

Regd. Office: Dr Lai PathLabs Ltd, Block-E, Sector-18, Rohini, New Delhi-110085 Web: www.laipathlabs.com, CIN: L74899DL1995PLC065388

Name : Ms. AKSHITA Lab No. : 179361202

Ref By SELF

Collected : 2/6/2024 11:38:00AM

A/c Status : P

Collected at : MAHESH NAGAR CC

Raja Park Chowk, Opp Easy Day, Jagadhari Road

Mb - 9896752154 Ambala Cantt Age : 24 Years Gender : Female

Reported : 2/6/2024 2:57:02PM Report Status : Final

Processed at: Dr. Lal PathLabs Ltd.

Ambala Haryana

Test Report

Test Name	Results	Units	Bio. Ref. Interval
VITAMIN D, 25 - HYDROXY, SERUM	107.10	nmol/L	75.00 - 250.00
(ECLIA)			

Interpretation

LEVEL	REFERENCE RANGE IN nmol/L	COMMENTS
Deficient	< 50 	High risk for developing bone disease
Insufficient 	 50-74 	Vitamin D concentration Which normalizes Parathyroid hormone concentration
Sufficient	75-250 	Optimal concentration for maximal health benefit
Potential intoxication	>250 	 High risk for toxic

Note

- The assay measures both D2 (Ergocalciferol) and D3 (Cholecalciferol) metabolites of vitamin D.
- 25 (OH)D is influenced by sunlight, latitude, skin pigmentation, sunscreen use and hepatic function.
- Optimal calcium absorption requires vitamin D 25 (OH) levels exceeding 75 nmol/L.
- It shows seasonal variation, with values being 40-50% lower in winter than in summer.
- · Levels vary with age and are increased in pregnancy.
- A new test Vitamin D, Ultrasensitive by LC-MS/MS is also available

Comments

Vitamin D promotes absorption of calcium and phosphorus and mineralization of bones and teeth. Deficiency in children causes Rickets and in adults leads to Osteomalacia. It can also lead to Hypocalcemia and Tetany. Vitamin D status is best determined by measurement of 25 hydroxy vitamin D, as it is the major circulating form and has longer half life (2-3 weeks) than 1,25 Dihydroxy vitamin D (5-8 hrs).

Decreased Levels

· Inadequate exposure to sunlight



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Test Report

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- Dietary deficiency
- Vitamin D malabsorption
- Severe Hepatocellular disease
- Drugs like Anticonvulsants
- Nephrotic syndrome

Increased levels

Vitamin D intoxication





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Test Report

Test Name	Results	Units	Bio. Ref. Interval
HEPATITIS B SURFACE ANTIGEN (HBsAg), RAPID	Non-Reactive		
SCREENING TEST, SERUM			
(Immunochromatography)			

Interpretation

RESULT	REMARKS
Reactive	Indicates presence of Hepatitis B Surface Antigen.
Non-Reactive	Indicates absence of Hepatitis B Surface Antigen.

^{*} All reactive results should be subjected to HBsAg Neutralization test which can be requested as Test Code S116.

Note

- 1. Reactive test result indicates presence of Hepatitis B Surface Antigen. It cannot differentiate between the stages of Hepatitis B viral infection.
- 2. Non-Reactive test result indicates absence of Hepatitis B Surface Antigen.
- 3. False positive results may be observed in presence of heterophilic antibodies in serum or after HBV vaccination for transient period of time.
- 4. False negative reaction may be due to processing of sample collected early in the course of disease or presence of mutant forms of HBsAg.
- 5. For monitoring HBsAg levels, HBsAg Quantitative assay is recommended.



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Test Report

Test Name	Results	Units	Bio. Ref. Interval
HEPATITIS C VIRUS (HCV), RAPID SCREENING	Non-Reactive		
TEST, SERUM			
(Immunochromatography)			

Interpretation

	RESULTS		REMARKS	ļ
ļ	Reactive		Indicates presence of antibodies to Hepatitis C virus	ļ
ļ	Non-Reactive		Indicates absence of antibodies to Hepatitis C virus	İ

^{*} It is recommended to confirm all reactive results with the HCV antibody confirmatory test (S314).*

Note

- Reactive test result indicates presence of Hepatitis C virus infection. Active infection to be confirmed by HCV RNA PCR test. It cannot differentiate between the stages of Hepatitis C viral infection nor used to monitor the efficacy of treatment.
- 2. Non-Reactive test result indicates Hepatitis C virus infection is unlikely.
- 3. False positive results may be observed in patients receiving mouse monoclonal antibodies, on heparin therapy, on biotin supplements for diagnosis or therapy or presence of heterophilic antibodies in serum
- 4. False negative reaction may be due to processing of sample collected early in the course of disease, Prozone phenomenon, Immunosuppression & Immuno-incompetence.
- 5. Test conducted on serum.

Uses

- To diagnose suspected HCV infection in risk group.
- Prenatal Screening of pregnant women and pre surgical/interventional procedures work up.



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Ambala Haryana

Test Report

Test Name		Results	Units	Bio. Ref. Interval
HIV 1 & 2 ANTIBODI (Immunochromatogra	ES SCREENING TEST, SERUM aphy)	Non-Reactive		
Final Result	: Negative		Negative	
REMARKS	INTERPRETATION			
Reactive	Indicates Presence of ant	:ibodies to HIV 1/2	virus	
Non-Reactive	Indicates absence of anti	bodies to HIV 1/2	virus	

It is advised to verify all positive results by conducting the supplemental HIV 1 and HIV2 antibody confirmation and differentiation, LIA (S315).

Note

- 1. Positive test result indicates antibody detected against HIV-1/2.
- 2. Negative test result indicates antibody is not detected against HIV- 1/2.
- 3. Indeterminate test result indicates antibody to HIV-1/2 have been detected in the sample by two of three
- 4. False positive results may be observed in Autoimmune diseases, Alcoholic hepatitis, Primary biliary cirrhosis, Leprosy, Multiple pregnancies, Rheumatoid factor, and due to presence of heterophile
- 5. False negative results may occur during the window period and during the end stage of the disease.

Recommendations

1. Post-test counseling available between 9 am to 5 pm at LPL laboratories.



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Dr.Samriti Gupta MBBS, MD (Pathology) Chief of Lab

-----End of report -----



IMPORTANT INSTRUCTIONS

•Test results released pertain to the specimen submitted. •All test results are dependent on the quality of the sample received by the Laboratory.
•Laboratory investigations are only a tool to facilitate in arriving at a diagnosis and should be clinically correlated by the Referring Physician. •Report delivery may be delayed due to unforeseen circumstances. Inconvenience is regretted. •Certain tests may require further testing at additional cost for derivation of exact value. Kindly submit request within 72 hours post reporting. •Test results may show interlaboratory variations. •The Courts/Forum at Delhi shall have exclusive jurisdiction in all disputes/claims concerning the test(s). • or results of test(s). •Test results are not valid for medico legal purposes. •This is computer generated medical diagnostic report that has been validated by Authorized Medical Practitioner/Doctor. •The report does not need physical signature.

(#) Sample drawn from outside source.

If Test results are alarming or unexpected, client is advised to contact the Customer Care immediately for possible remedial action.

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