Patient Name UHID Age/Gender	Mrs. SHARDA 40001135 37 Yrs/Female			Lab No Collection Date Receiving Date	4001317 11/03/2023 11:1 11/03/2023 11:2	
IP/OP Location	O-OPD			Report Date	11/03/2023 4:38	BPM
Referred By	Dr. DIWANSHU KHATANA			Report Status	Final	
Mobile No.	9588082315					
			BIOCHEMIST	RY		
Test Name		Result	Unit	Biolog	gical Ref. Range	
BLOOD GLUCOSE (F	ASTING)					Sample: Fl. Plasma
BLOOD GLUCOSE FASTING		121.2				
Method: Hexokinase Interpretation:-D various diseases.	e assay. iagnosis and monitoring of	treatment in d	iabetes mellitu	s and evaluation of o	carbohydrate metabol	ism in
BLOOD GLUCOSE (P	PP )					Sample: PLASMA
BLOOD GLUCOSE (P	Ρ)	143.3	mg/dl	Pre – Diabe	etic: - < 140 mg/dl tic: - 140-199 mg/dl ⊱=200 mg/dl	
Method: Hexokinas Interpretation:-D various diseases.	e assay. iagnosis and monitoring of	treatment in d	iabetes mellitu	s and evaluation of a	carbohydrate metabol	ism in

THYROID T3 T4 TSH				Sample: Serum
Т3	1.19	ng/mL	0.970 - 1.690	
Τ4	7.22	ug/dl	5.53 - 11.00	
TSH	1.772	μIU/mL	0.40 - 4.05	

**RESULT ENTERED BY : NEETU SHARMA** 

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Dr. MUDITA SHARMA

Patient Name	Mrs. SHARDA	Lab No	4001317
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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 acompetitive test principle with an antibody specifically directed against T4.

TSH - THYROID STIMULATING HORMONE :- ElectroChemiLuminescenceImmunoAssay - ECLIA

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

#### LFT (LIVER FUNCTION TEST)

BILIRUBIN TOTAL	0.56	mg/dl	0.00 - 1.20
BILIRUBIN INDIRECT	0.40	mg/dl	0.20 - 1.00
BILIRUBIN DIRECT	0.16	mg/dl	0.00 - 0.40
SGOT	20.0	U/L	0.0 - 40.0
SGPT	17.4	U/L	0.0 - 40.0
TOTAL PROTEIN	7.70	g/dl	6.6 - 8.7
ALBUMIN	5.27 H	g/dl	3.5 - 5.2
GLOBULIN	2.4		1.8 - 3.6
ALKALINE PHOSPHATASE	66.8	U/L	42 - 98
A/G RATIO	2.2	Ratio	1.5 - 2.5
GGTP	25.4	U/L	6.0 - 38.0

#### Sample: Serum

**RESULT ENTERED BY : NEETU SHARMA** 

Concerto to

Dr. MUDITA SHARMA

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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.

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

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	274		<200 mg/dl :- Desirable 200-240 mg/dl :- Borderline >240 mg/dl :- High
HDL CHOLESTEROL	68.3		High Risk :-<40 mg/dl (Male), <40 mg/dl (Female) Low Risk :->=60 mg/dl (Male), >=60 mg/dl (Female)
LDL CHOLESTEROL	169.1		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	19	mg/dl	10 - 50
TRIGLYCERIDES	97.0		Normal :- <150 mg/dl Border Line:- 150 - 199 mg/dl High :- 200 - 499 mg/dl Very high :- > 500 mg/dl
CHOLESTEROL/HDL RATIO	4.0	%	

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**Dr. MUDITA SHARMA** 

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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

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

UREA	29.5	mg/dl	16.60 - 48.50
BUN	13.8	mg/dl	6 - 20
CREATININE	0.49 L	mg/dl	0.50 - 0.90
SODIUM	141.1	mmol/L	136 - 145
POTASSIUM	4.65	mmol/L	3.50 - 5.50
CHLORIDE	103.7	mmol/L	98 - 107
URIC ACID	2.19 L	mg/dl	2.6 - 6.0
CALCIUM	9.27	mg/dl	8.60 - 10.30

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Dr. MUDITA SHARMA

MBBS | MD | PATHOLOGY

#### Sample: Serum

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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 andkidney 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 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.

HBA1C

5.9

%

< 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 HbAlC and mean blood glucose values during the preceding 2 to 3 months.

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**Dr. MUDITA SHARMA** 

MBBS | MD | PATHOLOGY

Sample: WHOLE BLOOD EDTA

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### **BLOOD BANK INVESTIGATION**

Test Name	Result	Unit	Biological Ref. Range
BLOOD GROUPING	"A" Rh Positive		

**BLOOD GROUPING** 

Note :

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

**RESULT ENTERED BY : NEETU SHARMA** 

Dr. MUDITA SHARMA

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### **CLINICAL PATHOLOGY**

	C			
Test Name	Result	Unit	<b>Biological Ref. Range</b>	
URINE SUGAR (POST PRANDIAL)				Sample: Urine
URINE SUGAR (POST PRANDIAL)	NEGATIVE			
URINE SUGAR (RANDOM)				Sample: Urine
URINE SUGAR (RANDOM)	NEGATIVE			
<b>ROUTINE EXAMINATION - URINE</b>				Sample: Urine
PHYSICAL EXAMINATION				
VOLUME	20	ml		
COLOUR	PALE YELLOW		P YELLOW	
APPEARANCE	CLEAR		CLEAR	
CHEMICAL EXAMINATION				
РН	6.5		5.5 - 7.0	
SPECIFIC GRAVITY	1.000		1.016-1.022	
PROTEIN	NEGATIVE		NEGATIVE	
SUGAR	NEGATIVE		NEGATIVE	
BILIRUBIN	NEGATIVE		NEGATIVE	
BLOOD	NEGATIVE			
KETONES	NEGATIVE		NEGATIVE	
NITRITE	NEGATIVE		NEGATIVE	
UROBILINOGEN	NEGATIVE		NEGATIVE	
LEUCOCYTE	NEGATIVE		NEGATIVE	
MICROSCOPIC EXAMINATION				
WBCS/HPF	1-2	/hpf	0 - 3	
RBCS/HPF	0-0	/hpf	0 - 2	
EPITHELIAL CELLS/HPF	2-3	/hpf	0 - 1	
CASTS	NIL		NIL	
CRYSTALS	NIL		NIL	

### **RESULT ENTERED BY : NEETU SHARMA**

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Dr. MUDITA SHARMA

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### **CLINICAL PATHOLOGY**

BACTERIA	NIL	NIL
OHTERS	NIL	NIL

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

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Dr. MUDITA SHARMA

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### HEMATOLOGY

Test Name	Result	Unit	Biological Ref. Ran	ge
CBC (COMPLETE BLOOD COUNT)				Sample: WHOLE BLOOD EDTA
HAEMOGLOBIN	11.4 L	g/dl	12.0 - 15.0	
PACKED CELL VOLUME(PCV)	36.9	%	36.0 - 46.0	
MCV	87.2	fl	82 - 92	
МСН	27.0	pg	27 - 32	
МСНС	30.9 L	g/dl	32 - 36	
RBC COUNT	4.23	millions/cu.mm	3.80 - 4.80	
TLC (TOTAL WBC COUNT)	7.21	10^3/ uL	4 - 10	
DIFFERENTIAL LEUCOCYTE COUNT				
NEUTROPHILS	59.5	%	40 - 80	
LYMPHOCYTE	32.6	%	20 - 40	
EOSINOPHILS	2.2	%	1 - 6	
MONOCYTES	5.3	%	2 - 10	
BASOPHIL	0.4 L	%	1 - 2	
PLATELET COUNT	2.88	lakh/cumm	1.500 - 4.500	

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.

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)

25 H

mm/1st hr 0 - 15

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**Dr. MUDITA SHARMA** 

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

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Unit

**Test Name** 

Result

**Biological Ref. Range** 

## **USG REPORT - ABDOMEN AND PELVIS**

### LIVER:

Is borderline enlarged (~155 mm) and shows diffuse increased echogenicity. No obvious focal lesion seen. No intra - Hepatic biliary radical dilatation seen.

## GALL BLADDER:

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

### PANCREAS:

Appears normal in size and it shows uniform echo texture.

## SPLEEN:

Is normal in size (~101 mm) and shows uniform echogenicity.

### **RIGHT KIDNEY:**

Right kidney measures 96 x 45 mm.

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

Corticomedullary differentiation is maintained. No evidence of pelvicalyceal dilatation.

No calculi seen.

### LEFT KIDNEY:

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Left kidney measures **108 x 56 mm**.

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

Corticomedullary differentiation is maintained. No evidence of pelvicalyceal dilatation.

No calculi seen.

## **BLADDER**:

Is normal contour. No intraluminal echoes are seen.

## **UTERUS**:

Uterus measures  $\sim$  57 x 50 x 101 mm, anteverted.

Endometrial thickness measures  $\sim$  10 mm.

## An intramural fibroid size of 16 x 15 mm is seen in posterior myometrium.

### OVARIES:

Both ovaries are normal in size and echoes.

Right ovary measures ~ 26 x 17 mm.

Left ovary measures ~ 38 x 19 mm.

## **RIGHT ILIAC FOSSA:**

No focal fluid collections seen.

### **IMPRESSION:**

Borderline hepatomegaly with diffuse grade I fatty liver.

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USG

## Small intramural uterine fibroid.

**RESULT ENTERED BY : NEETU SHARMA** 

Rendered

Dr. RENU JADIYA MBBS, DNB RADIOLOGIST

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X Ray

Unit

**Test Name** 

Result

**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\*\*

**RESULT ENTERED BY : NEETU SHARMA** 

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Dr. RENU JADIYA MBBS, DNB RADIOLOGIST