**Patient Name** Mr. BHAGIRATH SINGH Lab No 4001316 UHID 40001134 **Collection Date** 11/03/2023 11:19AM 11/03/2023 11:20AM Age/Gender 41 Yrs/Male **Receiving Date Report Date IP/OP Location** O-OPD 11/03/2023 4:20PM

**Referred By** Dr. DIWANSHU KHATANA **Report Status** Final

Mobile No. 7506363465

### **BIOCHEMISTRY**

**Test Name** Result Unit **Biological Ref. Range BLOOD GLUCOSE (FASTING)** Sample: Fl. Plasma BLOOD GLUCOSE FASTING 155.5

Method: Hexokinase assay.

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

**BLOOD GLUCOSE (PP)** Sample: PLASMA

BLOOD GLUCOSE (PP) 164.7 Non – Diabetic: - < 140 mg/dl mg/dl

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

Method: Hexokinase assay.

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

**THYROID T3 T4 TSH** Sample: Serum

T3	1.25	ng/mL	0.970 - 1.690
T4	5.28 L	ug/dl	5.53 - 11.00
TSH	2.543	μIU/mL	0.40 - 4.05

**RESULT ENTERED BY: NEETU SHARMA** Os garrie.

Dr. MUDITA SHARMA

Patient Name	Mr. BHAGIRATH SINGH	Lab No	4001316
UHID	40001134	Collection Date	11/03/2023 11:19AM
Age/Gender IP/OP Location	41 Yrs/Male	Receiving Date	11/03/2023 11:20AM
	O-OPD	Report Date	11/03/2023 4:20PM
Referred By	Dr. DIWANSHU KHATANA	Report Status	Final
Mobile No.	7506363465		

#### **BIOCHEMISTRY**

T3:- Method: ElectroChemiLuminescence ImmunoAssay - ECLIA

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

T4:- Method: ElectroChemiLuminescence ImmunoAssay - ECLIA

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

TSH - THYROID STIMULATING HORMONE :- ElectroChemiLuminescenceImmunoAssay - ECLIA

2.4

22.4

Interpretation: - The determination of TSH serves as theinitial 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	0.62	mg/dl	0.00 - 1.20	
BILIRUBIN INDIRECT	0.34	mg/dl	0.20 - 1.00	
BILIRUBIN DIRECT	0.28	mg/dl	0.00 - 0.40	
SGOT	24.6	U/L	0.0 - 40.0	
SGPT	27.9	U/L	0.0 - 40.0	
TOTAL PROTEIN	7.68	g/dl	6.6 - 8.7	
ALBUMIN	5.40 H	g/dl	3.5 - 5.2	
GLOBULIN	2.3		1.8 - 3.6	
ALKALINE PHOSPHATASE	95.5	U/L	53 - 128	

Ratio

U/L

1.5 - 2.5

10.0 - 55.0

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A/G RATIO

**GGTP** 

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**Patient Name** Mr. BHAGIRATH SINGH Lab No 4001316 UHID **Collection Date** 11/03/2023 11:19AM 40001134 11/03/2023 11:20AM Age/Gender **Receiving Date** 41 Yrs/Male Report Date O-OPD **IP/OP Location** 11/03/2023 4:20PM

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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 structive.

BILLRUBIN 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.

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: Enzymetic colorimetric assay. Interpretation:-y-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	184		<200 mg/dl :- Desirable 200-240 mg/dl :- Borderline >240 mg/dl :- High
HDL CHOLESTEROL	47.1		High Risk :-<40 mg/dl (Male), <40 mg/dl (Female) Low Risk :->=60 mg/dl (Male), >=60 mg/dl (Female)
LDL CHOLESTEROL	122.1		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	15	mg/dl	10 - 50
TRIGLYCERIDES	75.1		Normal :- <150 mg/dl Border Line:- 150 - 199 mg/dl High :- 200 - 499 mg/dl Very high :- > 500 mg/dl
CHOLESTEROL/HDL RATIO	3.9	%	

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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 form 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 triglycerde 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	31.4	mg/dl	16.60 - 48.50
BUN	14.7	mg/dl	6 - 20
CREATININE	0.80	mg/dl	0.60 - 1.10
SODIUM	141.2	mmol/L	136 - 145
POTASSIUM	4.27	mmol/L	3.50 - 5.50
CHLORIDE	103.8	mmol/L	98 - 107
URIC ACID	3.53	mg/dl	3.5 - 7.2
CALCIUM	9.72	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, diminished reabsorption in the kidney and excessive fluid retention. Increase: excessive fluid loss, high salt intake and kidney reabsorption.

POTASSIUM:- Method: ISE electrode. Intrpretation:-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 usually associated with hypercalcemia. Increased serum calcium levels may also be observed in multiple myeloma and other neoplastic diseases. Hypocalcemia may

beobserved in hypoparathyroidism, nephrosis, and pancreatitis.

Sample: WHOLE BLOOD EDTA

HBA1C 5.8 % < 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**

**Biological Ref. Range Test Name** Result Unit

**BLOOD GROUPING** "AB" Rh Positive

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

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1-2

# **CLINICAL PATHOLOGY**

**Test Name** Result Unit **Biological Ref. Range** Sample: Urine **URINE SUGAR (POST PRANDIAL)** URINE SUGAR (POST PRANDIAL) NEGATIVE **URINE SUGAR (RANDOM)** Sample: Urine **URINE SUGAR (RANDOM) NEGATIVE ROUTINE EXAMINATION - URINE** Sample: Urine **PHYSICAL EXAMINATION VOLUME** 20 ml COLOUR PALE YELLOW P YELLOW **APPEARANCE** CLEAR CLEAR **CHEMICAL EXAMINATION** РΗ 7.0 5.5 - 7.0 SPECIFIC GRAVITY 1.000 1.016-1.022 **PROTEIN** NIL NEGATIVE NEGATIVE **SUGAR** NIL **BILIRUBIN** NIL **NEGATIVE BLOOD** NIL **KETONES** NIL NEGATIVE NITRITE **NEGATIVE** NIL UROBILINOGEN NIL NEGATIVE NEGATIVE LEUCOCYTE NIL MICROSCOPIC EXAMINATION WBCS/HPF 1-2 /hpf 0 - 3

/hpf

0 - 1

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**EPITHELIAL CELLS/HPF** 

Mobile No.

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#### Methodology:-

Glucose: GOD-POD, Bilirubin: Diazo-Azo-coupling reaction with a diazonium, Ketone: Nitro Pruside reaction, Specific Gravity: Proton re; ease 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

**RESULT ENTERED BY: NEETU SHARMA** 

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#### **HEMATOLOGY**

Result	Unit	Biological Ref. Range
		Sample: WHOLE BLOOD EDTA
15.6	g/dl	13.0 - 17.0
46.5	%	40.0 - 50.0
86.3	fl	82 - 92
28.9	pg	27 - 32
33.5	g/dl	32 - 36
5.39	millions/cu.mm	4.50 - 5.50
3.89 L	10^3/ uL	4 - 10
70.2	%	40 - 80
20.6	%	20 - 40
1.0	%	1 - 6
7.7	%	2 - 10
0.5 L	%	1 - 2
2.40	lakh/cumm	1.500 - 4.500
	15.6 46.5 86.3 28.9 33.5 5.39 <b>3.89 L</b> 70.2 20.6 1.0 7.7 <b>0.5 L</b>	15.6 g/dl 46.5 % 86.3 fl 28.9 pg 33.5 g/dl 5.39 millions/cu.mm 3.89 L 10^3/ uL  70.2 % 20.6 % 1.0 % 7.7 % 0.5 L %

HAEMOGLOBIN :- Method:-SLS HemoglobinMethodology by Cell Counter.Interpretation:-Low-Anemia, High-Polycythemia.

MCV :- Method:- Calculation bysysmex.

MCH: - Method: - Calculation bysysmex.

MCHC: - Method: - Calculation bysysmex.

MCHC: - Method: - Calculation bysysmex.

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.

0 - 15

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)** 05 mm/1st hr

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Method:-Modified Westergrens. Interpretation:-Increased in infections, sepsis, and malignancy.

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Test Name Result Unit Biological Ref. Range

# **USG REPORT - ABDOMEN AND PELVIS**

### LIVER:

Is normal in size (~140 mm) and diffuse increased echogenicity. No obvious focal lesion seen. No intra hepatic biliary radical dilatation seen.

# **GALL BLADDER:**

Adequately distended with no obvious wall thickening/pericholecystic fat stranding/fluid. No obvious calculus/polyp/mass seen within.

### PANCREAS:

Appears normal in size and shows uniform echo texture. The pancreatic duct is normal. No calcifications are seen.

### SPLEEN:

Appears normal in size and it shows uniform echo texture. It measures 87 mm in long axis.

# **RIGHTKIDNEY:**

Right kidney measures 88 x 52 mm.

The shape, size and contour of the right kidney appear normal.

Corticomedullary differentiation is maintained. No evidence of pelvicalyceal dilatation.

No calculi seen.

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USG

# **LEFT KIDNEY:**

Left kidney measures 93 x 54 mm.

The shape, size and contour of the left kidney appear normal.

Corticomedullary differentiation is maintained. No evidence of pelvicalyceal dilatation.

No calculi seen.

# **URINARY BLADDER:**

Is normal in contour. No intraluminal echoes are seen. No calculus or diverticulum is seen.

# PROSTATE:

Measures 26 x 33 x 35 mm, 17cc in volume. Normal.

# **RIGHT ILIAC FOSSA:**

No focal fluid collections seen.

### **IMPRESSION:**

Diffuse grade I fatty liver.

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Patient Name Mr. BHAGIRATH SINGH Lab No 4001316

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X Ray

Test Name Result Unit Biological Ref. Range

# X-RAY - CHEST PA VIEW

### **OBSERVATION:**

The trachea is central.

The mediastinal and cardiac silhouette are normal.

Cardiothoracic ratio is normal.

Cardiophrenic and costophrenic angles are normal.

Both hila are normal.

The lung fields are clear.

Bones of the thoracic cage are normal.

Soft tissues of the chest wall are normal.

# **IMPRESSION:**

No significant abnormality seen.

\*\*End Of Report\*\*

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