



PLEASE SCAN QR CODE

Name	: Mr . B RAMANJANEYULU	TID	: UMR1964396
Age/Gender	: 38 Years/Male	Registered On	: 14-Sep-2024 10:45 AM
Ref By	: ARCOFEMI HEALTH CARE LTD - MEDI WHEELS	Reported On	: 14-Sep-2024 05:44 PM
Reg.No	: BIL4711588	Reference	: Arcofemi Health Care Ltd - Medi Whe

Clinical details: General checkup.

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 seen.

*** End Of Report ***



Dr Mahesh M S
Consultant Radiologist



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Age/Gender	: 38 Years/Male	Registered On	: 14-Sep-2024 10:45 AM
Ref By	: ARCOFEMI HEALTH CARE LTD - MEDI WHEELS	Reported On	: 14-Sep-2024 06:28 PM
Reg.No	: BIL4711588	Reference	: Arcofemi Health Care Ltd - Medi Whe

2D ECHOCARDIOGRAPHIC STUDY

M mode measurement:

AORTA	:	3.31	cms
LEFT ATRIUM	:	3.31	cms
AVS	:	1.02	cms
LEFT VENTRICLE (DIASTOLE)	:	4.25	cms
(SYSTOLE)	:	2.67	cms
VENTRICULAR SEPTUM (DIASTOLE)	:	0.95	cms
(SYSTOLE)	:	1.27	cms
POSTERIOR WALL (DIASTOLE)	:	0.91	cms
(SYSTOLE)	:	1.34	cms
EDV	:	81	ml
ESV	:	26	ml
FRACTIONAL SHORTENING	:	35	%
EJECTION FRACTION	:	65	%
EPSS	:		cms
RVID	:	1.41	cms

DOPPLER MEASUREMENTS:

MITRAL VALVE	:	'E' - 0.81 m/s	'A' - 0.47 m/s	TRIVIAL MR
AORTIC VALVE	:	1.68 m/s		NO AR
TRICUSPID VALVE	:	PASP 22mmHg		TRIVIAL TR
PULMONARY VALVE	:	0.84 m/s		NO PR



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:2:

2D ECHOCARDIOGRAPHY FINDINGS:

Left Ventricle : Normal size, Normal systolic function.
No regional wall motion abnormalities.

Left Atrium : Normal.

Right Ventricle : Normal

Right Atrium : Normal.

Mitral valve : Normal, No mitral valve prolapse.

Aortic valve : Normal, Trileaflet.

Tricuspid valve : Normal.

Pulmonary valve : Normal.

IAS : Intact.

IVS : Intact.

Pericardium : No Pericardial effusion.

IMPRESSION :

- **TRIVIAL MITRAL REGURGITATION**
- **TRIVIAL TRICUSPID REGURGITATION. PASP : 22 mmHg**
- **NORMAL SIZED CARDIAC CHAMBERS.**
- **NORMAL LV SYSTOLIC FUNCTION. EF : 65 %**
- **NO REGIONAL WALL MOTION ABNORMALITIES.**
- **NO CLOTS / PERICARDIAL EFFUSION/ VEGETATION.**

(KINDLY CORRELATE CLINICALLY AND WITH ECG)

(BRADYCARDIA OBSERVED DURING THE STUDY)



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*** End Of Report ***

Dr Sridhar L
Consultant Cardiologist



Name	: Mr . B RAMANJANEYULU	TID	: UMR1964396
Age/Gender	: 38 Years/Male	Registered On	: 14-Sep-2024 10:45 AM
Ref By	: ARCOFEMI HEALTH CARE LTD - MEDI WHEELS	Reported On	: 14-Sep-2024 11:45 AM
Reg.No	: BIL4711588	Reference	: Arcofemi Health Care Ltd - Medi Whe

*Sonographic examination was performed on **GE Voluson - S8** machine with curvilinear probe.
Clinical details: General checkup. Quality of the scan is adequate.*

ABDOMINO-PELVIC ULTRASONOGRAPHY

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

GALL BLADDER is partially distended - post prandial status. 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 (8.4cm) and echopattern. No evidence of calcifications or focal lesions. Splenic hilum is normal.

KIDNEYS: Both kidneys are normal in size, shape, location and echopattern. Cortico-medullary differentiation is well maintained. No evidence of calculus or hydronephrosis.

Right kidney measures 9.3 x 1.4 cm

Left kidney measures 9.4 x 1.5 cm

URINARY BLADDER is well distended with normal contour and wall thickness.

No evidence of calculi / diverticuli.

PROSTATE is normal in size and echotexture measuring 12 cc in volume.

No evidence of cysts / focal lesion.

No evidence of ascites.

IMPRESSION:

- **Grade I hepatic steatosis.**

*** End Of Report ***



Dr Renya B S
Consultant Radiologist



Name	: MR.B RAMANJANEYULU	TID/SID	: UMR1964396/ 28247656
Age / Gender	: 38 Years / Male	Registered on	: 14-Sep-2024 / 10:45 AM
Ref.By	: ARCOFEMI HEALTH CARE LTD - MEDI WHEELS	Collected on	: 14-Sep-2024 / 10:48 AM
Req.No	: BIL4711588	Reported on	: 14-Sep-2024 / 12:57 PM
		Reference	: Arcofemi Health Care Ltd -

TEST REPORT

DEPARTMENT OF CLINICAL PATHOLOGY

Complete Urine Examination (CUE), Urine

Investigation	Observed Value	Biological Reference Intervals
Physical Examination		
Colour Method:Physical	Pale Yellow	Straw to Yellow
Appearance Method:Physical	Clear	Clear
Chemical Examination		
Reaction and pH Method:pH- Methyl red & Bromothymol blue	6.5	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	Negative	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
Microscopic Examination		
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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TEST REPORT

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**





Name : **MR.B RAMANJANEYULU** TID/SID : UMR1964396/ 28247657
Age / Gender : 38 Years / Male Registered on : 14-Sep-2024 / 10:45 AM
Ref.By : ARCOFEMI HEALTH CARE LTD - MEDI WHEELS Collected on : 14-Sep-2024 / 10:48 AM
Req.No : BIL4711588 Reported on : 14-Sep-2024 / 13:41 PM
Reference : Arcofemi Health Care Ltd -

TEST REPORT

DEPARTMENT OF HEMATOPATHOLOGY

Blood Grouping ABO And Rh Typing, EDTA Whole Blood

Parameter	Results
Blood Grouping (ABO)	O
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.

* 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	: BIL4711588	Reported on	: 14-Sep-2024 / 11:56 AM
		Reference	: Arcofemi Health Care Ltd -

TEST REPORT

DEPARTMENT OF HEMATOPATHOLOGY

Erythrocyte Sedimentation Rate (ESR), Whole Blood

Investigation	Observed Value	Biological Reference Intervals
ESR 1st Hour Method:Modified Westergren	02	<=15 mm/hour

Complete Blood Count (CBC), EDTA Whole Blood

Investigation	Observed Value	Biological Reference Interval
Hemoglobin Method:Spectrophotometry	14.4	13.0-18.0 g/dL
Packed Cell Volume Method:Derived from Impedance	43.0	40-54 %
Red Blood Cell Count. Method:Impedance Variation	4.45	4.3-6.0 Mill/Cumm
Mean Corpuscular Volume Method:Derived from Impedance	96.7	78-100 fL
Mean Corpuscular Hemoglobin Method:Derived from Impedance	32.3	27-32 pg
Mean Corpuscular Hemoglobin Concentration Method:Derived from Impedance	33.4	31.5-36 g/dL
Red Cell Distribution Width - CV Method:Derived from Impedance	14.2	11.5-16.0 %
Red Cell Distribution Width - SD Method:Derived from Impedance	49.8	39-46 fL
Total WBC Count. Method:Impedance Variation	6790	4000-11000 cells/cumm
Neutrophils Method:Impedance Variation, Flowcytometry	50.8	40-75 %
Lymphocytes Method:Microscopy	32.8	20-45 %
Eosinophils Method:Impedance Variation,Method_Desc= Flow Cytometry	5.4	01-06 %
Monocytes Method:Impedance Variation, Flowcytometry	10.3	01-10 %
Basophils. Method:Impedance Variation,Method_Desc= Flow Cytometry	0.7	00-02 %



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TEST REPORT		Reference	: Arcofemi Health Care Ltd -

Absolute Neutrophils Count. Method:Calculated	3449	1500-6600 cells/cumm
Absolute Lymphocyte Count Method:Calculated	2227	1500-3500 cells/cumm
Absolute Eosinophils count. Method:Calculated	367	40-440 cells/cumm
Absolute Monocytes Count. Method:Calculated	699	<1000 cells/cumm
Absolute Basophils count. Method:Calculated	48	<200 cells/cumm
Platelet Count. Method:Impedance Variation	3.85	1.4-4.4 lakhs/cumm
Mean Platelet Volume. Method:Derived from Impedance	8.0	7.9-13.7 fL
Plateletcrit. Method:Derived from Impedance	0.30	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





Name	: MR.B RAMANJANEYULU	TID/SID	: UMR1964396/ 28247659F
Age / Gender	: 38 Years / Male	Registered on	: 14-Sep-2024 / 10:45 AM
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Req.No	: BIL4711588	Reported on	: 14-Sep-2024 / 12:15 PM
		Reference	: Arcofemi Health Care Ltd -

TEST REPORT

DEPARTMENT OF CLINICAL CHEMISTRY I

Blood Urea Nitrogen (BUN), Serum

Investigation	Observed Value	Biological Reference Interval
Blood Urea Nitrogen.	8	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.	0.76	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	84	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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Ref.By	: ARCOFEMI HEALTH CARE LTD - MEDI WHEELS	Collected on	: 14-Sep-2024 / 10:48 AM
Req.No	: BIL4711588	Reported on	: 14-Sep-2024 / 12:02 PM
		Reference	: Arcofemi Health Care Ltd -

TEST REPORT

Glycosylated Hemoglobin (HbA1C), EDTA Whole Blood

Investigation	Observed Value	Biological Reference Interval
Glycosylated Hemoglobin (HbA1c) Method:High-Performance Liquid Chromatography	5.7	Non-diabetic: <= 5.6 % Pre-diabetic: 5.7 - 6.4 % Diabetic: >= 6.5 %
Estimated Average Glucose (eAG) Method:High-Performance Liquid Chromatography	117	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 glycemc 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	11

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.

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

--- End Of Report ---

Debleena Thakur

**Dr Debleena Thakur
Consultant Pathologist**





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TEST REPORT

DEPARTMENT OF CLINICAL CHEMISTRY I

Lipid Profile, Serum

Investigation	Observed Value	Biological Reference Interval
Total Cholesterol Method:Spectrophotometry , CHOD - POD	196	Desirable: < 200 mg/dL Borderline: 200-239 mg/dL High: >= 240 mg/dL
HDL Cholesterol Method:Spectrophotometry , Direct Measurement	38	Optimal : >=60 mg/dL Borderline : 40-59 mg/dL High Risk <40 mg/dL
Non HDL Cholesterol Method:Calculated	158	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	138.0	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	20	<30 mg/dL
Total Cholesterol/HDL Ratio Method:Calculated	5.16	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	3.63	Optimal : 0.5-3.0 Borderline : 3.1-6.0 High Risk : >6.0
Triglycerides Method:Spectrophotometry, Enzymatic - GPO/POD	100	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.

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

--- End Of Report ---



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TEST REPORT

Debleena Thakur

Dr Debleena Thakur
Consultant Pathologist





Name : **MR.B RAMANJANEYULU** TID/SID : UMR1964396/ 28247658
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 Ref.By : ARCOFEMI HEALTH CARE LTD - MEDI WHEELS Collected on : 14-Sep-2024 / 10:48 AM
 Req.No : BIL4711588 Reported on : 14-Sep-2024 / 12:15 PM
 Reference : Arcofemi Health Care Ltd -

TEST REPORT

DEPARTMENT OF CLINICAL CHEMISTRY I

Liver Function Test (LFT), Serum

Investigation	Result	Biological Reference Interval
Total Bilirubin. Method:Spectrophotometry, Diazo method	0.70	Neonates: <=15.0 mg/dL Adults: <=1.2 mg/dL
Direct Bilirubin. Method:Spectrophotometry, Diazo method	0.34	<=0.30 mg/dL
Indirect Bilirubin. Method:Calculated	0.36	Neonates: <= 14.7 mg/dL Adults: <= 1.0 mg/dL
Alanine Aminotransferase ,(ALT/SGPT) Method: IFCC without pyridoxal phosphate activation	23	<=41 U/L
Aspartate Aminotransferase,(AST/SGOT) Method: IFCC without pyridoxal phosphate activation	20	<=40 U/L
ALP (Alkaline Phosphatase). Method:Spectrophotometry , IFCC	91	40-129 U/L
Gamma GT. Method:Spectrophotometry , IFCC	22	<60 U/L
Total Protein. Method:Spectrophotometry, Biuret	7.4	6.4-8.3 g/dL
Albumin. Method:Spectrophotometry, Bromcresol Green	4.4	3.5-5.2 g/dL
Globulin. Method:Spectrophotometry, Bromcresol Green	3	2.0-3.5 g/dL
A/GRatio. Method:Calculated	1.47	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.

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

--- End Of Report ---

Dr.M.G.Satish
Consultant Pathologist



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Req.No	: BIL4711588	Reported on	: 14-Sep-2024 / 12:25 PM
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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	1.33	0.80-2.00 ng/mL Note: Biological Reference Ranges are changed due to change in method of testing.
Thyroxine Total (T4) Method:ECLIA	9.68	4.6-12.0 µg/dL
Thyroid Stimulating Hormone (TSH) Method:ECLIA	2.58	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.

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

--- End Of Report ---

Dr.M.G.Satish
Consultant Pathologist





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DEPARTMENT OF CLINICAL CHEMISTRY I

Uric Acid, Serum

Investigation	Observed Value	Biological Reference Interval
Uric Acid. Method:Enzymatic	5.6	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.

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

--- End Of Report ---

Debleena Thakur

Dr Debleena Thakur
Consultant Pathologist



ID: 4/11388
MR RAMANJAYULU
Male 38Years
T ID : ARCOFEMI HEALTH CARE

14-09-2024 11:22:49 AM
HR : 61 bpm
P : 124 ms
PR : 160 ms
QRS : 94 ms

QT/QTcBz : 390/393 ms
P/QRS/T : 49/62/38 °
RV5/SV1 : 0.961/0.432 mV

Diagnosis Information:

HR: 61/min

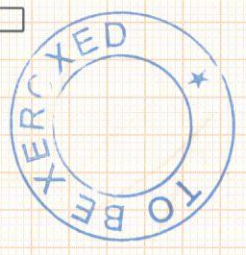
HR

Technician : SUMMMAIYA

Report Confirmed by:

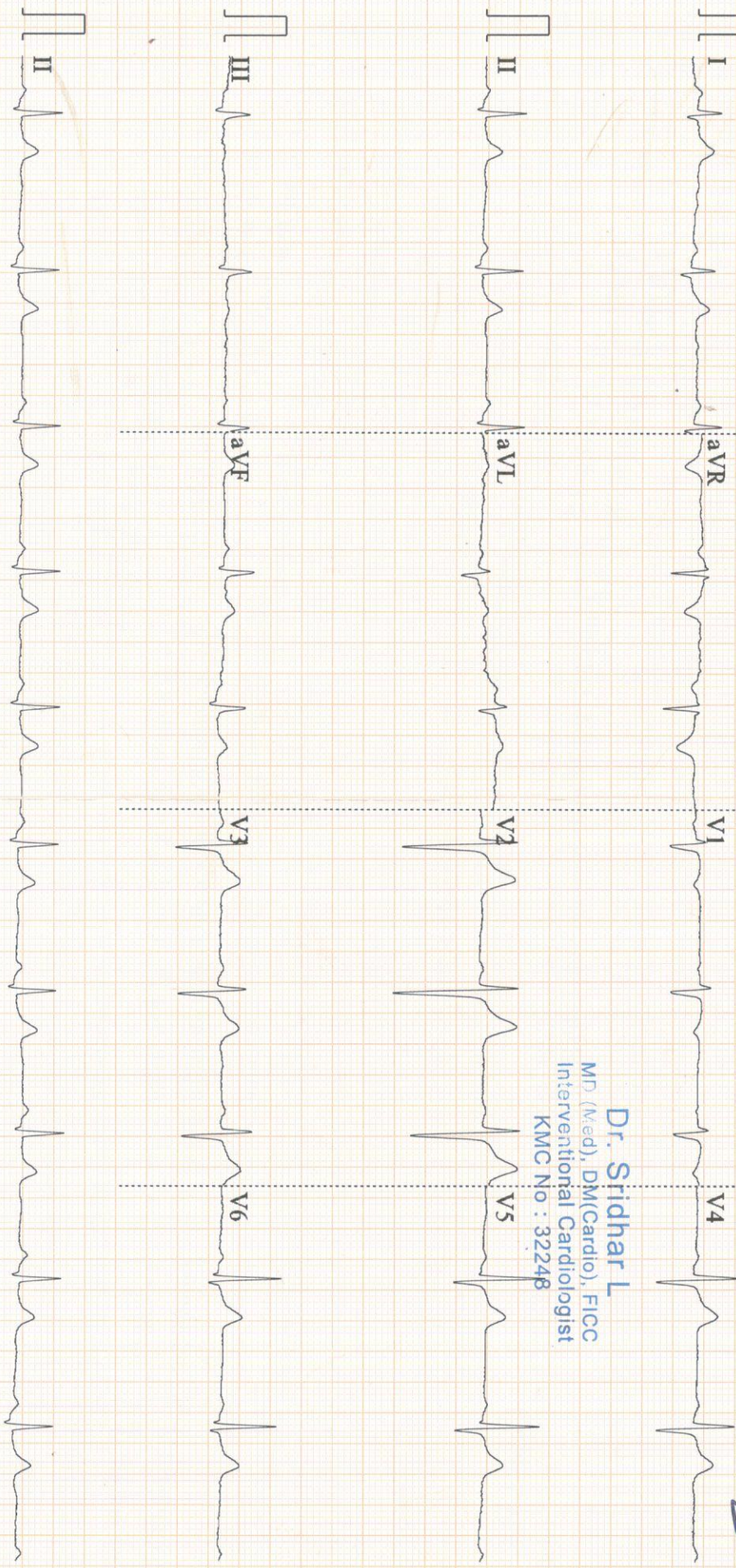
HR P 78°

Normal ECG



NEEDS CLINICAL CORRELATION
FOR FURTHER MANAGEMENT

Dr. Sridhar L
MD (Med), DM(Cardio), FICC
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बैंक ऑफ बरोडा
Bank of Baroda

नाम : श्री. रमनंजयुलु

Name : B. Ramananjayulu

E.C.No : 167131





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