DEPARTMENT OF CARDIOLOGY

UHID / IP NO	40001604 (1792)	**RISNo./Status :**	4002093/
Patient Name :	Mrs. MANJU SHARMA	**Age/Gender :**	56 Y/F
Referred By :	EHS CONSUTANT	**Ward/Bed No :**	OPD
Bill Date/No :	17/04/2023 8:05AM/ OPSCR23-24/121	**Scan Date :**	
Report Date :	17/04/2023 10:06AM	**Company Name:**	Provisional

REFERRAL REASON: - HEALTH CHECK UP

2D ECHOCARDIOGRAPHY WITH COLOR DOPPLER

M MODE DIMENSIONS: -

		Normal			Normal
IVSD	**9.9**	**6-12mm**	**LVIDS**	**27.2**	**20-40mm**
LVIDD	**41.2**	**32-57mm**	**LVPWS**	**16.3**	**mm**
LVPWD	**10.4**	**6-12mm**	**AO**	**30.8**	**19-37mm**
IVSS	**17.2**	**mm**	**LA**	**32.2**	**19-40mm**
LVEF	**62-64**	**>55%**	**RA**	**-**	**mm**

DOPPLER MEASUREMENTS & CALCULATIONS:

STRUCTURE	MORPHOLOGY	VELOCITY (m/s)				GRADIENT (mmHg)	REGURGITATION
MITRAL VALVE	**NORMAL**	**E**	**0.88**	**e'**		**-**	**TRIVIAL MR**
		A	**0.76**	**E/e'**			
TRICUSPID VALVE	**NORMAL**	**E**		**0.57**		_	**NIL**
		A		**0.45**			
AORTIC VALVE	**NORMAL**	**1.13**				-	**NIL**
PULMONARY VALVE	**NORMAL**	**0.67**				-	**NIL**

COMMENTS & CONCLUSION: -

- NO RWMA, LVEF 62-64%
- NORMAL LV DIASTOLIC FUNCTIONS
- TRIVIAL MR, OTHER CARDIAC VALVES ARE NORMAL
- ALL CARDIAC CHAMBERS ARE NORMAL
- NO EVIDENCE OF CLOT/VEGETATION/PE
- INTACT IVS/IAS

IMPRESSION: - TRIVIAL MR, NORMAL BI VENTRICULAR FUNCTIONS

DR ROOPAM SHARMA
MBBS, PGDCC, FIAE
CONSULTANT $ INCHARGE
EMERGENCY, PREVENTIVE CARDIOLOGY AND WELLNESS CENTER.

ETERNAL HOSPITAL MEDICAL TESTING LABORATORY

Patient Name	Mrs. MANJU SHARMA	**Lab No**	4002093
UHID	40001604	**Collection Date**	17/04/2023 9:16AM
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BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range
BLOOD GLUCOSE (FASTING)			Sample: Fl. Plasma
BLOOD GLUCOSE (FASTING)	**113.6 H**	mg/dl	74 - 106

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)	145.8	mg/dl	Non – Diabetic: - < 140 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	1.250	ng/mL	0.970 - 1.690
T4	8.42	ug/dl	5.53 - 11.00
TSH	1.94	µIU/mL	0.40 - 4.05

RESULT ENTERED BY : SUNIL EHS

Dr. MUDITA SHARMA

MBBS|MD| PATHOLOGY

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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

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) Sample: Serum

BILIRUBIN TOTAL	0.58	mg/dl	0.00 - 1.20
BILIRUBIN INDIRECT	0.46	mg/dl	0.20 - 1.00
BILIRUBIN DIRECT	0.12	mg/dl	0.00 - 0.40
SGOT	30.6	U/L	0.0 - 40.0
SGPT	**42.9 H**	U/L	0.0 - 40.0
TOTAL PROTEIN	7.6	g/dl	6.6 - 8.7
ALBUMIN	4.5	g/dl	3.5 - 5.2
GLOBULIN	3.1		1.8 - 3.6
ALKALINE PHOSPHATASE	114.0	U/L	39 - 118
A/G RATIO	1.5	Ratio	1.5 - 2.5
GGTP	17.8	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 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 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	247		<200 mg/dl :- Desirable 200-240 mg/dl :- Borderline >240 mg/dl :- High
HDL CHOLESTEROL	49.7		High Risk :-<40 mg/dl (Male), <40 mg/dl (Female) Low Risk :->=60 mg/dl (Male), >=60 mg/dl (Female)
LDL CHOLESTEROL	175.6		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	**61 H**	mg/dl	10 - 50
TRIGLYCERIDES	307.3		Normal :- <150 mg/dl Border Line:- 150 - 199 mg/dl High :- 200 - 499 mg/dl Very high :- > 500 mg/dl
CHOLESTEROL/HDL RATIO	5.0	%	

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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	**12.7 L**	mg/dl	16.60 - 48.50
BUN	**5.9 L**	mg/dl	6 - 20
CREATININE	0.59	mg/dl	0.50 - 0.90
SODIUM	144.6	mmol/L	136 - 145
POTASSIUM	5.42	mmol/L	3.50 - 5.50
CHLORIDE	104.8	mmol/L	98 - 107
URIC ACID	3.77	mg/dl	2.6 - 6.0
CALCIUM	**10.47 H**	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 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.

Sample: WHOLE BLOOD EDTA

HBA1C	4.0	%	< 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
BLOOD GROUPING	"O" Rh Positive		

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	
URINE SUGAR (POST PRANDIAL)				Sample: Urine
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				
PH	6.0		5.5 - 7.0	
SPECIFIC GRAVITY	1.005		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	

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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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HEMATOLOGY

Test Name	Result	Unit	Biological Ref. Range
CBC (COMPLETE BLOOD COUNT)			Sample: WHOLE BLOOD EDTA
HAEMOGLOBIN	13.6	g/dl	12.0 - 15.0
PACKED CELL VOLUME(PCV)	42.6	%	36.0 - 46.0
MCV	84.9	fl	82 - 92
MCH	27.1	pg	27 - 32
MCHC	**31.9 L**	g/dl	32 - 36
RBC COUNT	**5.02 H**	millions/cu.mm	3.80 - 4.80
TLC (TOTAL WBC COUNT)	9.02	10^3/ uL	4 - 10
DIFFERENTIAL LEUCOCYTE COUNT			
NEUTROPHILS	60.6	%	40 - 80
LYMPHOCYTE	30.3	%	20 - 40
EOSINOPHILS	4.2	%	1 - 6
MONOCYTES	4.5	%	2 - 10
BASOPHIL	**0.4 L**	%	1 - 2
PLATELET COUNT	2.29	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.
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.

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Test Name	Result	Unit	Biological Ref. Range

USG REPORT - ABDOMEN AND PELVIS

LIVER:

Is normal in size **measure121 mm and 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 **measure 86 mm** and shows uniform echogenicity.

RIGHTKIDNEY:

Right kidney measures **86 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.

LEFTKIDNEY:

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USG

Left kidney measures **90 x 53 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 intra luminal echoes are seen.

UTERUS:

Uterus measures ~ **24 x 39 x 72 mm**, anteverted.

Endometrial thickness measures ~ **2 mm**.

No focal lesion noted.

ADNEXA:

No adnexal mass lesion is seen.

RIGHT ILIAC FOSSA:

No focal fluid collections seen.

IMPRESSION:

Diffuse grade I fatty liver.

RESULT ENTERED BY : SUNIL EHS

Dr. RENU JADIYA
MBBS, DNB
RADIOLOGIST

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USG REPORT - BOTH BREASTS

RIGHT BREAST:

Parenchyma

Skin Thickness normal

Sub cutaneous fat normal.

No ductal Dilatation.

No focal lesion seen.

Fibroglandulare echogenicity normal.

Nipple areolar complex normal.

Retromammary

Retromammary area appeared normal.

Axillary Tail

Axillary Tail: Normal.

Axillary Nodes

Few small volume lymph nodes with intact fatty hilum are seen in right axilla, largest 3 mm in short axis - unlikely to have clinical significance.

LEFT BREAST:

Parenchyma

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Skin Thickness normal

No ductal Dilatation.

No focal lesion seen.

Fibroglandulare echogenicity normal.

Nipple areolar complex normal.

Retromammary

Retromammary area appeared normal

Axillary Tail

Axillary Tail: Normal.

Axillary Nodes

Few small volume lymph nodes with intact fatty hilum are seen in right axila, largest 4 mm in short axis - unlikely to have clinical significance.

IMPRESSION:

Right breast parenchyma is normal.

Left breast parenchyma is normal.

Radiologically benign appearing bilateral axillary lymph nodes.

- **Suggested clinical correlation for further evaluation.**

BI - RADS SCORE IS: RIGHT BREAST: I LEFT BREAST : I

RESULT ENTERED BY : SUNIL EHS

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USG

NOTE: BI -RADS SCORING KEY

O – Needs additional evaluation, I – Negative, II – Benign findings, III – Probably benign

IV – Suspicious abnormality –Biopsy to be considered, V – Highly suggestive of malignancy,

VI – Known biopsy provenmalignancy.

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RADIOLOGIST

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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

RESULT ENTERED BY : SUNIL EHS

Dr. RENU JADIYA
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RADIOLOGIST