

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

Name : Mr. DUMMY Lab No. : WM17SPF

Ref By : SELF Collected : 7/11/2023 11:08:00AM

A/c Status : P

Collected at : LPL-ROHINI (NATIONAL REFERENCE LAB)

National Reference laboratory, Block E, Sector

18, ROHINI DELHI 110085 Age : 25 Years Gender : Male

Reported : 17/1/2024 10:55:48AM

Report Status : Revised

Processed at : LPL-NATIONAL REFERENCE LAB

National Reference laboratory, Block E.

Sector 18, Rohini, New Delhi -110085

Test Report

| Test Name | Results | Units | o. Re interval |
|-------------------|---------|-------|----------------|
| SWASTHFIT SUPER 4 | | | |

| LIVER & KIDNEY PANEL, SERUM | | | |
|--|--------|----------------|----------------|
| Creatinine | 1.00 | mg/dL | 0.70 - 1.30 |
| (Modified Jaffe,Kinetic) | 1.00 | mg/dL | 0.70 - 1.30 |
| GFR Estimated | 107 | r_L/min/1.73m. | >59 |
| (CKD EPI Equation 2021) | | | |
| GFR Category (KDIGO Guideline 2012) | G1 | | |
| Urea (Urease UV) | 40.00 | mg/dL | 13.00 - 43.00 |
| Urea Nitrogen Blood (Calculated) | 18.68 | mg/dL | 6.00 - 20.00 |
| BUN/Creatinine Ratio (Calculated) | 19 | | |
| Uric Acid (Uricase) | | mg/dL | 3.50 - 7.20 |
| AST (SGOT) (IFCC without P5P) | 30.0 | U/L | 15.00 - 40.00 |
| ALT (SGPT) | 40.0 | U/L | 10.00 - 49.00 |
| (IFCC without P5P) GGTP | 50.0 | U/L | 0 - 73 |
| (IFCC) | 00.0 | 0/L | 0-75 |
| Alkaline Phosphatase (/ ∠P) | 100.00 | U/L | 30.00 - 120.00 |
| (IFCC-AMP) | | | |
| Bilirubin Total | 1.00 | mg/dL | 0.30 - 1.20 |
| (Oxidation) | | | |
| Pilitubin Direct (Oxidation) | 0.20 | mg/dL | <0.3 |
| Bilirut | 0.80 | mg/dL | <1.10 |
| rotal Protei | 8.00 | g/dL | 5.70 - 8.20 |
| (Bitret) | 0.00 | g/uL | 5.70 - 6.20 |
| Albunan | 4.00 | g/dL | 3.20 - 4.80 |
| (BCG) | | S | |
| A : G Ratio | 1.00 | | 0.90 - 2.00 |
| (Calculated) | | | |
| Globulin(Calculated) | 4.00 | gm/dL | 2.0 - 3.5 |
| Calcium, Total (Arsenazo III) | 9.00 | mg/dL | 8.70 - 10.40 |



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| Phosphorus (Molybdate UV) | 4.00 | mg/dL | 2 0 - 5.10 |
| Sodium (Indirect ISE) | 140.00 | mEq/L | 136.00 - 145.00 |
| Potassium (Indirect ISE) | 4.00 | mEq/L | 3.50 - 5.10 |
| Chloride (Indirect ISE) | 100.00 | m .q/L | 98.00 - 107.00 |

| LIPID SCREEN, SERUM | | | |
|--|--------|-------|---------|
| Cholesterol, Total (CHO-POD) | 100.00 | mg/dL | <200.00 |
| Triglycerides (GPO-POD) | 100,00 | mg/dL | <150.00 |
| HDL Cholesterol (Enz Immunoinhibition) | 2,000 | mg/dL | >40.00 |
| LDL Cholesterol, Calculated (Calculated) | 50.00 | mg/dL | <100.00 |
| VLDL Cholesterol,Calculated (Calculated) | 20.00 | mg/dL | <30.00 |
| Non-HDL Cholesterol (Calculated) | 70 | mg/dL | <130 |

Note

- 1. Measurements in the same patient can show physiological & analytical variations. Three serial samples 1 that apart are recommended for Total Cholesterol, Triglycerides, HDL& LDL Cholesterol.
- 2. Additional tests for Apolipoprotein B, hsCRP, Lp(a) & LP-PLA2 should be considered among tients with moderate risk for ASCVD for risk refinement.

Treatment Gols sper Lipid Association of India 2020

| ATEGOR | TREATMI | ENT GOAL | CONSIDER THERAPY | | | |
|-------------------------------------|-----------------------------------|--|------------------|--|--|--|
| CATEGOR | LDL CHOLESTEROL (LDL-C)(mg/dL) | NON HDL CHLOESTEROL (NON HDL-C) (mg/dL) | | NON HDL CHLOESTEROL (NON HDL-C) (mg/dL) | | |
| Extreme Risk Group Category A | <50 (Optional goal ≤30) | <80 (Optional goal ≤60) | ≥50 | ≥80 | | |
| Extreme Risk Group Category A | ≤30 | ≤60 | >30 | >60 | | |
| Very High | <50 | <80 | ≥50 | ≥80 | | |



Page 2 of 7



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| 1 | Test Name | l | Results | Units | 1 | o. Re interv | val |
|---|-----------|------|---------|-------|---|--------------|-----|
| | High | <70 | <100 | ≥70 | | ≥100 | |
| | Moderate | <100 | <130 | ≥100 | | | |
| | Low | <100 | <130 | ≥130* | | ≥160* | |

^{*}In low risk patient, consider therapy after an initial non-pharmacological intervention for at least 2 nonths

| GLUCOSE, FASTING (F) | | V | |
|---------------------------------|-------|-------|----------|
| Glucose Fasting (Hexokinase) | 80.00 | mg/dL | 70 - 100 |

| VITAMIN B12; CYANOCOBALAMIN | | | |
|-----------------------------|------|-------|---------------|
| (CLIA) | | | |
| Vitamin B12: Cyanocohalamin | 0.00 | na/ml | 211.00 011.00 |

Vitamin B12; Cyanocobalamin pg/mL 211.00 - 911.00

VITAMIN D, 25 - HYDROXY, SERUM (CLIA)

Vitamin D, 25 Hydroxy 150.00 nmol/L 75.00 - 250.00

Interpretation

| LEVEL | | REFER | RANGE IN nmol/L | COMMENTS |
|---------------------------|---|-------|-----------------|--|
| Deficient | | | < 50 | High risk for developing bone disease |
| ⊤nsufficier. | | | 50-74 | Vitamin D concentration which normalizes Parathyroid hormone concentration |
| s' int | - | | 75-250 | Optimal concentration for maximal health benefit |
| Potenti I intoxi ation | | | >250 | High risk for toxic effects |

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.



Page 3 of 7



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Test Name Results Units . Reunterval

Levels vary with age and are increased in pregnancy.

• A new test Vitamin D, Ultrasensitive by LC-MS/MS is also available

| THYROID PROFILE,TOTAL, SERUM (CLIA) | | |
|-------------------------------------|------------|-----------------|
| T3, Total | 1.00 | 0.60 - 1.81 |
| T4, Total | 7.00 µg/dL | 5.01 - 12.45 |
| TSH | 3.00 µIU/m | L 0.550 - 4.780 |

Note

- 1. TSH levels are subject to circum variation, reading peak levels between 2 4.a.m. and at a minimum between 6-10 pm. The miation continued the order of 50%. Hence time of the day has influence on the measured serum Tamporcentrations.
- 2. Alteration in concentration of a roid harmone binding protein can profoundly affect Total T3 and/or Total T4 levels especified in pregnancy and in patients on steroid therapy.
- 3. Unbound fraction (Free, T3) of thyroid hormone is biologically active form and correlate more closely with light state of the patient than total T4/T3 concentration
- 4. Values <0 TU/mL to be unically correlated due to presence of a rare TSH variant in some intrividua.





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| Test Name | Results | Units | Re' Interval |
|---|---------|--------|--------------|
| HbA1c (GLYCOSYLATED HEMOGLOBIN), BLOOD (HPLC, NGSP certified) | | | |
| HbA1c | 10.0 | % | 4.00 - 5.60 |
| Estimated average glucose (eAG) | 240 | IIIs - | |
| | | | |

Interpretation

HbA1c result is suggestive of Diabetes/ Higher than glycem Lin and Diabetic patient.

Please note, Glycemic goal should be individualized based on duration of diabetes, age/life expectancy, comorbid conditions, known CVD or advanced microvascular complications, hypoglycaemia unawareness, and individual patient considerations

Interpretation as per American Diabete ssocia pp (ADA) Guidelines

| | Reference Group | Non diabetic adults >=18 | risk ediabetes) | Diagnosing Diabetes | Therapeutic goals for glycemic control |
|---|-----------------|-----------------------------|--------------------|--------------------------|---|
| l | HbA1c in % | 4.0-5.6 | 5.7-6.4 | >= 6.5 | <7.0 |

Note: Presence of Hemoglations rights a particularly when the Highest results as not arrelate with the patient's blood glucose levels.

| MEASUREM | OF HBA1C RESULTS |
|--|--|
| temoglo bin veriants, elevated fetal hemoglobic (H. and chemically modified myatives of hemoglobin lated Hb in patients all failure) can affect the accuracy of HbAlc measurements | Any condition that shortens erythrocyte survival or decreases mean erythrocyte age (e.g.,recovery from acute blood loss, hemolytic anemia, HbSS, HbCC, and HbSC) will falsely lower HbAlc test results regardless of the assay method used.Iron deficiency anemia is associated with |

| higher HbA1c



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|--|---------|----------|-----------------|
| COMPLETE BLOOD COUNT; CBC | | | |
| Hemoglobin (Photometry) | 15.00 | M. | 13.00 - 17.00 |
| Packed Cell Volume (PCV) (Calculated) | 45.00 | % | 40.00 - 50.00 |
| RBC Count (Electrical Impedence) | 5.00 | mir/mm3 | 4.50 - 5.50 |
| MCV (Electrical Impedence) | 10° | fL | 83.00 - 101.00 |
| MCH (Calculated) | 30.00 | pg | 27.00 - 32.00 |
| MCHC (Calculated) | 33.00 | g/dL | 31.50 - 34.50 |
| Red Cell Distribution Width (RDW) (Electrical Impedence) | 12.00 | % | 11.60 - 14.00 |
| Total Leukocyte Count (TLC) (Electrical Impedence) | 5.00 | thou/mm3 | 4.00 - 10.00 |
| Differential Leucocyte Count (DLC) (VCS Technology) | | | |
| Segmented Neutrophils | 50.00 | % | 40.00 - 80.00 |
| Lymphocytes | 40.00 | % | 20.00 - 40.00 |
| Monocytes | 5.00 | % | 2.00 - 10.00 |
| Eosinophils | 4.00 | % | 1.00 - 6.00 |
| Basophils | 1.00 | % | <2.00 |
| Absolute Level Count (Colculated | | | |
| Neurophils | 2.50 | thou/mm3 | 2.00 - 7.00 |
| Lymr | 2.00 | thou/mm3 | 1.00 - 3.00 |
| ocytes | 0.25 | thou/mm3 | 0.20 - 1.00 |
| Eosinophil | 0.20 | thou/mm3 | 0.02 - 0.50 |
| Basopals | 0.05 | thou/mm3 | 0.02 - 0.10 |
| Platelet Count (Electrical impedence) | 151 | thou/mm3 | 150.00 - 410.00 |
| Mean Platelet Volume (Electrical Impedence) | 6.5 | fL | 6.5 - 12.0 |

Note

1. As per the recommendation of International council for Standardization in Hematology, the differential



Page 6 of 7



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Test Report

o. R... Interval **Test Name** Results Units leucocyte counts are additionally being reported as absolute numbers of each cell in per unit v ume of

blood

2. Test conducted on EDTA whole blood

May Low

MCI - 24779

Dr Aiav Gupta MD, Pathology

Technical Director - Hematology &

Immunology

Dr.Kamal Modi

NRL - Dr Lal PathLabs Ltd

NRL - Dr Lal PathLabs Ltd

Hematopathology NRL - Dr Lal PathLabs Ltd

Senior Consultant and Lead

MD. Biochemistry Consultant Biochemist

Technical Director

DMC - 9550 Dr Nimmi Kans

Dr Gurleen Oberoi

MD.DNB.MNAMS

DM(Hematopathology)

MD. Biochemi

& Biochemical Geneti NRL - Dr Lal PathLabs Ltd

DMC - 89819

Dr Hi

Consulta Biochemist

DMC - 24201

Dr Sarita Kumari Lal b, Pathology

onsultant Pathologist Dr Lal PathLabs Ltd

DMC - 46663

Dr Sunanda

MD, Pathology

Consultant Pathologist Dr Lal PathLabs Ltd

MD, Pathology

Sr. Consultant Pathologist -Hematology & Immunology

NRL - Dr Lal PathLabs Ltd

This is a revised report & supersedes all he preusly issued reports



IMPORTANT INSTRUCTIONS

eleased pertain to the specimen submitted. All test results are dependent on the quality of the sample received by the Laboratory. estigations are only a tool to facilitate in arriving at a diagnosis and should be clinically correlated by the Referring Physician .•Report Laboratory 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 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 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.

Tel: +91-11-49885050,Fax: - +91-11-2788-2134, E-mail: lalpathlabs@lalpathlabs.com

National Reference lab, Delhi, a CAP (7171001) Accredited, ISO 9001:2015 (FS60411) & ISO 27001:2013 (616691) Certified laboratory.



Page 7 of 7