



PLEASE SCAN QR CODE  
TO VERIFY THE REPORT ONLINE

Name	: MR.K RAVICHANDRAN	TID/SID	: UMR2016473/ 28325696
Age / Gender	: 55 Years / Male	Registered on	: 28-Sep-2024 / 09:31 AM
Ref.By	: ARCOFEMI HEALTH CARE LTD - MEDI WHEELS	Collected on	: 28-Sep-2024 / 09:48 AM
Req.No	: BIL4768168	Reported on	: 28-Sep-2024 / 15:39 PM
		Reference	: Arcofemi Health Care Ltd -

TEST REPORT

DEPARTMENT OF CLINICAL PATHOLOGY

Complete Urine Examination (CUE), Urine

Investigation	Observed Value	Biological Reference Intervals
<b>Physical Examination</b>		
Colour Method:Physical	Pale yellow	Straw to Yellow
Appearance Method:Physical	Clear	Clear
<b>Chemical Examination</b>		
Reaction and pH Method:pH- Methyl red & Bromothymol blue	6.0	4.6-8.0
Specific gravity Method:Bromothymol Blue	1.015	1.003-1.035
Protein Method:Tetrabromophenol blue	Negative	Negative
Glucose Method:Glucose oxidase/Peroxidase	Positive(+++)	Negative
Blood Method:Peroxidase	Negative	Negative
Ketones Method:Sodium Nitroprusside	Negative	Negative
Bilirubin Method:Dichloroanilinediazonium	Negative	Negative
Leucocytes Method:3 hydroxy5 phenylpyrrole + diazonium	Negative	Negative
Nitrites Method:Diazonium + 1,2,3,4 tetrahydrobenzo (h) quinolin 3-ol	Negative	Negative
Urobilinogen Method:Dimethyl aminobenzaldehyde	0.2	0.2-1.0 mg/dl
<b>Microscopic Examination</b>		
Pus cells (leukocytes) Method:Microscopy	0-1	2 - 3 /hpf
Epithelial cells Method:Microscopy	0-1	2 - 5 /hpf
RBC (erythrocytes) Method:Microscopy	Absent	Absent
Casts Method:Microscopy	Absent	Occasional hyaline casts may be seen



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Crystals	Absent	Phosphate, oxalate, or urate crystals may be seen
Method:Microscopy		
Others	Nil	Nil
Method:Microscopy		

**Method: Semi Quantitative test ,For CUE**

**Reference:** Godkar Clinical Diagnosis and Management by Laboratory Methods, First South Asia edition. Product kit literature.

**Interpretation:**

The complete urinalysis provides a number of measurements which look for abnormalities in the urine. Abnormal results from this test can be indicative of a number of conditions including kidney disease, urinary tract infection or elevated levels of substances which the body is trying to remove through the urine . A urinalysis test can help identify potential health problems even when a person is asymptomatic. All the abnormal results are to be correlated clinically.

\* Sample processed at Regional Reference Laboratory, Tenet Diagnostics, Bangalore

--- End Of Report ---

*Debleena Thakur*

**Dr Debleena Thakur  
Consultant Pathologist**



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Req.No : BIL4768168 Reported on : 28-Sep-2024 / 14:57 PM  
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TEST REPORT

DEPARTMENT OF HEMATOPATHOLOGY

Blood Grouping ABO And Rh Typing, EDTA Whole Blood

Parameter	Results
Blood Grouping (ABO)	B
Rh Typing (D)	POSITIVE

**Method:** Hemagglutination Tube Method by Forward & Reverse Grouping

**Reference:** Tulip kit literature

**Interpretation:** The ABO grouping and Rh typing test determines blood type grouping (A,B, AB, O ) and the Rh factor (positive or negative). A person's blood type is based on the presence or absence of certain antigens on the surface of their red blood cells and certain antibodies in the plasma. ABO antigens are poorly expressed at birth, increase gradually in strength and become fully expressed around 1 year of age.

**Note:** Records of previous blood grouping/Rh typing not available. Please verify before transfusion.

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--- End Of Report ---

*Kavya SN*

**Dr.Kavya S N**  
Consultant Pathologist



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Req.No	: BIL4768168	Reported on	: 28-Sep-2024 / 12:41 PM
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TEST REPORT

DEPARTMENT OF HEMATOPATHOLOGY

**Erythrocyte Sedimentation Rate (ESR), Whole Blood**

Investigation	Observed Value	Biological Reference Intervals
ESR 1st Hour Method:Modified Westergren	05	<=20 mm/hour

**Complete Blood Count (CBC), EDTA Whole Blood**

Investigation	Observed Value	Biological Reference Interval
Hemoglobin Method:Spectrophotometry	15.1	13.0-18.0 g/dL
Packed Cell Volume Method:Derived from Impedance	44.9	40-54 %
Red Blood Cell Count. Method:Impedance Variation	4.99	4.3-6.0 Mill/Cumm
Mean Corpuscular Volume Method:Derived from Impedance	90.0	78-100 fL
Mean Corpuscular Hemoglobin Method:Derived from Impedance	30.3	27-32 pg
Mean Corpuscular Hemoglobin Concentration Method:Derived from Impedance	33.7	31.5-36 g/dL
Red Cell Distribution Width - CV Method:Derived from Impedance	14.3	11.5-16.0 %
Red Cell Distribution Width - SD Method:Derived from Impedance	44.9	39-46 fL
Total WBC Count. Method:Impedance Variation	6290	4000-11000 cells/cumm
Neutrophils Method:Impedance Variation, Flowcytometry	56.5	40-75 %
Lymphocytes Method:Microscopy	32.2	20-45 %
Eosinophils Method:Impedance Variation,Method_Desc= Flow Cytometry	1.5	01-06 %
Monocytes Method:Impedance Variation, Flowcytometry	9.3	01-10 %
Basophils. Method:Impedance Variation,Method_Desc= Flow Cytometry	0.5	00-02 %



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Absolute Neutrophils Count. Method:Calculated	3554	1500-6600 cells/cumm
Absolute Lymphocyte Count Method:Calculated	2025	1500-3500 cells/cumm
Absolute Eosinophils count. Method:Calculated	94	40-440 cells/cumm
Absolute Monocytes Count. Method:Calculated	585	<1000 cells/cumm
Absolute Basophils count. Method:Calculated	31	<200 cells/cumm
Platelet Count. Method:Impedance Variation	3.20	1.4-4.4 lakhs/cumm
Mean Platelet Volume. Method:Derived from Impedance	7.6	7.9-13.7 fL
Plateletcrit. Method:Derived from Impedance	0.24	0.18-0.28 %

**Method:** Automated Hematology Analyzer, Microscopy

**Reference:** Dacie and Lewis Practical Hematology, 12th Edition

**Interpretation:** A Complete Blood Picture (CBP) is a screening test which can aid in the diagnosis of a variety of conditions and diseases such as anemia, leukemia, bleeding disorders and infections. This test is also useful in monitoring a person's reaction to treatment when a condition which affects blood cells has been diagnosed. All the abnormal results are to be correlated clinically.

\* Sample processed at Regional Reference Laboratory, Tenet Diagnostics, Bangalore

--- End Of Report ---

*Debleena Thakur*

**Dr Debleena Thakur**  
Consultant Pathologist



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<b>TEST REPORT</b>		Reference	: Arcofemi Health Care Ltd -

**DEPARTMENT OF CLINICAL CHEMISTRY I**

**Blood Urea Nitrogen (BUN), Serum**

Investigation	Observed Value	Biological Reference Interval
Blood Urea Nitrogen.	7	6-20 mg/dL

Method: Kinetic, Urease - GLDH, Calculated

**Interpretation:** Urea is a waste product formed in the liver when protein is metabolized. Urea is released by the liver into the blood and is carried to the kidneys, where it is filtered out of the blood and released into the urine. Since this is a continuous process, there is usually a small but stable amount of urea nitrogen in the blood. However, when the kidneys cannot filter wastes out of the blood due to disease or damage, then the level of urea in the blood will rise. The blood urea nitrogen (BUN) evaluates kidney function in a wide range of circumstances, to diagnose kidney disease, and to monitor people with acute or chronic kidney dysfunction or failure. It also may be used to evaluate a person's general health status as well.

**Reference:** Tietz Fundamentals of Clinical Chemistry and Molecular Diagnostics

**Creatinine, Serum**

Investigation	Observed Value	Biological Reference Interval
Creatinine.	0.73	0.7-1.3 mg/dL

Method: Spectrophotometry, Jaffe - IDMS Traceable

**Interpretation:**

Creatinine is a nitrogenous waste product produced by muscles from creatine. Creatinine is majorly filtered from the blood by the kidneys and released into the urine, so serum creatinine levels are usually a good indicator of kidney function. Serum creatinine is more specific and more sensitive indicator of renal function as compared to BUN because it is produced from muscle at a constant rate and its level in blood is not affected by protein catabolism or other exogenous products. It is also not reabsorbed and very little is secreted by tubules making it a reliable marker. Serum creatinine levels are increased in pre renal, renal and post renal azotemia, active acromegaly and gigantism. Decreased serum creatinine levels are seen in pregnancy and increasing age.

Biological reference interval changed; Reference: Tietz Textbook of Clinical Chemistry & Molecular Diagnostics, Fifth Edition.

**Glucose Fasting (FBS), Sodium Fluoride Plasma**

Investigation	Observed Value	Biological Reference Interval
Glucose Fasting	163	Normal: <100 mg/dL Impaired FG: 100-125 mg/dL Diabetes mellitus: >=126 mg/dL

Method: Hexokinase

**Interpretation:** It measures the Glucose levels in the blood with a prior fasting of 9-12 hours. The test helps screen a symptomatic/ asymptomatic person who is at risk for Diabetes. It is also used for regular monitoring of glucose levels in people with Diabetes.

**Reference:** American Diabetes Association. Standards of Medical Care in Diabetes-2022



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**Glucose Post Prandial (PPBS), Sodium Fluoride Plasma**

Investigation	Observed Value	Biological Reference Interval
Glucose Post Prandial Method:Hexokinase	<b>236</b>	Normal : <140 mg/dL Impaired PG: 140-199 mg/dL Diabetes mellitus: >=200 mg/dL

**Interpretation:** This test measures the blood sugar levels 2 hours after a normal meal. Abnormally high blood sugars 2 hours after a meal reflect that the body is not producing sufficient insulin which is indicative of Diabetes.

**Reference:** American Diabetes Association. Standards of Medical Care in Diabetes-2020.

**Glycosylated Hemoglobin (HbA1C), EDTA Whole Blood**

Investigation	Observed Value	Biological Reference Interval
Glycosylated Hemoglobin (HbA1c) Method:High-Performance Liquid Chromatography	<b>9.5</b>	Non-diabetic: <= 5.6 % Pre-diabetic: 5.7 - 6.4 % Diabetic: >= 6.5 %
Estimated Average Glucose (eAG) Method:High-Performance Liquid Chromatography	<b>226</b>	mg/dL

**Interpretation:** It is an index of long-term blood glucose concentrations and a measure of the risk for developing microvascular complications in patients with diabetes. Absolute risks of retinopathy and nephropathy are directly proportional to the mean HbA1c concentration. In persons without diabetes, HbA1c is directly related to risk of cardiovascular disease.

In known diabetic patients, HbA1c can be considered as a tool for monitoring the glycemic control.

- Excellent Control - 6 to 7 %,
- Fair to Good Control - 7 to 8 %,
- Unsatisfactory Control - 8 to 10 %
- and Poor Control - More than 10 %.

**Reference:** American Diabetes Association. Standards of Medical Care in Diabetes-2018.

**Bun/Creatinine Ratio, Serum**

Investigation	Observed Value
BUN/Creatinine Ratio Method:Calculated	<b>10</b>

**Reference:**

A Manual of Laboratory Diagnostic Tests. Edition 7, Lippincott Williams and Wilkins, By Frances Talaska Fischbach, RN, BSN, MSN, and Marshall Barnett Dunning 111, BS, MS, Ph.D.

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*Debleena Thakur*  
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**DEPARTMENT OF CLINICAL CHEMISTRY I**

**Lipid Profile, Serum**

Investigation	Observed Value	Biological Reference Interval
Total Cholesterol Method:Spectrophotometry , CHOD - POD	219	Desirable: < 200 mg/dL Borderline: 200-239 mg/dL High: >= 240 mg/dL
HDL Cholesterol Method:Spectrophotometry , Direct Measurement	42	Optimal : >=60 mg/dL Borderline : 40-59 mg/dL High Risk <40 mg/dL
Non HDL Cholesterol Method:Calculated	177	Optimal : <130 mg/dL Above Optimal : 130-159 mg/dL Borderline : 160-189 mg/dL High Risk : 190-219 mg/dL Very high Risk : >=220 mg/dL
LDL Cholesterol Method:Calculated	119.4	Optimum: <100 mg/dL Near/above optimum: 100-129 mg/dL Borderline: 130-159 mg/dL High: 160-189 mg/dL Very high: >=190 mg/dL
VLDL Cholesterol Method:Calculated	<b>57.60</b>	<30 mg/dL
Total Cholesterol/HDL Ratio Method:Calculated	5.21	Optimal : <3.3 Low Risk : 3.4-4.4 Average Risk : 4.5-7.1 Moderate Risk : 7.2-11.0 High Risk : >11.0
LDL/HDL Ratio Method:Calculated	2.84	Optimal : 0.5-3.0 Borderline : 3.1-6.0 High Risk : >6.0
Triglycerides Method:Spectrophotometry, Enzymatic - GPO/POD	288	Normal:<150 mg/dL Borderline: 150-199 mg/dL High: 200-499 mg/dL Very high: >=500 mg/dL mg/dl #

**Interpretation:** Lipids are fats and fat-like substances which are important constituents of cells and are rich sources of energy. A lipid profile typically includes total cholesterol, high density lipoproteins (HDL), low density lipoprotein (LDL), chylomicrons, triglycerides, very low density lipoproteins (VLDL), Cholesterol/HDL ratio .The lipid profile is used to assess the risk of developing a heart disease and to monitor its treatment. The results of the lipid profile are evaluated along with other known risk factors associated with heart disease to plan and monitor treatment. Treatment options require clinical correlation.**Reference:** Third Report of the National Cholesterol Education program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III), JAMA 2001.

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*Debleena Thakur*

**Dr Debleena Thakur**  
Consultant Pathologist



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TEST REPORT

DEPARTMENT OF CLINICAL CHEMISTRY I

Liver Function Test (LFT), Serum

Investigation	Result	Biological Reference Interval
Total Bilirubin. Method:Spectrophotometry, Diazo method	0.75	Neonates: <=15.0 mg/dL Adults: <=1.2 mg/dL
Direct Bilirubin. Method:Spectrophotometry, Diazo method	0.32	<=0.30 mg/dL
Indirect Bilirubin. Method:Calculated	0.43	Neonates: <= 14.7 mg/dL Adults: <= 1.0 mg/dL
Alanine Aminotransferase ,(ALT/SGPT) Method: IFCC without pyridoxal phosphate activation	32	<=41 U/L
Aspartate Aminotransferase,(AST/SGOT) Method: IFCC without pyridoxal phosphate activation	20	<=40 U/L
ALP (Alkaline Phosphatase). Method:Spectrophotometry , IFCC	87	40-129 U/L
Gamma GT. Method:Spectrophotometry , IFCC	39	<60 U/L
Total Protein. Method:Spectrophotometry, Biuret	7.2	6.4-8.3 g/dL
Albumin. Method:Spectrophotometry, Bromcresol Green	4.5	3.5-5.2 g/dL
Globulin. Method:Spectrophotometry, Bromcresol Green	2.70	2.0-3.5 g/dL
A/GRatio. Method:Calculated	1.67	1.1-2.5

**Interpretation:** Liver functions tests help to identify liver disease, its severity, and its type. Generally these tests are performed in combination, are abnormal in liver disease, and the pattern of abnormality is indicative of the nature of liver disease. An isolated abnormality of a single liver function test usually means a non-hepatic cause. If several liver function tests are simultaneously abnormal, then hepatic etiology is likely.

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--- End Of Report ---

*Debleena Thakur*

**Dr Debleena Thakur**  
Consultant Pathologist



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TEST REPORT

DEPARTMENT OF CLINICAL CHEMISTRY I

**Prostate Specific Antigen (PSA) Total, Serum**

Investigation	Observed Value	Biological Reference Interval
Prostate Specific Antigen (PSA) Total Method:ECLIA	0.807	0.0-4.0 ng/mL

**Interpretation:** PSA is a protein produced by cells in the prostate and is used to screen men for prostate cancer. PSA levels are elevated in Prostate cancer, and other conditions such as benign prostatic hyperplasia (BPH) and inflammation of the prostate. An elevated PSA may be followed by a biopsy and other tests like urinalysis and ultrasound to rule out urinary tract infections and for an accurate diagnosis. PSA levels are vital to determine the effectiveness of treatment and to detect recurrence in diagnosed cases of prostate cancer.

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**Dr.M.G.Satish**  
Consultant Pathologist



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TEST REPORT

DEPARTMENT OF CLINICAL CHEMISTRY I

Thyroid Profile (T3,T4,TSH), Serum

Investigation	Observed Value	Biological Reference Interval
Triiodothyronine Total (T3) Method:ECLIA	0.842	0.80-2.00 ng/mL <b>Note:</b> Biological Reference Ranges are changed due to change in method of testing.
Thyroxine Total (T4) Method:ECLIA	6.84	4.6-12.0 µg/dL
Thyroid Stimulating Hormone (TSH) Method:ECLIA	1.52	0.27-4.20 µIU/mL

**Interpretation:** A thyroid profile is used to evaluate thyroid function and/or help diagnose hypothyroidism and hyperthyroidism due to various thyroid disorders. T4 and T3 are hormones produced by the thyroid gland. They help control the rate at which the body uses energy, and are regulated by a feedback system. TSH from the pituitary gland stimulates the production and release of T4 (primarily) and T3 by the thyroid. Most of the T4 and T3 circulate in the blood bound to protein. A small percentage is free (not bound) and is the biologically active form of the hormones.

**Reference:** Tietz Fundamentals of Clinical Chemistry and Molecular Diagnostics, Carl A. Burtis, David E. Bruns.

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DEPARTMENT OF CLINICAL CHEMISTRY I

Uric Acid, Serum

Investigation	Observed Value	Biological Reference Interval
Uric Acid. Method:Enzymatic	6	3.4-7.0 mg/dL

**Interpretation:** It is the major product of purine catabolism. Hyperuricemia can result due to increased formation or decreased excretion of uric acid which can be due to several causes like metabolic disorders, psoriasis, tissue hypoxia, pre-eclampsia, alcohol, lead poisoning, acute or chronic kidney disease, etc. Hypouricemia may be seen in severe hepato cellular disease and defective renal tubular reabsorption of uric acid.

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**Dr.M.G.Satish**  
Consultant Pathologist



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- Medi Whe

**X - RAY CHEST PA VIEW**

Bilateral lung fields appear normal.

Cardiac size is within normal limits.

Bilateral hilar regions appear normal.


Bilateral domes of diaphragm and costophrenic angles are normal.

Visualised bones and soft tissues appear normal.

**IMPRESSION:**

- **No significant abnormality detected.**

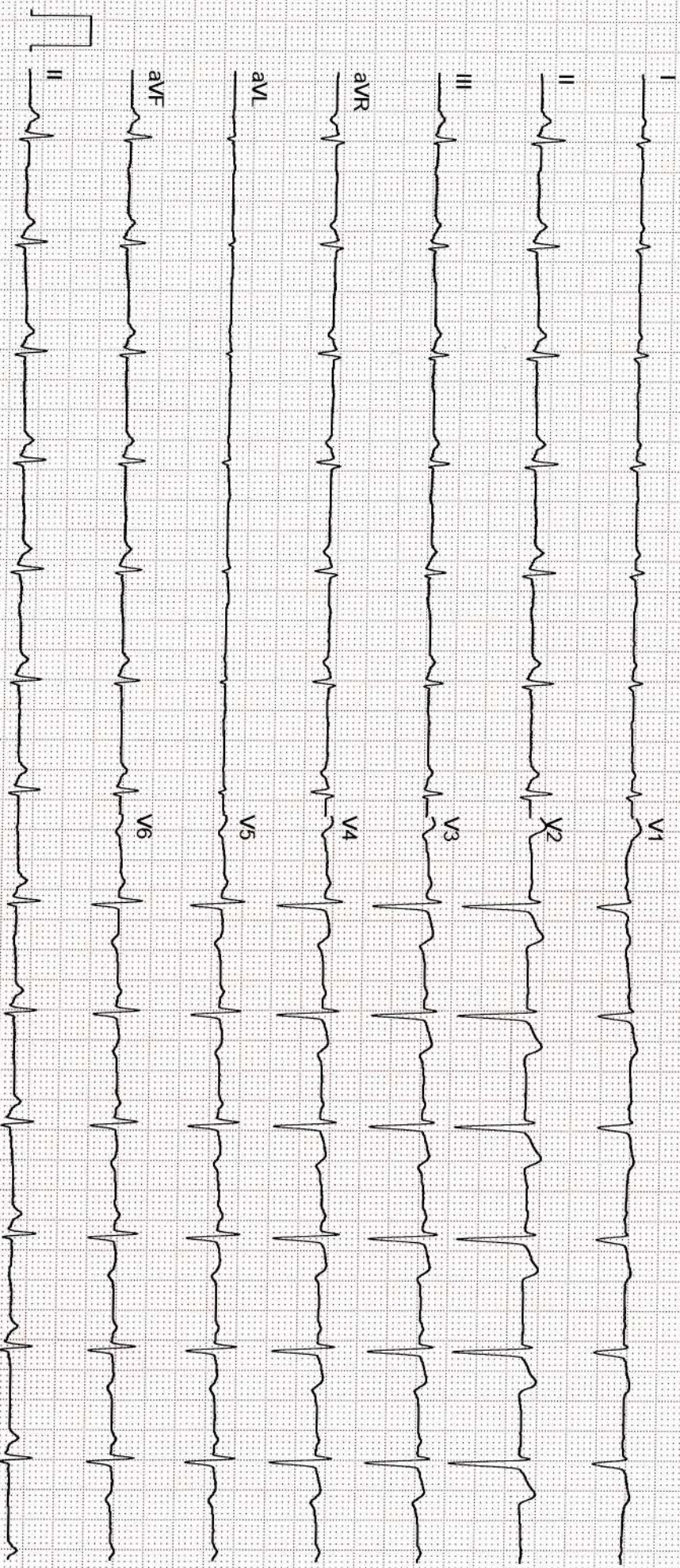
\*\*\* End Of Report \*\*\*

  
**Dr. Roohi Singh**  
Consultant Radiologist

QRS : 80 ms  
QT / QTcBaz : 380 / 441 ms  
PR : 150 ms  
P : 100 ms  
RR / PP : 738 / 740 ms  
P / QRS / T : 64 / 78 / 73 degrees

Normal sinus rhythm  
Nonspecific T wave abnormality  
Abnormal ECG

*Normal sinus rhythm,  
SVC empty catheter,  
VSI 20/10/20/7/10/10/5*







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Reg.No	: BIL4768168	Reference	: Arcofemi Health Care Ltd - Medi Whe

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**ECHOCARDIOGRAM REPORT**

**MESUREMENTS**

IVS (D):0.7 CM      LVID (D):4.2 CM      LVPW (D): 0.8CM  
IVS(S): 0.9CM      LVID (S):3.1 CM      LVPW(S): 1.0CM  
AO: 2.9 CM      LA: 2.8CM      RVID (D):2.6 CM  
EF: 60%

**VALVES:**

MITRAL VALVE                               :    NORMAL  
AORTIC VALVE                               :    NORMAL  
TRICUSPID VALVE                           :    NORMAL  
PULMONARY VALVE                         :    NORMAL

**CHAMBERS:**

LEFT ARTIUM                                :    NORMAL  
RIGHT ARTIUM                               :    NORMAL  
LEFT VENTRICLE                             :    NORMAL  
RIGHT VENTRICLE                          :    NORMAL

**SEPTAE:**

IVS   :    INTACT  
IAS   :    INTACT

**GREAT ARTERIES:**

AORTA                                        :    NORMAL  
PULMONARY ARTERY                        :    NORMAL

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Registered On : 28-Sep-2024 09:31 AM  
Reported On : 28-Sep-2024 11:23 AM  
Reference : Arcofemi Health Care Ltd  
- Medi Whe

**DOPPLER STUDY:**

MITRAL VALVE : E -0.5/ A -0.7M/S  
AORTIC VALVE : 1.1 M/S  
TRICUSPID VALVE : E -0.4/ A -0.6 M/S  
PULMONARY VALVE : 0.9 M/S

**WALL MOTION ABNORMALITIES: NO RWMA PRESENT**

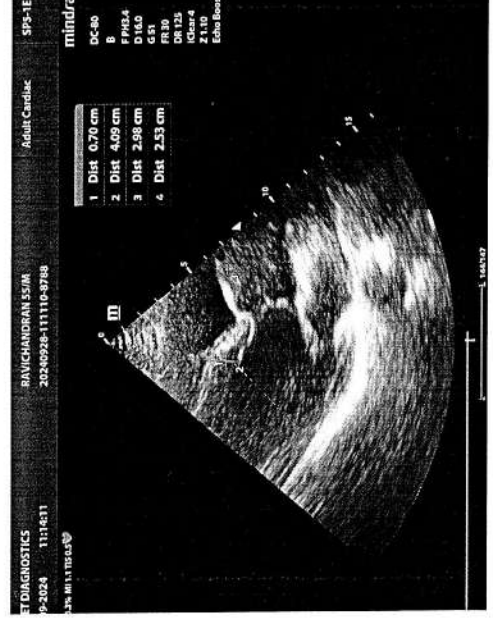
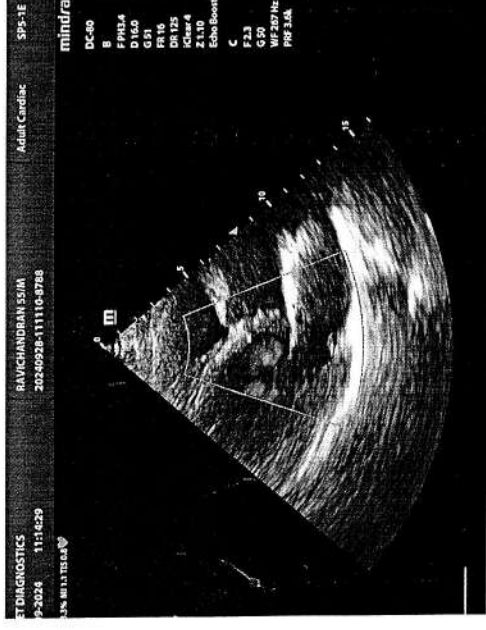
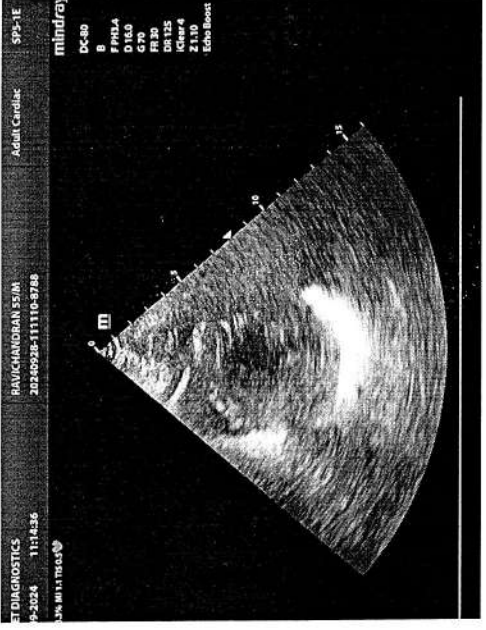
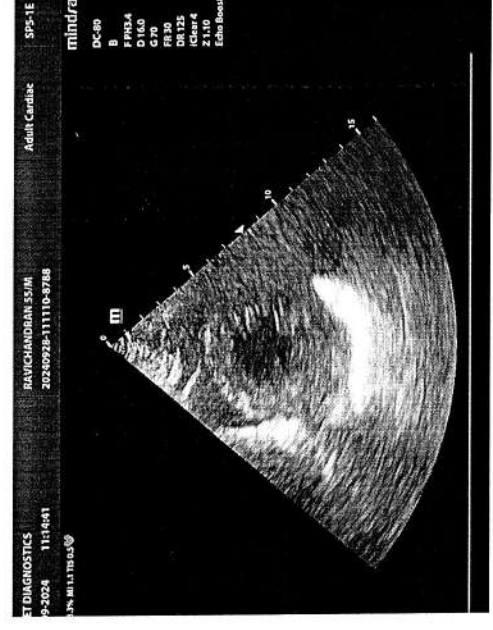
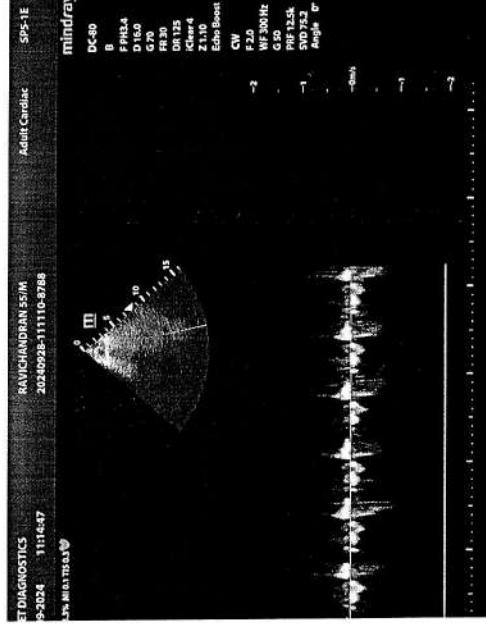
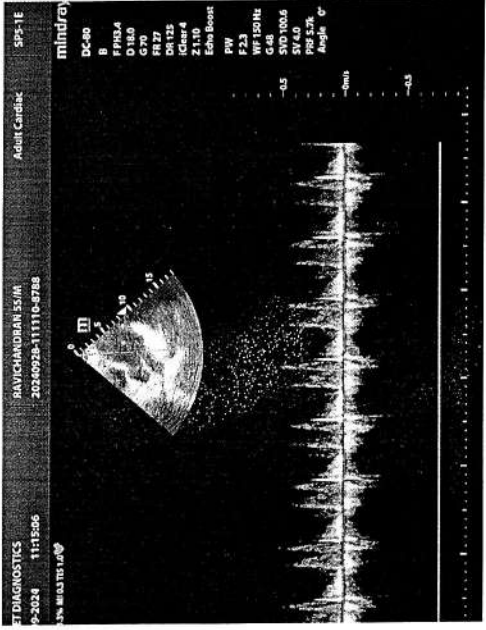
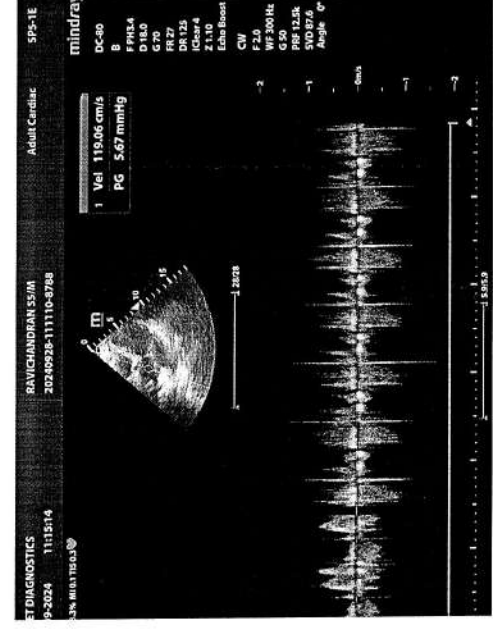
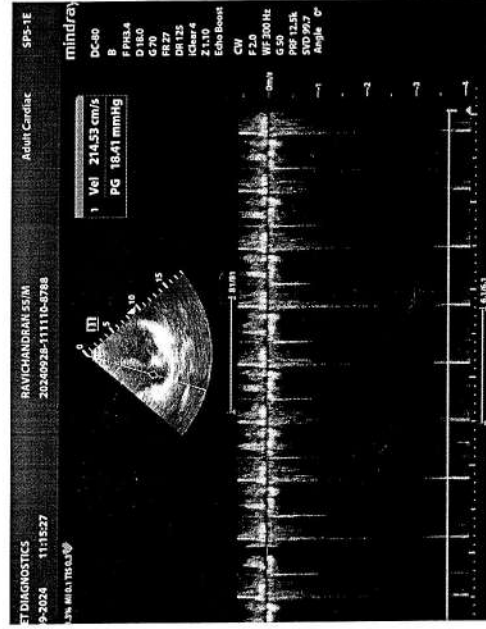
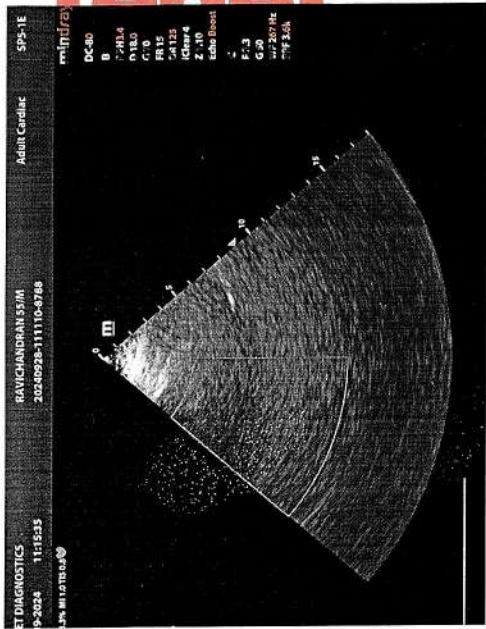
PERICARDIUM : NORMAL  
VEGETATION / THROMBUS : NO

**FINAL DIAGNOSIS:**

- NORMAL CARDIAC CHAMBERS.
- NORMAL LV SYSTOLIC FUNCTION.
- LVEF-60%.
- NO RWMA PRESENT.
- GRADE I LVDD.
- TRIVIAL MR.
- TRIVIAL TR (PASP-26 mmHg)
- NO PE / CLOT / VEGETATION SEEN.

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

**Dr.Sendil G**  
Consultant Cardiologist





Name : Mr . K RAVICHANDRAN  
Age/Gender : 55 Years/Male  
Ref By : ARCOFEMI HEALTH CARE LTD - MEDI WHEELS  
Reg.No : BIL4768168

TID : UMR2016473  
Registered On : 28-Sep-2024 09:31 AM  
Reported On : 28-Sep-2024 11:50 AM  
Reference : Arcofemi Health Care Ltd  
- Medi Whe

**ABDOMINO-PELVIC ULTRASONOGRAPHY**

**LIVER** is normal in shape, size and has increased echogenicity. No evidence of focal lesion or intrahepatic biliary ductal dilatation. Hepatic and portal vein radicals are normal.

**GALL BLADDER** is distended. No evidence of calculus or wall thickening. No pericholecystic fluid collection. CBD is of normal calibre.

**PANCREAS** has normal shape, size and uniform echopattern. No evidence of ductal dilatation or calcification.

**SPLEEN** shows normal shape, size and echopattern.

**Right kidney:** Normal in shape, size and echopattern. Cortico-medullary differentiation preserved. No evidence of calculus or hydronephrosis.

**Left kidney:** Normal in shape, size and echopattern. Cortico-medullary differentiation preserved. No evidence of calculus or hydronephrosis.

The kidney measures as follows:

	Bipolar length (cm)	Parenchymal thickness (cm)
Right Kidney	9.9	1.7
Left Kidney	10.0	2.4

**URINARY BLADDER** shows normal shape and wall thickness. It has clear contents. No evidence of diverticula.

**PROSTATE** shows normal shape, size and echopattern. It measures 3.7 x 3.9 x 3.1 cm volume: 24 cc.

No evidence of ascites.

**IMPRESSION:**

- **Grade I fatty infiltration of liver.**

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

**Dr Ramachandra C R**  
Consultant Radiologist

