

## ETERNAL HOSPITAL MEDICAL TESTING LABORATORY

<b>Patient Name</b>	Mrs. VIBHA MATHUR	<b>Lab No</b>	4001322
<b>UHID</b>	40001141	<b>Collection Date</b>	11/03/2023 11:17AM
<b>Age/Gender</b>	33 Yrs/Female	<b>Receiving Date</b>	11/03/2023 11:22AM
<b>IP/OP Location</b>	O-OPD	<b>Report Date</b>	12/03/2023 8:42PM
<b>Referred By</b>	Dr. DIWANSHU KHATANA	<b>Report Status</b>	Final
<b>Mobile No.</b>	9079965069		

### BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range	Sample: Fl. Plasma
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#### BLOOD GLUCOSE (FASTING)

BLOOD GLUCOSE FASTING 130.1

Method: Hexokinase assay.

Interpretation:-Diagnosis and monitoring of treatment in diabetes mellitus and evaluation of carbohydrate metabolism in various diseases.

#### BLOOD GLUCOSE (PP )

BLOOD GLUCOSE (PP ) 100.6 mg/dl

Non – Diabetic: - < 140 mg/dl  
Pre – Diabetic: - 140-199 mg/dl  
Diabetic: - >=200 mg/dl

Sample: PLASMA

Method: Hexokinase assay.

Interpretation:-Diagnosis and monitoring of treatment in diabetes mellitus and evaluation of carbohydrate metabolism in various diseases.

#### THYROID T3 T4 TSH

	Result	Unit	Biological Ref. Range	Sample: Serum
T3	1.41	ng/mL	0.970 - 1.690	
T4	9.56	ug/dl	5.53 - 11.00	
TSH	1.352	μIU/mL	0.40 - 4.05	

RESULT ENTERED BY : NEETU SHARMA



Dr. MUDITA SHARMA

MBBS|MD| PATHOLOGY

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### BIOCHEMISTRY

**T3**:- Method: ElectroChemiLuminescence ImmunoAssay - ECLIA

Interpretation:-The determination of T3 is utilized in the diagnosis of T3-hyperthyroidism the detection of early stages of hyperthyroidism and for indicating a diagnosis of thyrotoxicosis factitia.

**T4**:- Method: ElectroChemiLuminescence ImmunoAssay - ECLIA

Interpretation:-The determination of T4 assay employs a competitive test principle with an antibody specifically directed against T4.

**TSH - THYROID STIMULATING HORMONE** :- ElectroChemiLuminescenceImmunoAssay - ECLIA

Interpretation:-The determination of TSH serves as the initial test in thyroid diagnostics. Even very slight changes in the concentrations of the free thyroid hormones bring about much greater opposite changes in the TSH levels.

### LFT (LIVER FUNCTION TEST)

Sample: Serum

BILIRUBIN TOTAL	0.37	mg/dl	0.00 - 1.20
BILIRUBIN INDIRECT	0.24	mg/dl	0.20 - 1.00
BILIRUBIN DIRECT	0.13	mg/dl	0.00 - 0.40
SGOT	14.2	U/L	0.0 - 40.0
SGPT	15.8	U/L	0.0 - 40.0
TOTAL PROTEIN	7.90	g/dl	6.6 - 8.7
ALBUMIN	4.95	g/dl	3.5 - 5.2
GLOBULIN	3.0		1.8 - 3.6
ALKALINE PHOSPHATASE	54.2	U/L	42 - 98
A/G RATIO	1.7	Ratio	1.5 - 2.5
GGTP	12.1	U/L	6.0 - 38.0

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### BIOCHEMISTRY

**BILIRUBIN TOTAL** :- Method: DPD assay. Interpretation:-Total Bilirubin measurements are used in the diagnosis and treatment of various liver diseases, and of haemolytic and metabolic disorders in adults and newborns. Both obstruction damage to hepatocellular structure.

**BILIRUBIN DIRECT** :- Method: Diazo method Interpretation:-Determinations of direct bilirubin measure mainly conjugated, water soluble bilirubin.

**SGOT - AST** :- Method: IFCC without pyridoxal phosphate activation. Interpretation:-SGOT(AST) measurements are used in the diagnosis and treatment of certain types of liver and heart disease.

**SGPT - ALT** :- Method: IFCC without pyridoxal phosphate activation. Interpretation:-SGPT(ALT) Ratio Is Used For Differential Diagnosis In Liver Diseases.

**TOTAL PROTEINS** :- Method: Biuret colorimetric assay. Interpretation:-Total protein measurements are used in the diagnosis and treatment of a variety of liver and kidney diseases and bone marrow as well as metabolic and nutritional disorder.

**ALBUMIN** :- Method: Colorimetric (BCP) assay. Interpretation:-For Diagnosis and monitoring of liver diseases, e.g. liver cirrhosis, nutritional status.

**ALKALINE PHOSPHATASE** :- Method: Colorimetric assay according to IFCC. Interpretation:-Elevated serum ALT is found in hepatitis, cirrhosis, obstructive jaundice, carcinoma of the liver, and chronic alcohol abuse. ALT is only slightly elevated in patients who have an uncomplicated myocardial infarction. **GGTP-GAMMA GLUTAMYL TRANSPEPTIDASE** :- Method:

Enzymatic colorimetric assay. Interpretation:- $\gamma$ -glutamyltransferase is used in the diagnosis and monitoring of hepatobiliary disease. Enzymatic activity of GGT is often the only parameter with increased values when testing for such diseases and is one of the most sensitive indicator known.

### LIPID PROFILE

TOTAL CHOLESTEROL	171		<200 mg/dl :- Desirable 200-240 mg/dl :- Borderline >240 mg/dl :- High
HDL CHOLESTEROL	51.0		High Risk :-<40 mg/dl (Male), <40 mg/dl (Female) Low Risk :->=60 mg/dl (Male), >=60 mg/dl (Female)
LDL CHOLESTEROL	94.4		Optimal :- <100 mg/dl Near or Above Optimal :- 100-129 mg/dl Borderline :- 130-159 mg/dl High :- 160-189 mg/dl Very High :- >190 mg/dl
CHOLESTERO VLDL	13	mg/dl	10 - 50
TRIGLYCERIDES	67.2		Normal :- <150 mg/dl Border Line:- 150 - 199 mg/dl High :- 200 - 499 mg/dl Very high :- > 500 mg/dl
CHOLESTEROL/HDL RATIO	3.4	%	

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### BIOCHEMISTRY

CHOLESTEROL TOTAL :- Method: CHOD-PAP enzymatic colorimetric assay.

interpretation:-The determination of the individual total cholesterol (TC) level is used for screening purposes while for a better risk assessment it is necessary to measure additionally lipid & lipoprotein metabolic disorders.

HDL CHOLESTEROL :- Method:-Homogenous enzymetic colorimetric method.

Interpretation:-HDL-cholesterol has a protective against coronary heart disease, while reduced HDL-cholesterol concentrations, particularly in conjunction with elevated triglycerides, increase the cardiovascular disease.

LDL CHOLESTEROL :- Method: Homogenous enzymatic colorimetric assay.

Interpretation:-LDL play a key role in causing and influencing the progression of atherosclerosis and in particular coronary sclerosis. The LDL are derived from VLDL rich in TG by the action of various lipolytic enzymes and are synthesized in the liver.

CHOLESTEROL VLDL :- Method: VLDL Calculative

TRIGLYCERIDES :- Method: GPO-PAP enzymatic colorimetric assay.

Interpretation:-High triglyceride levels also occur in various diseases of liver, kidneys and pancreas. DM, nephrosis, liver obstruction.

CHOLESTEROL/HDL RATIO :- Method: Cholesterol/HDL Ratio Calculative

### RENAL PROFILE TEST

Sample: Serum

UREA	14.9 L	mg/dl	16.60 - 48.50
BUN	7.0	mg/dl	6 - 20
CREATININE	0.58	mg/dl	0.50 - 0.90
SODIUM	138.7	mmol/L	136 - 145
POTASSIUM	4.08	mmol/L	3.50 - 5.50
CHLORIDE	102.3	mmol/L	98 - 107
URIC ACID	1.69 L	mg/dl	2.6 - 6.0
CALCIUM	8.79	mg/dl	8.60 - 10.30

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### BIOCHEMISTRY

**CREATININE - SERUM** :- Method:-Jaffe method, Interpretation:-To differentiate acute and chronic kidneydisease.

**URIC ACID** :- Method: Enzymatic colorimetric assay. Interpretation:- Elevated blood concentrations of uricacid are renal diseases with decreased excretion of waste products, starvation,drug abuse and increased alcohol consume.

**SODIUM**:- Method: ISE electrode. Interpretation:-Decrease: Prolonged vomiting or diarrhea,diminshed reabsorption in the kidney and excessive fluid retention. Increase: excessive fluid loss, high salt intake andkidney reabsorption.

**POTASSIUM** :- Method: ISE electrode. Intrapretation:-Low level: Intake excessive loss formbodydue to diarrhea, vomiting renal failure, High level: Dehydration, shock severe burns, DKA, renalfailure.

**CHLORIDE - SERUM** :- Method: ISE electrode. Interpretation:-Decrease: reduced dietary intake,prolonged vomiting and reduced renal reabsorption as well as forms of acidosisand alkalosis.

Increase: dehydration, kidney failure, some form ofacidosis, high dietary or parenteral chloride intake, and salicylate poisoning.

**UREA**:- Method: Urease/GLDH kinetic assay. Interpretation:-Elevations in blood urea nitrogenconcentration are seen in inadequate renal perfusion, shock, diminished bloodvolume, chronic nephritis, nephrosclerosis, tubular necrosis, glomerularnephritis and UTI.

**CALCIUM TOTAL** :- Method: O-Cresolphthaleine complexone. Interpretation:-Increase in serum PTH or vit-D are usuallyassociated with hypercalcemia. Increased serum calcium levels may also beobserved in multiple myeloma and other neoplastic diseases. Hypocalcemia may beobserved in hypoparathyroidism, nephrosis, and pancreatitis.

Sample: WHOLE BLOOD EDTA

<b>HBA1C</b>	5.3	%		
			< 5.7%	Nondiabetic
			5.7-6.4%	Pre-diabetic
			> 6.4%	Indicate Diabetes
				Known Diabetic Patients
			< 7 %	Excellent Control
			7 - 8 %	Good Control
			> 8 %	Poor Control

Method : - High - performance liquid chromatography HPLC

Interpretation:-Monitoring long term glycemic control, testing every 3 to 4 months is generally sufficient. The approximate relationship between HbA1C and mean blood glucose values during the preceding 2 to 3 months.

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### BLOOD BANK INVESTIGATION

Test Name	Result	Unit	Biological Ref. Range
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BLOOD GROUPING	"B" Rh Negative		
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Note :

1. Both forward and reverse grouping performed.
2. Test conducted on EDTA whole blood.

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### CLINICAL PATHOLOGY

Test Name	Result	Unit	Biological Ref. Range	Sample: Urine
<u>URINE SUGAR (RANDOM)</u>				
URINE SUGAR (RANDOM)	NEGATIVE			

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### CLINICAL PATHOLOGY

Test Name	Result	Unit	Biological Ref. Range	Sample: Urine
<b><u>ROUTINE EXAMINATION - URINE</u></b>				
<b>PHYSICAL EXAMINATION</b>				
VOLUME	25	ml		
COLOUR	PALE YELLOW		P YELLOW	
APPEARANCE	CLEAR		CLEAR	
<b>CHEMICAL EXAMINATION</b>				
PH	6.0		5.5 - 7.0	
SPECIFIC GRAVITY	1.000		1.016-1.022	
PROTEIN	NIL		NEGATIVE	
SUGAR	NIL		NEGATIVE	
BILIRUBIN	NIL		NEGATIVE	
BLOOD	NIL			
KETONES	NIL		NEGATIVE	
NITRITE	NIL		NEGATIVE	
UROBILINOGEN	NIL		NEGATIVE	
LEUCOCYTE	NIL		NEGATIVE	
<b>MICROSCOPIC EXAMINATION</b>				
WBCS/HPF	3-4	/hpf	0 - 3	
RBCS/HPF	00	/hpf	0 - 2	
EPITHELIAL CELLS/HPF	10-15	/hpf	0 - 1	
CASTS	NIL		NIL	
CRYSTALS	NIL		NIL	
BACTERIA	NIL		NIL	
OHTERS	NIL		NIL	

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Methodology:-

Glucose: GOD-POD, Bilirubin: Diazo-Azo-coupling reaction with a diazonium, Ketone: Nitro Pruside reaction, Specific Gravity: Proton release from ions, Blood: Psuedo-Peroxidase activity oh Haem moiety, pH: Methye Red-Bromothymol Blue (Double indicator system), Protein: H+ Release by buffer, microscopic & chemical method. interpretation: Diagnosis of Kidney function, UTI, Presence of Protein, Glucoses, Blood. Vocubulary syntax: Kit insert

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### HEMATOLOGY

Test Name	Result	Unit	Biological Ref. Range
<b><u>CBC (COMPLETE BLOOD COUNT)</u></b>			
Sample: WHOLE BLOOD EDTA			
HAEMOGLOBIN	<b>11.8 L</b>	g/dl	12.0 - 15.0
PACKED CELL VOLUME(PCV)	36.6	%	36.0 - 46.0
MCV	87.6	fl	82 - 92
MCH	28.2	pg	27 - 32
MCHC	32.2	g/dl	32 - 36
RBC COUNT	4.18	millions/cu.mm	3.80 - 4.80
TLC (TOTAL WBC COUNT)	5.09	10 <sup>3</sup> / uL	4 - 10
<b><u>DIFFERENTIAL LEUCOCYTE COUNT</u></b>			
NEUTROPHILS	70.1	%	40 - 80
LYMPHOCYTE	21.4	%	20 - 40
EOSINOPHILS	<b>0.4 L</b>	%	1 - 6
MONOCYTES	7.5	%	2 - 10
BASOPHIL	<b>0.6 L</b>	%	1 - 2
PLATELET COUNT	2.14	lakh/cumm	1.500 - 4.500

**HAEMOGLOBIN** :- Method:-SLS HemoglobinMethodology by Cell Counter.Interpretation:-Low-Anemia, High-Polycythemia.  
**MCV** :- Method:- Calculation bysystemex.  
**MCH** :- Method:- Calculation bysystemex.  
**MCHC** :- Method:- Calculation bysystemex.  
**RBC COUNT** :- Method:-Hydrodynamicfocusing.Interpretation:-Low-Anemia,High-Polycythemia.  
**TLC (TOTAL WBC COUNT)** :- Method:-Optical Detectorblock based on Flowcytometry.Interpretation:-High-Leucocytosis, Low-Leucopenia.  
**NEUTROPHILS** :- Method: Optical detectorblock based on Flowcytometry  
**LYMPHOCYTS** :- Method: Optical detectorblock based on Flowcytometry  
**EOSINOPHILS** :- Method: Optical detectorblock based on Flowcytometry  
**MONOCYTES** :- Method: Optical detectorblock based on Flowcytometry  
**BASOPHIL** :- Method: Optical detectorblock based on Flowcytometry  
**PLATELET COUNT** :- Method:-Hydrodynamicfocusing method.Interpretation:-Low-Thrombocytopenia, High-Thrombocytosis.  
**HCT**: Method:- Pulse Height Detection. Interpretation:-Low-Anemia, High-Polycythemia.  
**NOTE**: CH- CRITICAL HIGH, CL: CRITICAL LOW, L: LOW, H: HIGH

ESR (ERYTHROCYTE SEDIMENTATION RATE)	15	mm/1st hr	0 - 15
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Method:-Modified Westergrens.

Interpretation:-Increased in infections, sepsis, and malignancy.

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

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