBASHSHIR Sample Received ON : 11/Mar/2025 06:23PM Refer Lab/Hosp : CHARAK NA Report Generated ON : 11/Mar/2025 08:16PM

Doctor Advice : PLAT COUNT, HB, URINE COM. EXMAMINATION, PT/PC/INR, BTCT, BLOOD GROUP, HCV, HBSAg, HIV, TSH, RANDOM

Test Name	Result	Unit	Bio. Ref. Range	Method

**BLOOD GROUP** 

P.R.

Blood Group "O" Rh (Anti -D) POSITIVE

PT/PC/INR		A contract of the contract of	
PROTHROMBIN TIME	13 Second	13 Second	Clotting Assay
Protrhromin concentration	100 %	100 %	
INR (International Normalized Ratio	) 1.00	1.0	





3 hade



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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.



**PATHOLOGIST** 



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Test Name	Result	Unit	Bio. Ref. Range	Method
HIV				

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

HCV

NON REACTIVE Anti-Hepatitis C Virus Antibodies. < 1.0: NON REACTIVE Sandwich Assay

> 1.0 : REACTIVE

Done by: Vitros ECI (Sandwich Assay)

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

URINE EXAMINATION REPORT				
Colour-U	YELLOW		Light Yellow	
Appearance (Urine)	CLEAR		Clear	
Specific Gravity	1.020		1.005 - 1.025	
pH-Urine	Acidic (6.0)		4.5 - 8.0	
PROTEIN	Absent	mg/dl	ABSENT	Dipstick
Glucose	Absent			
Ketones	Absent		Absent	
Bilirubin-U	Absent		Absent	
Blood-U	Absent	71 77 7	Absent	
Urobilinogen-U	0.20	EU/dL	0.2 - 1.0	
Leukocytes-U	Absent		Absent	
NITRITE	Absent		Absent	
MICROSCOPIC EXAMINATION				
Pus cells / hpf	Occasional	/hpf	< 5/hpf	
Epithelial Cells	4-6	/hpf	0 - 5	
RBC / hpf	Nil		< 3/hpf	





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Test Name	Result	Unit	Bio. Ref. Range	Method
HAEMOGLOBIN				
Hb	12.8	g/dl	12 - 15	Non Cyanide

## Comment:

Hemoglobin screening helps to diagnose conditions that affect RBCs such as anemia or polycythemia.

PLATELET COUNT					
PLATELET COUNT		231,000	/cmm	150000 - 450000	Elect Imped
PLATELET COUNT (MANUAL	_)	231000	/cmm	150000 - 450000	Microscopy.
BLOOD SUGAR RANDOM					
BLOOD SUGAR RANDOM		81.5	mg/dl	70 - 170	Hexokinase
	la de la companya de	7 /			
TSH					
TSH		3.78	ulU/ml	0.7 - 6.4	ECLIA

## Note

- (1) Patients having low T3 & T4 levels but high TSH levels suffer from primary hypothyroidism, cretinism, juvenile mysedema or autoimmune disorders
- (2) Patients having low T3 & T4 levels but high TSH levels suffer from grave~s disease, toxic adenoma or sub-acute thyroiditis.
- (3) Patients having either low or normal T3 & T4 levels but low TSH values suffer from iodine deficiency or secondary hypothyroidism.
- (4) Patients having high T3 & T4 levels but normal TSH levels may suffer from toxic multinodular goitre. This condition is mostly asymptomatic and may cause transient hyperthyroidism but no persistent symptoms.
- (5) Patient with high or normal T3 & T4 levels and low or normal TSH levels suffer either from T3 toxicosis or T4 Toxicosis respectively.
- (6) In patients with non thyroidal illness abnormal test results are not necessarily indicative of thyroidism but may be due to adaptation to the cacabolic state and may revert tonormal when the patient recovers.
- (7) There are many drugs for eg.Glucocorticoids ,dopamine,Lithium,iodides ,oral radiographic dyes,ets.Which may affect the thyroid function tests.
- (8) Generally when total T3& T4 results are indecisive then Free T3 & Free T4 test are recommended for further confirmation along with
- (1 Beckman DxI-600 2. ELECTRO-CHEMILUMINISCENCE TECHINIQUE BY ELECSYSYS -E411)

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





DR SHADARKHAN