Name	: Mr. DUMMY		
Lab No.	: WM15SPF	Age	: 25 Years
Ref By	: SELF	Gender	: Male
Collected	: 7/11/2023 11:08:00AM	Reported	: 11/11/2023 11:54:11AM
A/c Status	: P	Report Status	: Final
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MC-2113

Test Report

SWASTHFIT SUPER 2

Test Name	Results	Units	Bio. Ref. Interval
LIVER & KIDNEY PANEL, SERUM			
Creatinine (Modified Jaffe,Kinetic)	0.72	mg/dL	0.70 - 1.30
GFR Estimated (CKD EPI Equation 2021)	130	mL/min/1.73m²	>59
GFR Category (KDIGO Guideline 2012)	G1		
Urea (Urease UV)	14.00	mg/dL	13.00 - 43.00
Urea Nitrogen Blood (Calculated)	6.54	mg/dL	6.00 - 20.00
BUN/Creatinine Ratio (Calculated)	9		
Uric Acid (Uricase)		mg/dL	3.50 - 7.20
AST (SGOT) (IFCC without P5P)	16.0	U/L	15.00 - 40.00
ALT (SGPT) (IFCC without P5P)	11.0	U/L	10.00 - 49.00
GGTP (IFCC)	72.0	U/L	0 - 73
Alkaline Phosphatase (ALP) (IFCC-AMP)	31.00	U/L	30.00 - 120.00
Bilirubin Total (Oxidation)	1.00	mg/dL	0.30 - 1.20
Bilirubin Direct (Oxidation)	0.20	mg/dL	<0.3
Bilirubin Indirect (Calculated)	0.80	mg/dL	<1.10
Total Protein (Biuret)	**3.50**	g/dL	5.70 - 8.20
Albumin (BCG)	**3.10**	g/dL	3.20 - 4.80
A : G Ratio (Calculated)	**7.75**		0.90 - 2.00
Globulin(Calculated)	**0.40**	gm/dL	2.0 - 3.5
Calcium, Total (Arsenazo III)	10.00	mg/dL	8.70 - 10.40

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Test Report

Test Name	Results	Units	Bio. Ref. Interval
Phosphorus (Molybdate UV)	3.40	mg/dL	2.50 - 5.10
Sodium (Indirect ISE)	137.00	mEq/L	136.00 - 145.00
Potassium (Indirect ISE)	4.00	mEq/L	3.50 - 5.10
Chloride (Indirect ISE)	99.00	mEq/L	98.00 - 107.00

LIPID SCREEN, SERUM

Cholesterol, Total (CHO-POD)	151.00	mg/dL	<200.00
Triglycerides (GPO-POD)	145.00	mg/dL	<150.00
HDL Cholesterol (Enz Immunoinhibition)	41.00	mg/dL	>40.00
LDL Cholesterol, Calculated (Calculated)	81.00	mg/dL	<100.00
VLDL Cholesterol,Calculated (Calculated)	29.00	mg/dL	<30.00
Non-HDL Cholesterol (Calculated)	110	mg/dL	<130

Note

1. Measurements in the same patient can show physiological & analytical variations. Three serial samples 1 week 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 patients with moderate risk for ASCVD for risk refinement.

Treatment Goals as per Lipid Association of India 2020

RISK CATEGORY	TREATMENT GOAL		CONSIDER THERAPY	
	LDL CHOLESTEROL (LDL-C)(mg/dL)	NON HDL CHLOESTEROL (NON HDL-C) (mg/dL)	LDL CHOLESTEROL (LDL-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

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Test Name		Results	Units	Bio. Ref. Interval
High	<70	<100	≥70	≥100
Moderate	<100	<130	≥100	[illegible]0
Low	<100	<130	≥130*	≥160*

*In low risk patient, consider therapy after an initial non-pharmacological intervention for at l[illegible] months

GLUCOSE, FASTING (F)

Glucose Fasting (Hexokinase)	100.00	mg/dL	70 - 100

THYROID PROFILE,TOTAL, SERUM
(CLIA)

T3, Total	1.[illegible]0	ng/mL	0.60 - 1.81
T4, Total	5.[illegible]	µg/dL	5.01 - 12.45
TSH	4.00	µIU/mL	0.550 - 4.780

Note

1. TSH levels are subject to circadian variation, reaching peak levels between 2 - 4.a.m. and at a minimum between 6-[illegible]. The variation is of the order of 50% . hence time of the day has influenced on the measured serum TSH concentrations.
2. Alteration in concentration of Thyroid hormone binding protein can profoundly affect Total T3 and/or Total T4 levels especially in pregnancy and in patients on steroid therapy.
3. Unbound fraction (Free,T4 /Free,T3) of thyroid hormone is biologically active form and correlate more closely with clinical status of the patient than total T4/T3 concentration
Values <0.03 uIU/mL need to be clinically correlated due to presence of a rare TSH variant in some individuals

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Test Report

Test Name	Results	Units	Ref Interval
HbA1c (GLYCOSYLATED HEMOGLOBIN), BLOOD (HPLC, NGSP certified)			
HbA1c	5.0	%	4.00 - 5.60
Estimated average glucose (eAG)	97	mg/dL	

Interpretation

HbA1c result is suggestive of non diabetic adults (>=18 years) and could be labelled Diabetes in a known Diabetic

Interpretation as per American Diabetes Association (ADA) Guidelines

Reference Group	Non diabetic adults >=18 years	At risk (Pre-diabetes)	Diagnosing Diabetes	Therapeutic goals for glycemic control
HbA1c in %	4.0-5.6	5.7-6.4	>= 6.5	<7.0

Note: Presence of Hemoglobin variants and/or conditions that affect red cell turnover must be considered, particularly when the HbA1C result does not correlate with the patient's blood glucose levels.

FACTORS THAT INTERFERE WITH HbA1c MEASUREMENT	FACTORS THAT AFFECT INTERPRETATION OF HBA1C RESULTS
Hemoglobin variants, elevated fetal hemoglobin (HbF), and chemically modified derivatives of hemoglobin (e.g. carbamylated Hb in patients with renal failure) can affect the accuracy of HbA1c 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 HbA1c test results regardless of the assay method used.Iron deficiency anemia is associated with higher HbA1c

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Test Report

Test Name	Results	Units	Bio. Ref. Interval
COMPLETE BLOOD COUNT; CBC			
Hemoglobin (Photometry)	14.00	g/dL	13.00 - 17.00
Packed Cell Volume (PCV) (Calculated)	41.00	%	40.00 - 50.00
RBC Count (Electrical Impedence)	5.00	mill/mm3	4.50 - 5.50
MCV (Electrical Impedence)	84	fL	83.00 - 101.00
MCH (Calculated)	28.00	pg	27.00 - 32.00
MCHC (Calculated)	32.00	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	8.00	%	2.00 - 10.00
Eosinophils	1.00	%	1.00 - 6.00
Basophils	1.00	%	<2.00
Absolute Leucocyte Count (Calculated)			
Neutrophils	2.50	thou/mm3	2.00 - 7.00
Lymphocytes	2.00	thou/mm3	1.00 - 3.00
Monocytes	0.40	thou/mm3	0.20 - 1.00
Eosinophils	0.05	thou/mm3	0.02 - 0.50
Basophils	0.05	thou/mm3	0.02 - 0.10
Platelet Count (Electrical impedence)	151	thou/mm3	150.00 - 410.00
Mean Platelet Volume (Electrical Impedence)	11.0	fL	6.5 - 12.0

Note

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

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Test Report

Test Name	Results	Units	Bio. Ref. Interval

leucocyte counts are additionally being reported as absolute numbers of each cell in per unit volume of blood

2. Test conducted on EDTA whole blood

MCI – 24779
Dr Ajay Gupta
MD, Pathology
Technical Director - Hematology & Immunology
NRL - Dr Lal PathLabs Ltd

DMC – 77091
Dr Gurleen Oberoi
DM(Hematopathology), MD,DNB,MNAMS
Senior Consultant and Lead-Hematopathology
NRL - Dr Lal PathLabs Ltd

DMC - 89819
Dr Hi Mazumda
MD Biochemistry
Consultant Biochemist
Dr Lal Ltd

Dr Ja Munjal
MD,Pathology
Consultant Pathologist
Dr Lal PathLabs Ltd

DMC – 89936
Dr.Kamal Modi
MD, Biochemistry
Consultant Biochemist
NRL - Dr Lal PathLabs Ltd

DMC – 9550
Dr Nimmi Kansa
MD, Biochemi
Technical Director - l Chemi
& Biochemical Genetics
NRL - Dr Lal PathLabs Ltd

DMC - 24201
Dr Sarita Kumari Lal
D, Pathology
Consultant Pathologist
Dr Lal PathLabs Ltd

DMC – 46663
Dr Sunanda
MD, Pathology
Sr. Consultant Pathologist - Hematology & Immunology
NRL - Dr Lal PathLabs Ltd

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

IMPORTANT INSTRUCTIONS

•Test relea pertain to the specimen submitted.•All test results are dependent on the quality of the sample received by the Laboratory. Labo estigations 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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National Reference lab, Delhi, a CAP (7171001) Accredited, ISO 9001:2015 (FS60411) & ISO 27001:2013 (616691) Certified laboratory.