

Patient Name : Mr.ASHAD	Visit No : CHA250043958
Age/Gender : 25 Y/M	Registration ON : 11/Mar/2025 02: 18PM
<b>Lab No : 10141253</b>	Sample Collected ON : 11/Mar/2025 02: 20PM
Referred By : Dr.MANISH TANDON	Sample Received ON : 11/Mar/2025 02: 41PM
Refer Lab/Hosp : CHARAK NA	Report Generated ON : 11/Mar/2025 04: 40PM
Doctor Advice : T3T4TSH,FOLIC ACID,VIT B12,HCV,HBSAg,HIV,CHEST PA,USG WHOLE ABDOMEN	



Test Name	Result	Unit	Bio. Ref. Range	Method
<b>VITAMIN B12</b>				
VITAMIN B12	124.0	pg/mL	180 - 814 Normal 145 - 180 Intermediate 145.0 Deficient pg/ml	CLIA

**Summary :-**

Nutritional & macrocytic anemias can be caused by a deficiency of vitamin B12. This deficiency can result from diets devoid of meat & bacterial products, from alcoholism or from structural / functional damage to digestive or absorptive processes. Malabsorption is the major cause of this deficiency.

<b>FOLIC ACID</b>				
FOLIC ACID	8.02	ng/ml	3.89 26.8	CMIA

Method: Electrochemiluminescence

COMMENTS: Folate deficiency causes megaloblastic anemia and eventually leukopenia and thrombocytopenia. Folic acid is believed to play a role in birth defects such as spina bifida, anencephaly, and oro-facial clefts as well as in inducing cardiovascular morbidity and mortality. Symptoms of deficiency take about 3 months to appear and can be caused by inadequate intake, increased body demand or folate antagonism by drugs. For diagnostics purposes, the folate findings should always be assessed in conjunction with the patient's medical history, clinical examination and other findings. This deficiency can result from diets devoid of raw fruits, vegetables or other foods rich in folic acid, as may be the case with chronic alcoholics, drug addicts, the elderly or persons of low socioeconomic status, etc. In addition, low serum also occurs during pregnancy. Folate assays are affected by hemolysis within the specimen.

CHARAK

[Checked By]



Print.Date/Time: 11-03-2025 17:39:57

\*Patient Identity Has Not Been Verified. Not For Medicolegal

DR. NISHANT SHARMA DR. SHADAB DR. ADITI D AGARWAL  
PATHOLOGIST PATHOLOGIST PATHOLOGIST

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

HEPATITIS B SURFACE ANTIGEN	NON REACTIVE		<1 - Non Reactive >1 - Reactive	CMIA
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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 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	Result	Unit	Bio. Ref. Range	Method
<b>HIV</b>				
HIV-SEROLOGY	NON REACTIVE		<1.0 : NON REACTIVE >1.0 : REACTIVE	

Done by: Vitros ECI ( Sandwich Assay)

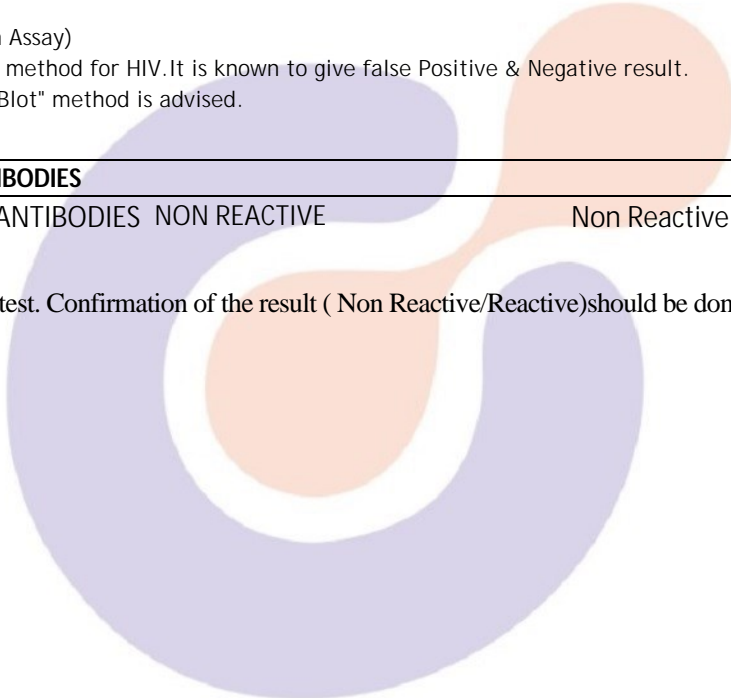
Note:-Elisa test is a screening method for HIV.It is known to give false Positive & Negative result.  
Hence confirmation:"Western Blot" method is advised.

**HEPATITIS C VIRUS (HCV) ANTIBODIES**

HEPATITIS C VIRUS (HCV) ANTIBODIES NON REACTIVE Non Reactive

(TRIO DOT ASSAY)

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*Aditi D Agarwal*

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Test Name	Result	Unit	Bio. Ref. Range	Method
<b>T3T4TSH</b>				
T3	1.99	nmol/L	1.49-2.96	ECLIA
T4	127.10	n mol/l	63 - 177	ECLIA
TSH	2.60	uIU/ml	0.47 - 4.52	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 \*\*\*



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*Signature*