

## DEPARTMENT OF RADIO DIAGNOSIS

<b>UHID / IP NO</b>	40022110 (41653)	<b>RISNo./Status :</b>	4058009/
<b>Patient Name :</b>	Mrs. JYOTI SACHDEVA	<b>Age/Gender :</b>	43 Y/F
<b>Referred By :</b>	Dr. EHS CONSULTANT	<b>Ward/Bed No :</b>	OPD
<b>Bill Date/No :</b>	18/10/2024 9:36AM/ OPSCR24-25/24142	<b>Scan Date :</b>	
<b>Report Date :</b>	18/10/2024 10:23AM	<b>Company Name:</b>	Mediwheel - Arcofemi Health Care Ltd.

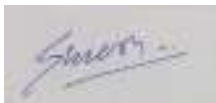
### ULTRASOUND STUDY OF WHOLE ABDOMEN

- Liver:** Normal in size & **shows increased parenchymal echotexture.** No obvious significant focal parenchymal mass lesion noted. Intrahepatic biliary radicals are not dilated. Portal vein is normal.
- Gall Bladder:** **A calculus of size approx. 15.9mm seen within lumen.** Wall thickness is normal. CBD is normal.
- Pancreas:** Normal in size & echotexture.
- Spleen:** Normal in size & echotexture. No focal lesion seen.
- Right Kidney:** Normal in shape, size & location. Echotexture is normal. Corticomedullary differentiation is maintained. No evidence of significant hydronephrosis or obstructive calculus noted.
- Left Kidney:** Normal in shape, size & location. Echotexture is normal. Corticomedullary differentiation is maintained. No evidence of significant hydronephrosis or obstructive calculus noted.
- Urinary Bladder:** Normal in size, shape & volume. No obvious calculus or mass lesion is seen. Wall thickness is normal.
- Uterus:** Not seen.  
No obvious adnexal mass seen.
- Others:** No significant free fluid is seen in pelvic peritoneal cavity.

**IMPRESSION: USG findings are suggestive of**

- Fatty liver grade – I.
- Cholelithiasis.

**Correlate clinically & with other related investigations.**



**DR. SURESH KUMAR SAINI**  
**RADIOLOGIST**  
**MBBS, MD.**  
**Reg. No. 22597, 36208.**

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### ULTRASOUND BOTH BREASTS

Both the breasts were examined by high frequency linear transducer.

Both breasts show predominant fatty component with less fibroglandular tissue.

No evidence of any focal lesion was seen on either side in the breast parenchyma in any of the quadrants.

No evidence of any ductal dilatation or pathological calcification is seen on either side.

Skin subcutaneous tissue appears normal.

Nipple areolar complex appears normal.

The axillae are unremarkable bilaterally.

#### IMPRESSION:

- **No significant sonographic abnormality noted.**

**Correlate clinically & with other related investigations.**



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**Reg. No. 22597, 36208.**

## DEPARTMENT OF CARDIOLOGY

UHID / IP NO	40022110 (41653)	RISNo./Status :	4058009/
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Referred By :	Dr. EHS CONSULTANT	Ward/Bed No :	OPD
Bill Date/No :	18/10/2024 9:36AM/ OPSCR24-25/24142	Scan Date :	
Report Date :	18/10/2024 12:26PM	Company Name:	Final

REFERRAL REASON: HTN, HEALTH CHECKUP

### 2D ECHOCARDIOGRAPHY WITH COLOR DOPPLER

#### M MODE DIMENSIONS: -

		Normal			Normal
IVSD	10.9	6-12mm	LVIDS	31.3	20-40mm
LVIDD	44.9	32-57mm	LVPWS	17.2	mm
LVPWD	10.9	6-12mm	AO	26.7	19-37mm
IVSS	17.2	mm	LA	34.4	19-40mm
LVEF	58-60	>55%	RA	-	mm

#### DOPPLER MEASUREMENTS & CALCULATIONS:

STRUCTURE	MORPHOLOGY	VELOCITY (m/s)				GRADIENT (mmHg)	REGURGITATION
		E	0.83	e'	-		
MITRAL VALVE	NORMAL	A	0.94	E/e'	-	-	NIL
		E	0.57		-		
TRICUSPID VALVE	NORMAL	A	0.48			-	NIL
		E	1.25		-		
AORTIC VALVE	NORMAL	0.73				-	NIL
PULMONARY VALVE	NORMAL	0.73				-	NIL

#### COMMENTS & CONCLUSION: -

- ALL CARDIAC CHAMBERS ARE NORMAL
- NO RWMA, LVEF 58-60%
- NORMAL LV SYSTOLIC FUNCTION
- GRADE I LV DIASTOLIC DYSFUNCTION
- ALL CARDIAC VALVES ARE NORMAL
- NO EVIDENCE OF CLOT/VEGETATION/PE
- INTACT IVS/IAS

**IMPRESSION: - GRADE I LV DIASTOLIC DYSFUNCTION, NORMAL BI VENTRICULAR SYSTOLIC FUNCTION**

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EMERGENCY, PREV.  
CARDIOLOGY(NIC) & WELLNESS  
CENTER

## ETERNAL HOSPITAL MEDICAL TESTING LABORATORY

<b>Patient Name</b>	Mrs. JYOTI SACHDEVA	<b>Lab No</b>	4058009
<b>UHID</b>	40022110	<b>Collection Date</b>	18/10/2024 9:48AM
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<b>IP/OP Location</b>	O-OPD	<b>Report Date</b>	18/10/2024 6:04PM
<b>Referred By</b>	Dr. EHS CONSULTANT	<b>Report Status</b>	Final
<b>Mobile No.</b>	9783941307		

### BIOCHEMISTRY

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

BLOOD GLUCOSE (FASTING)	93.8	mg/dl	71 - 109	
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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 )	124.8	mg/dl	Non – Diabetic: - < 140 mg/dl Pre – Diabetic: - 140-199 mg/dl Diabetic: - >=200 mg/dl	Sample: PLASMA
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Method: Hexokinase assay.

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

**THYROID T3 T4 TSH**

T3	1.180	ng/mL	0.970 - 1.690	Sample: Serum
T4	8.61	ug/dl	5.53 - 11.00	
TSH	2.08	μIU/mL	0.40 - 4.05	

RESULT ENTERED BY : SUNIL EHS



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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	0.28	mg/dl	0.00 - 1.20
BILIRUBIN INDIRECT	<b>0.16 L</b>	mg/dl	0.20 - 1.00
BILIRUBIN DIRECT	0.12	mg/dl	0.00 - 0.30
SGOT	19.5	U/L	0.0 - 32.0
SGPT	25.2	U/L	0.0 - 33.0
TOTAL PROTEIN	7.2	g/dl	6.6 - 8.7
ALBUMIN	4.5	g/dl	3.5 - 5.2
GLOBULIN	2.7		1.8 - 3.6
ALKALINE PHOSPHATASE	42	U/L	35 - 104
A/G RATIO	1.7	Ratio	1.5 - 2.5
GGTP	16.0	U/L	0.0 - 40.0

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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	143.1		<200 mg/dl :- Desirable 200-240 mg/dl :- Borderline >240 mg/dl :- High
HDL CHOLESTEROL	41.1		High Risk :- <40 mg/dl (Male), <40 mg/dl (Female) Low Risk :- >=60 mg/dl (Male), >=60 mg/dl (Female)
LDL CHOLESTEROL	84.4		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	27	mg/dl	10 - 50
TRIGLYCERIDES	136.7		Normal :- <150 mg/dl Border Line:- 150 - 199 mg/dl High :- 200 - 499 mg/dl Very high :- > 500 mg/dl
CHOLESTEROL/HDL RATIO	3	%	

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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 enzymatic 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

Sample: Serum

UREA	19.80	mg/dl	16.60 - 48.50
BUN	9	mg/dl	6 - 20
CREATININE	0.88	mg/dl	0.50 - 0.90
SODIUM	138	mmol/L	136 - 145
POTASSIUM	4.65	mmol/L	3.50 - 5.50
CHLORIDE	101.7	mmol/L	98 - 107
URIC ACID	4.1	mg/dl	2.4 - 5.7
CALCIUM	9.78	mg/dl	8.60 - 10.00

**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. 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

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### BIOCHEMISTRY

HbA1C	5.6	%			
				< 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 :** - Turbidimetric inhibition immunoassay (TINIA), **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	"A" Rh Positive		
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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
<b><u>URINE SUGAR (POST PRANDIAL)</u></b>				
URINE SUGAR (POST PRANDIAL)	NEGATIVE		NEGATIVE	Sample: Urine
<b><u>URINE SUGAR (RANDOM)</u></b>				
URINE SUGAR (RANDOM)	NEGATIVE		NEGATIVE	Sample: Urine
<b>PHYSICAL EXAMINATION</b>				
VOLUME	20	ml		Sample: Urine
COLOUR	PALE YELLOW		P YELLOW	
APPEARANCE	CLEAR		CLEAR	
<b>CHEMICAL EXAMINATION</b>				
PH	5.0 L		5.5 - 7.0	
SPECIFIC GRAVITY	1.010		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	
<b>MICROSCOPIC EXAMINATION</b>				
WBCS/HPF	1-2	/hpf	0 - 3	
RBCS/HPF	0-0	/hpf	0 - 2	
EPITHELIAL CELLS/HPF	1-2	/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 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
Sample: WHOLE BLOOD EDTA			
HAEMOGLOBIN	13.0	g/dl	12.0 - 15.0
PACKED CELL VOLUME(PCV)	40.2	%	36.0 - 46.0
MCV	86.5	fl	82 - 92
MCH	28.0	pg	27 - 32
MCHC	32.3	g/dl	32 - 36
RBC COUNT	4.65	millions/cu.mm	3.80 - 4.80
TLC (TOTAL WBC COUNT)	8.09	10 <sup>3</sup> / uL	4 - 10
<b>DIFFERENTIAL LEUCOCYTE COUNT</b>			
NEUTROPHILS	62.8	%	40 - 80
LYMPHOCYTE	27.6	%	20 - 40
EOSINOPHILS	3.3	%	1 - 6
BASOPHIL	<b>0.4 L</b>	%	1 - 2
MONOCYTES	5.9	%	2 - 10
PLATELET COUNT	3.45	lakh/cumm	1.500 - 4.500

**HAEMOGLOBIN** :- Method:-SLS Hemoglobin Methodology by Cell Counter. Interpretation:-Low-Anemia, High-Polycythemia.

**MCV** :- Method:- Calculation by sysmex.

**MCH** :- Method:- Calculation by sysmex.

**MCHC** :- Method:- Calculation by sysmex.

**RBC COUNT** :- Method:-Hydrodynamic focusing. Interpretation:-Low-Anemia, High-Polycythemia.

**TLC (TOTAL WBC COUNT)** :- Method:-Optical Detector block based on Flowcytometry. Interpretation:-High-Leucocytosis, Low-Leucopenia.

**NEUTROPHILS** :- Method: Optical detector block based on Flowcytometry

**LYMPHOCYTES** :- Method: Optical detector block based on Flowcytometry

**EOSINOPHILS** :- Method: Optical detector block based on Flowcytometry

**MONOCYTES** :- Method: Optical detector block based on Flowcytometry

**BASOPHIL** :- Method: Optical detector block based on Flowcytometry

**PLATELET COUNT** :- Method:-Hydrodynamic focusing 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)	<b>30 H</b>	mm/1st hr	0 - 15
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<b>IP/OP Location</b>	O-OPD	<b>Report Date</b>	18/10/2024 6:04PM
<b>Referred By</b>	Dr. EHS CONSULTANT	<b>Report Status</b>	Final
<b>Mobile No.</b>	9783941307		

Method:-Modified Westergrens.

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

RESULT ENTERED BY : SUNIL EHS

## ETERNAL HOSPITAL MEDICAL TESTING LABORATORY

<b>Patient Name</b>	Mrs. JYOTI SACHDEVA	<b>Lab No</b>	4058009
<b>UHID</b>	40022110	<b>Collection Date</b>	18/10/2024 9:48AM
<b>Age/Gender</b>	43 Yrs/Female	<b>Receiving Date</b>	18/10/2024 10:17AM
<b>IP/OP Location</b>	O-OPD	<b>Report Date</b>	18/10/2024 6:04PM
<b>Referred By</b>	Dr. EHS CONSULTANT	<b>Report Status</b>	Final
<b>Mobile No.</b>	9783941307		

### X Ray

Test Name	Result	Unit	Biological Ref. Range
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#### X-RAY CHEST P. A. VIEW

Both lung fields are clear.

Both CP angles are clear.

Both hemi-diaphragms are normal in shape and outlines.

Cardiac shadow is within normal limits.

Visualized bony thorax is unremarkable.

**Correlate clinically & with other related investigations.**

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

RESULT ENTERED BY : SUNIL EHS



APOORVA JETWANI

Select