



भारत सरकार



आधार

भारतीय विशिष्ट पहचान प्राधिकरण

भारत सरकार  
Unique Identification Authority of India  
Government of India

नामांकन क्रम / Enrollment No.: 1408/41522/00768

To  
अर्चना  
Archana  
W/O: Jitendra Thakur  
A-40 satya nagar  
rajputana marg Jhotwara  
Jhotwara  
Jhotwara  
Jaipur Jaipur  
Rajasthan 302012  
9530210194

02/08/2016  
383789875



MA837898756FT



आपका आधार क्रमांक / Your Aadhaar No. :

~~6515 7558~~ 8542

मेरा आधार, मेरी पहचान

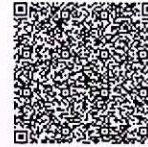


भारत सरकार

Government of India



अर्चना  
Archana  
पिता : जीतेन्द्र सिंह  
Father : Jitendra Singh  
जन्म तिथि / DOB : 10/08/1976  
महिला / Female



~~6515 7558~~ 8542

मेरा आधार मेरी पहचान

Dr. PIYUSH GOYAL  
MBBS, DMRD (Radiologist)  
RMC No.-037041



# P3 HEALTH SOLUTIONS LLP

(ASSOCIATES OF MAXCARE DIAGNOSTICS)

- B-14, Vidhyadhar Enclave-II, Near Axis Bank  
Central Spine, Vidhyadhar Nagar, Jaipur - 302023
- +91 141 4824885 ✉ maxcarediagnostics1@gmail.com



## General Physical Examination

Date of Examination: 13/04/2024

Name: ARCHANA Age: 47 YRS DOB: 10/08/1976 Sex: Female

Referred By: GANESH BARODA

Photo ID: ANNA BARODA ID #: 8540

Ht: 163 (cm)

Wt: 73 (Kg)

Chest (Expiration): 100 (cm)

Abdomen Circumference: 100 (cm)

Blood Pressure: 100/80 mm Hg

PR: 78/min

RR: 18/min

Temp: Afebrile

BMI 27.5

Eye Examination: R/E - GIG, NIG, NCB  
L/E - GIG, NIG, NCB

Other: NO

On examination he/she appears physically and mentally fit: Yes/No

Signature Of Examinee : [Signature] Name of Examinee: ARCHANA

Signature Medical Examiner : [Signature] Name Medical Examiner DR. PIYUSH GOYAL

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RMC No.-037041





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**Patient ID** 122476 Patient Mob No.9462962318  
**NAME** Mrs. ARCHANA  
Age 47 Yrs Sex F Marital Status Married  
Ref. By BANK OF BARODA  
Lab/Hosp Mr.MEDIWHEEL

Registered On 13/04/2024 08:48:49  
Collected On 13/04/2024 10:18:38  
Authorized On 13/04/2024 17:16:28  
Printed On 13/04/2024 17:16:34

## HAEMOGARAM

## HAEMATOTOLOGY

Test Name	Value	Unit	Biological Ref Interval
FULL BODY HEALTH CHECKUP ABOVE 40FEMALE			
<b>HAEMOGLOBIN (Hb)</b>	10.1 L	g/dL	12.0 - 15.0
<b>TOTAL LEUCOCYTE COUNT</b>	5.90	/cumm	4.00 - 10.00
<b>DIFFERENTIAL LEUCOCYTE COUNT</b>			
NEUTROPHIL	56.0	%	40.0 - 80.0
LYMPHOCYTE	39.0	%	20.0 - 40.0
EOSINOPHIL	2.0	%	1.0 - 6.0
MONOCYTE	3.0	%	2.0 - 10.0
BASOPHIL	0.0	%	0.0 - 2.0
TOTAL RED BLOOD CELL COUNT (RBC)	4.52	$\times 10^6/\mu\text{L}$	3.80 - 4.80
HEMATOCRIT (HCT)	32.90 L	%	36.00 - 46.00
MEAN CORP VOLUME (MCV)	73.0 L	fL	83.0 - 101.0
MEAN CORP HB (MCH)	22.2 L	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	30.5 L	g/dL	31.5 - 34.5
<b>PLATELET COUNT</b>	324	$\times 10^3/\mu\text{L}$	150 - 410
RDW-CV	16.2 H	%	11.6 - 14.0

Technologist  
Page No. 1 of 16

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MD (Pathology)  
RMC No. 17226



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## HAEMATOLOGY

### HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
<b>Erythrocyte Sedimentation Rate (ESR)</b> Method:- Westergreen	19	mm in 1st hr	00 - 20

The erythrocyte sedimentation rate (ESR or sed rate) is a relatively simple, inexpensive, non-specific test that has been used for many years to help detect inflammation associated with conditions such as infections, cancers, and autoimmune diseases. ESR is said to be a non-specific test because an elevated result often indicates the presence of inflammation but does not tell the health practitioner exactly where the inflammation is in the body or what is causing it. An ESR can be affected by other conditions besides inflammation. For this reason, the ESR is typically used in conjunction with other tests, such as C-reactive protein. ESR is used to help diagnose certain specific inflammatory diseases, including temporal arteritis, systemic vasculitis and polymyalgia rheumatica. (For more on these, read the article on Vasculitis.) A significantly elevated ESR is one of the main test results used to support the diagnosis. This test may also be used to monitor disease activity and response to therapy in both of the above diseases as well as



Technologist  
Page No. 2 of 16

*Tanu*

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(CBC): Methodology: TLC,DLC Fluorescent Flow cytometry, HB SLS method,TRBC,PCV,PLT Hydrodynamically focused Impedance, and MCH,MCV,MCHC,MENTZER INDEX are calculated. InstrumentName: Sysmex 6 part fully automatic analyzer XN-L,Japan





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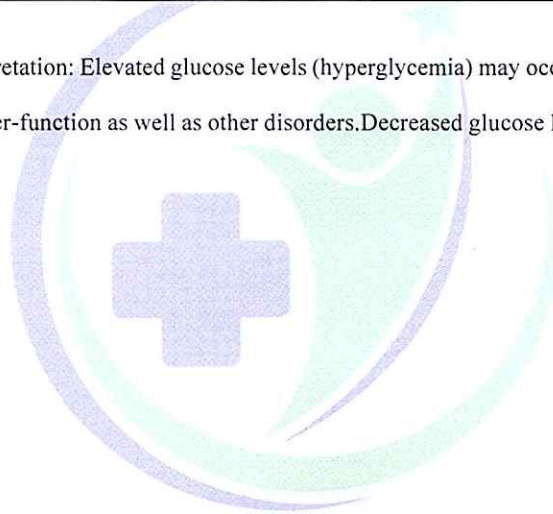


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## BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
FASTING BLOOD SUGAR (Plasma) Method:- GLUCOSE OXIDASE/PEROXIDASE	146.0 H	mg/dl	70.0 - 115.0
Impaired glucose tolerance (IGT)		111 - 125 mg/dL	
Diabetes Mellitus (DM)		> 126 mg/dL	

Instrument Name: HORIBA CA60 Interpretation: Elevated glucose levels (hyperglycemia) may occur with diabetes, pancreatic neoplasm, hyperthyroidism and adrenal cortical hyper-function as well as other disorders. Decreased glucose levels (hypoglycemia) may result from excessive insulin therapy or various liver diseases .



**Technologist**  
Page No. 4 of 16

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## HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
<b>GLYCOSYLATED HEMOGLOBIN (HbA1C)</b> Method:- CAPILLARY with EDTA	6.7	mg%	Non-Diabetic < 6.0 Good Control 6.0-7.0 Weak Control 7.0-8.0 Poor control > 8.0
<b>MEAN PLASMA GLUCOSE</b> Method:- Calculated Parameter	146 H	mg/dL	68 - 125

### INTERPRETATION

AS PER AMERICAN DIABETES ASSOCIATION (ADA)

Reference Group HbA1c in %

Non diabetic adults >=18 years < 5.7

At risk (Prediabetes) 5.7 - 6.4

Diagnosing Diabetes >= 6.5

### CLINICAL NOTES

In vitro quantitative determination of HbA1c in whole blood is utilized in long term monitoring of glycemia. The HbA1c level correlates with the mean glucose concentration prevailing in the course of the patient's recent history (approx - 6-8 weeks) and therefore provides much more reliable information for glycemia monitoring than do determinations of blood glucose or urinary glucose. It is recommended that the determination of HbA1c be performed at intervals of 4-6 weeks during Diabetes Mellitus therapy. Results of HbA1c should be assessed in conjunction with the patient's medical history, clinical examinations and other findings.

Some of the factors that influence HbA1c and its measurement [Adapted from Gallagher et al]

#### 1. Erythropoiesis

- Increased HbA1c: iron, vitamin B12 deficiency, decreased erythropoiesis.
- Decreased HbA1c: administration of erythropoietin, iron, vitamin B12, reticulocytosis, chronic liver disease.

#### 2. Altered Haemoglobin-Genetic or chemical alterations in hemoglobin: hemoglobinopathies, HbF, methemoglobin, may increase or decrease HbA1c.

#### 3. Glycation

- Increased HbA1c: alcoholism, chronic renal failure, decreased intraerythrocytic pH.
- Decreased HbA1c: certain hemoglobinopathies, increased intra-erythrocyte pH

#### 4. Erythrocyte destruction

- Increased HbA1c: increased erythrocyte life span: Splenectomy.
- Decreased A1c: decreased RBC life span: hemoglobinopathies, splenomegaly, rheumatoid arthritis or drugs such as antiretrovirals, ribavirin & dapsone.

#### 5. Others

- Increased HbA1c: hyperbilirubinemia, carbamylated hemoglobin, alcoholism, large doses of aspirin, chronic opiate use, chronic renal failure
- Decreased HbA1c: hypertriglyceridemia, reticulocytosis, chronic liver disease, aspirin, vitamin C and E, splenomegaly, rheumatoid arthritis or drugs

Technologist  
Page No: 5 of 16

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## HAEMATOLOGY

### HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
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BLOOD GROUP ABO  
 Method:- Haemagglutination reaction

"B" POSITIVE



Technologist  
Page No. 6 of 16

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MD (Pathology)  
RMC No. 17226





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## BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
<b>LIPID PROFILE</b>			
SERUM TOTAL CHOLESTEROL Method:- CHOLESTEROL OXIDASE/PEROXIDASE	132.00	mg/dl	Desirable <200 Borderline 200-239 High > 240
<b>InstrumentName:</b> HORIBA <b>Interpretation:</b> Cholesterol measurements are used in the diagnosis and treatments of lipid lipoprotein metabolism disorders.			
SERUM TRIGLYCERIDES Method:- GLYCEROL PHOSPHATE OXIDASE/PREOXIDASE	81.60	mg/dl	Normal <150 Borderline high 150-199 High 200-499 Very high >500
<b>InstrumentName:</b> Ranox Rx Imola <b>Interpretation :</b> Triglyceride measurements are used in the diagnosis and treatment of diseases involving lipid metabolism and various endocrine disorders e.g. diabetes mellitus, nephrosis and liver obstruction.			
DIRECT HDL CHOLESTEROL Method:- Direct clearance Method	37.90	mg/dl	MALE- 30-70 FEMALE - 30-85
<b>Instrument Name:</b> Rx Daytona plus <b>Interpretation:</b> An inverse relationship between HDL-cholesterol (HDL-C) levels in serum and the incidence/prevalence of coronary heart disease (CHD) has been demonstrated in a number of epidemiological studies. Accurate measurement of HDL-C is of vital importance when assessing patient risk from CHD. Direct measurement gives improved accuracy and reproducibility when compared to precipitation methods.			
LDL CHOLESTEROL Method:- Calculated Method	80.50	mg/dl	Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190
VLDL CHOLESTEROL Method:- Calculated	16.32	mg/dl	0.00 - 80.00
T.CHOLESTEROL/HDL CHOLESTEROL RATIO Method:- Calculated	3.48		0.00 - 4.90
LDL / HDL CHOLESTEROL RATIO Method:- Calculated	2.12		0.00 - 3.50
TOTAL LIPID Method:- CALCULATED	398.68	L mg/dl	400.00 - 1000.00

**Technologist**  
Page No. 7 of 16

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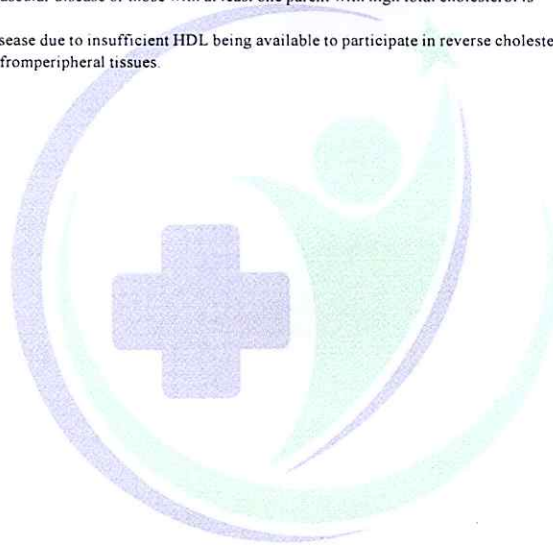
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## BIOCHEMISTRY

### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
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1. Measurements in the same patient can show physiological & analytical variations. Three serial samples 1 week apart are recommended for Total Cholesterol, Triglycerides, HDL & LDL Cholesterol.
2. As per NCEP guidelines, all adults above the age of 20 years should be screened for lipid status. Selective screening of children above the age of 2 years with a family history of premature cardiovascular disease or those with at least one parent with high total cholesterol is recommended.
3. Low HDL levels are associated with Coronary Heart Disease due to insufficient HDL being available to participate in reverse cholesterol transport, the process by which cholesterol is eliminated from peripheral tissues.



**Technologist**  
Page No. 8 of 16

*Tanu*

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## BIOCHEMISTRY

### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
<b>LIVER PROFILE WITH GGT</b>			
SERUM BILIRUBIN (TOTAL) Method:- DIAZOTIZED SULFANILIC	0.56	mg/dL	Infants : 0.2-8.0 mg/dL Adult - Up to - 1.2 mg/dL
SERUM BILIRUBIN (DIRECT) Method:- DIAZOTIZED SULFANILIC	0.20	mg/dL	Up to 0.40 mg/dL
SERUM BILIRUBIN (INDIRECT) Method:- Calculated	0.36	mg/dl	0.30-0.70
SGOT Method:- IFCC	26.1	U/L	0.0 - 40.0
SGPT Method:- IFCC	33.2	U/L	0.0 - 35.0
SERUM ALKALINE PHOSPHATASE Method:- IFCC	74.20	IU/L	53.00 - 141.00
SERUM GAMMA GT Method:- Szasz methodology Instrument Name Randox Rx Imola Interpretation: Elevations in GGT levels are seen earlier and more pronounced than those with other liver enzymes in cases of obstructive jaundice and metastatic neoplasms. It may reach 5 to 30 times normal levels in intra-or post-hepatic biliary obstruction. Only moderate elevations in the enzyme level (2 to 5 times normal) are observed with infectious hepatitis.	20.20	U/L	5.00 - 32.00
SERUM TOTAL PROTEIN Method:- BIURET	6.58	g/dl	6.00 - 8.40
SERUM ALBUMIN Method:- BROMOCRESOL GREEN	4.23	g/dl	3.50 - 5.50
SERUM GLOBULIN Method:- CALCULATION	2.35	gm/dl	2.20 - 3.50
A/G RATIO	1.80		1.30 - 2.50

**Interpretation :** Measurements obtained by this method are used in the diagnosis and treatment of a variety of diseases involving the liver, kidney and bone marrow as well as other metabolic or nutritional disorders.

Note :- These are group of tests that can be used to detect the presence of liver disease, distinguish among different types of liver disorders, gauge the extent of known liver damage, and monitor the response to treatment. Most liver diseases cause only mild symptoms initially, but these diseases must be detected early. Some tests are associated with functionality (e.g.,

**Technologist**  
Page No. 9 of 16

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MD (Pathology)  
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## BIOCHEMISTRY

### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
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albumin), some with cellular integrity (e.g., transaminase), and some with conditions linked to the biliary tract (gamma-glutamyl transferase and alkaline phosphatase). Conditions with elevated levels of ALT and AST include hepatitis A,B ,C ,paracetamol toxicity etc. Several biochemical tests are useful in the evaluation and management of patients with hepatic dysfunction. Some or all of these measurements are also carried out (usually about twice a year for routine cases) on those individuals taking certain medications, such as anticonvulsants, to ensure that the medications are not adversely impacting the person's liver.



Technologist  
Page No. 10 of 16

*Tanu*

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## BIOCHEMISTRY

### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
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#### RFT / KFT WITH ELECTROLYTES

SERUM UREA Method:- UREASE / GLUTAMATE DEHYDROGENASE	32.30	mg/dl	10.00 - 50.00
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InstrumentName: HORIBA CA 60 Interpretation : Urea measurements are used in the diagnosis and treatment of certain renal and metabolic discases.

SERUM CREATININE Method:- JAFFE	0.88	mg/dl	Males : 0.6-1.50 mg/dl Females : 0.6 -1.40 mg/dl
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Interpretation : Creatinine is measured primarily to assess kidney function and has certain advantages over the measurement of urea. The plasma level of creatinine is relatively independent of protein ingestion, water intake, rate of urine production and exercise. Depressed levels of plasma creatinine are rare and not clinically significant.

SERUM URIC ACID Method:- URICASE/PEROXIDASE	4.25	mg/dl	2.40 - 7.00
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InstrumentName:HORIBA YUMIZEN CA60 Daytona plus Interpretation: Elevated Urate:High purine diet,Alcohol• Renal insufficiency,Drugs , Polycythaemia vera, Malignancies,Hypothyroidism,Rare enzyme defects ,Downs syndrome,Metabolic syndrome, Pregnancy,Gout.

SODIUM Method:- ISE	138.9	mmol/L	135.0 - 150.0
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POTASSIUM Method:- ISE	4.60	mmol/L	3.50 - 5.50
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CHLORIDE Method:- ISE	102.3	mmol/L	94.0 - 110.0
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SERUM CALCIUM Method:- Arsenazo III Method	9.25	mg/dL	8.80 - 10.20
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InstrumentName:MISPA PLUS Interpretation: Serum calcium levels are believed to be controlled by parathyroid hormone and vitamin D. Increases in serum PTH or vitamin D are usually associated with hypercalcemia .Hypocalcemia may be observed in hypoparathyroidism, nephrosis and pancreatitis.

SERUM TOTAL PROTEIN Method:- BIURET	6.58	g/dl	6.00 - 8.40
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SERUM ALBUMIN Method:- BROMOCRESOL GREEN	4.23	g/dl	3.50 - 5.50
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Technologist  
Page No. 11 of 16

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Age 47 Yrs	Sex M	Authorized On	13/04/2024 17:16:28
Ref. By BANK OF BARODA		Printed On	13/04/2024 17:16:34
Lab/Hosp Mr.MEDIWHEEL			

## BIOCHEMISTRY

### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
SERUM GLOBULIN Method:- CALCULATION	2.35	gm/dl	2.20 - 3.50
A/G RATIO	1.80		1.30 - 2.50

**Interpretation :** Measurements obtained by this method are used in the diagnosis and treatment of a variety of diseases involving the liver, kidney and bone marrow as well as other metabolic or nutritional disorders.

#### INTERPRETATION

Kidney function tests are group of tests that can be used to evaluate how well the kidneys are functioning. Creatinine is a waste product that comes from protein in the diet and also comes from the normal wear and tear of muscles of the body. In blood, it is a marker of GFR. In urine, it can remove the need for 24-hour collections for many analytes or be used as a quality assurance tool to assess the accuracy of a 24-hour collection. Higher levels may be a sign that the kidneys are not working properly. As kidney disease progresses, the level of creatinine and urea in the blood increases. Certain drugs are nephrotoxic hence KFT is done before and after initiation of treatment with these drugs.

Low serum creatinine values are rare, they almost always reflect low muscle mass.

Apart from renal failure Blood Urea can increase in dehydration and GI bleed

Technologist  
Page No. 12 of 16

**DR. TANU RUNGTA**  
MD (Pathology)  
RMC No. 17226





# P3 HEALTH SOLUTIONS LLP

(ASSOCIATES OF MAXCARE DIAGNOSTICS)

📍 B-14, Vidhyadhar Enclave-II, Near Axix Bank  
Central Spine, Vidhyadhar Nagar, Jaipur - 302023  
☎️ +91 141 4824885 📧 maxcarediagnostics1@gmail.com



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## CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
<b>Urine Routine</b>			
<b><u>PHYSICAL EXAMINATION</u></b>			
COLOUR	PALE YELLOW		PALE YELLOW
APPEARANCE	Clear		Clear
<b><u>CHEMICAL EXAMINATION</u></b>			
REACTION(PH)	6.5		5.0 - 7.5
SPECIFIC GRAVITY	1.010		1.010 - 1.030
PROTEIN	NIL		NIL
SUGAR	NIL		NIL
BILIRUBIN	NEGATIVE		NEGATIVE
UROBILINOGEN	NORMAL		NORMAL
KETONES	NEGATIVE		NEGATIVE
NITRITE	NEGATIVE		NEGATIVE
<b><u>MICROSCOPY EXAMINATION</u></b>			
RBC/HPF	NIL	/HPF	NIL
WBC/HPF	2-3	/HPF	2-3
EPITHELIAL CELLS	2-3	/HPF	2-3
CRYSTALS/HPF	ABSENT		ABSENT
CAST/HPF	ABSENT		ABSENT
AMORPHOUS SEDIMENT	ABSENT		ABSENT
BACTERIAL FLORA	ABSENT		ABSENT
YEAST CELL	ABSENT		ABSENT
OTHER	ABSENT		ABSENT

**Technologist**  
Page No. 13 of 16

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MD (Pathology)  
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Lab/Hosp Mr.MEDIWHEEL			

## CLINICAL PATHOLOGY

## CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
URINE SUGAR (FASTING) Collected Sample Received	Nil		Nil



**Technologist**  
Page No. 14 of 16

**DR. TANU RUNGTA**  
MD (Pathology)  
RMC No. 17226





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<b>Patient ID</b> 122476	Patient Mob No.9462962318	Registered On	13/04/2024 08:48:49
<b>NAME</b> Mrs. ARCHANA		Collected On	13/04/2024 10:18:38
Age 47 Yrs	Sex M on 3 Feb 2024	Authorized On	13/04/2024 17:16:28
Ref. By BANK OF BARODA		Printed On	13/04/2024 17:16:34
Lab/Hosp Mr.MEDIWHEEL			

## IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
<b>TOTAL THYROID PROFILE</b>			
<b>THYROID-TRIiodothyronine T3</b> Method:- ECLIA	0.88	ng/mL	0.70 - 2.04
<b>THYROID - THYROXINE (T4)</b> Method:- ECLIA	7.46	ug/dl	5.10 - 14.10
<b>TSH</b> Method:- ECLIA	2.395	μIU/mL	0.350 - 5.500

4th Generation Assay, Reference ranges vary between laboratories

### PREGNANCY - REFERENCE RANGE for TSH IN uIU/mL (As per American Thyroid Association)

- 1st Trimester : 0.10-2.50 uIU/mL
- 2nd Trimester : 0.20-3.00 uIU/mL
- 3rd Trimester : 0.30-3.00 uIU/mL

The production, circulation, and disintegration of thyroid hormones are altered throughout the stages of pregnancy.

NOTE-TSH levels are subject to circadian variation, reaching peak levels between 2-4 AM and min between 6-10 PM. The variation is the order of 50% hence time of the day has influence on the measures serum TSH concentration. Dose and time of drug intake also influence the test result.

### INTERPRETATION

- 1.Primary hyperthyroidism is accompanied by ↑serum T3 & T4 values along with ↓ TSH level.
- 2.Primary hypothyroidism is accompanied by ↓ serum T3 and T4 values & ↑serum TSH levels
- 3.Normal T4 levels accompanied by ↑ T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis
- 4.Normal or ↓ T3 & ↑T4 levels indicate T4 Thyrotoxicosis ( problem is conversion of T4 to T3)
- 5.Normal T3 & T4 along with ↓ TSH indicate mild / Subclinical Hyperthyroidism

. **COMMENTS:** Assay results should be interpreted in context to the clinical condition and associated results of other investigations. Previous treatment with corticosteroid therapy may result in lower TSH levels while thyroid hormone levels are normal. Results are invalidated if the client has undergone a radionuclide scan within 7-14 days before the test.

. **Disclaimer:** TSH is an important marker for the diagnosis of thyroid dysfunction. Recent studies have shown that the TSH distribution progressively shifts to a higher concentration with age, and it is debatable whether this is due to a real change with age or an increasing proportion of unrecognized thyroid disease in the elderly

. **Reference ranges are from Teltz fundamental of clinical chemistry 8th ed (2018)**

Test performed by Instrument : Beckman coulter Dxi 800

**Note :** The result obtained relate only to the sample given/ received & tested. A single test result is not always indicative of a disease, it has to be correlated with

\*\*\* End of Report \*\*\*

**DR. TANU RUNGTA**  
MD (Pathology)  
RMC No. 17226

**Technologist**  
Page No: 16 of 16



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☎ +91 141 4824885 📧 maxcarediagnostics1@gmail.com



MRS. ARCHANA	Age : 47 Y/F
Registration Date: 13/04/2024	Ref. by: BANK OF BARODA

## Ultrasonography report: Breast and Axilla

### Right breast:-

Skin, subcutaneous tissue and retroareolar region is normal.  
Fibro glandular tissue shows normal architecture and echotexture.  
Pre and retro mammary regions are unremarkable.  
No obvious cyst, mass or architectural distortion visualized.  
Axillary lymph nodes are not significantly enlarged and their hilar shadows are preserved.

### Left breast: -

Skin, subcutaneous tissue and retroareolar region is normal.  
Fibro glandular tissue shows normal architecture and echotexture.  
Pre and retro mammary regions are unremarkable.  
No obvious cyst, mass or architectural distortion visualized.  
Axillary lymph nodes are not significantly enlarged and their hilar shadows are preserved.

**IMPRESSION: No significant abnormality is detected.**

DR.SHALINI GOEL  
M.B.B.S, D.N.B (Radiodiagnosis)  
RMC no.: 21954

**Dr. SHALINI GOEL**  
**MBBS, DNB (Radiologist)**  
**RMC No. 21954**  
**P-3 Health Solutions LLP**





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- B-14, Vidhyadhar Enclave-II, Near Axix Bank  
Central Spine, Vidhyadhar Nagar, Jaipur - 302023
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MRS. ARCHANA	Age : 47 Y/F
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## ULTRASOUND OF WHOLE ABDOMEN

**Liver** is of normal size (12.8 cm). Echo-texture is normal. No focal space occupying lesion is seen within liver parenchyma. Intrahepatic biliary channels are not dilated. Portal vein diameter is normal.

**Gall bladder** is partially distended. Common bile duct is not dilated.

**Pancreas** is of normal size and contour. Echo-pattern is normal. No focal lesion is seen within pancreas.

**Spleen** is of normal size and shape (10.0 cm). Echotexture is normal. No focal lesion is seen.

**Kidneys** are normally sited and are of normal size and shape. Cortico-medullary echoes are normal. No focal lesion is seen. Collecting system does not show any dilatation or calculus.

**Right kidney** is measuring approx. 10.6 x 4.2 cm.

**Left kidney** is measuring approx. 10.2 x 3.8 cm.

**Urinary bladder** does not show any calculus or mass lesion.

**Uterus** is anteverted and normal in size (measuring approx. 8.0 x 3.1 x 3.0 cm).

Myometrium shows normal echo -pattern. No focal space occupying lesion is seen. Endometrial echo is normal. Endometrial thickness is 3.6 mm.

Both ovaries are visualized and are normal. No adnexal mass lesion is seen.

No enlarged nodes are visualized. No retro-peritoneal lesion is identified.

No significant free fluid is seen in pouch of Douglas.

**IMPRESSION: No significant abnormality is detected**

DR. SHALINI GOEL  
M.B.B.S, D.N.B (Radiodiagnosis)  
RMC no.: 21954

Dr. SHALINI GOEL  
MBBS, DNB (Radiologist)  
RMC No. 21954  
P-3 Health Solutions LLP



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📍 B-14, Vidhyadhar Enclave-II, Near Axix Bank  
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MRS. ARCHANA	47 Y/F
Registration Date: 13/04/2024	Ref. by: BANK OF BARODA

**2D-ECHOCARDIOGRAPHY M.MODE WITH DOPPLER STUDY:**  
FAIR TRANSTHORACIC ECHOCARDIOGRAPHIC WINDOW MORPHOLOGY:

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

**M.MODE EXAMINATION:**

AO	2.6	Cm	LA	2.6	cm	IVS-D	0.9	cm
IVS-S	1.2	cm	LVID	4.0	cm	LVSD	3.6	cm
LVPW-D	0.8	cm	LVPW-S	1.2	cm	RV		cm
RVWT		cm	EDV		ml	LVVS		ml
LVEF	55-60%		RWMA			ABSENT		

**CHAMBERS:**

LA	NORMAL	RA	NORMAL
LV	NORMAL	RV	NORMAL
PERICARDIUM		NORMAL	

**COLOUR DOPPLER:**

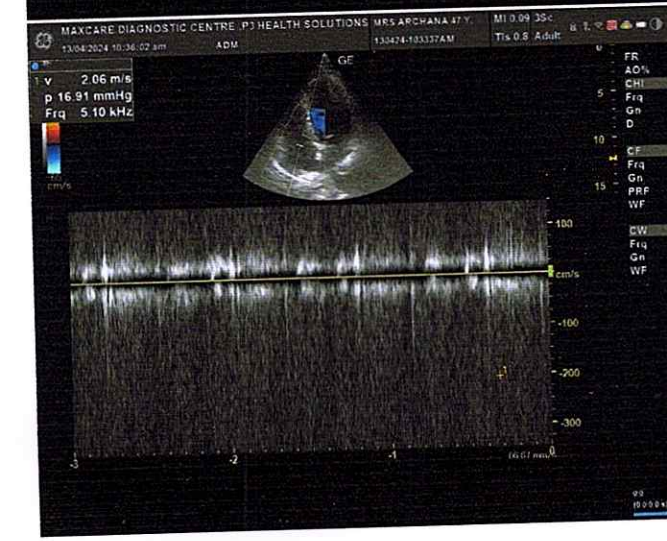
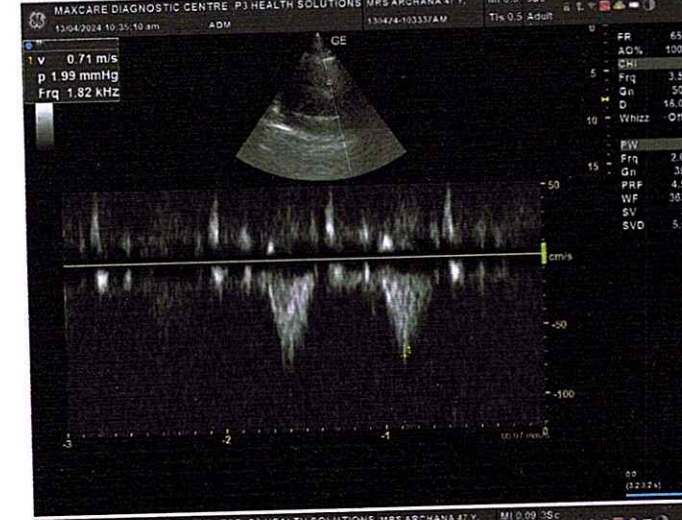
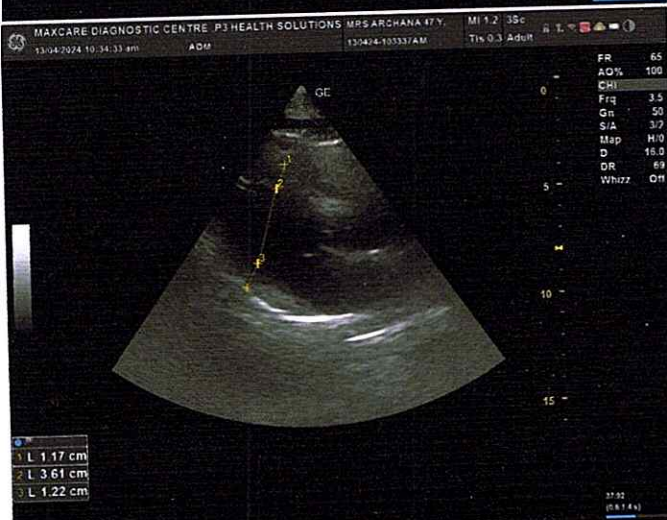
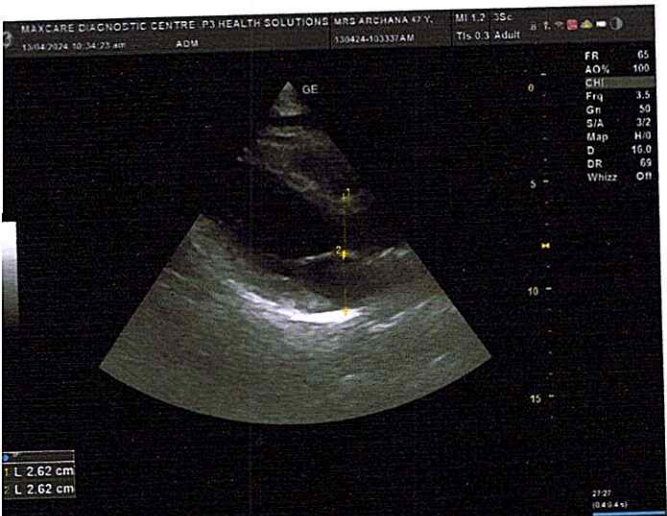
MITRAL VALVE				
E VELOCITY	0.83	m/sec	PEAK GRADIENT	Mm/hg
A VELOCITY	0.65	m/sec	MEAN GRADIENT	Mm/hg
MVA BY PHT		Cm2	MVA BY PLANIMETRY	Cm2
MITRAL REGURGITATION	ABSENT			
AORTIC VALVE				
PEAK VELOCITY	1.08	m/sec	PEAK GRADIENT	mm/hg
AR VMAX		m/sec	MEAN GRADIENT	mm/hg
AORTIC REGURGITATION	ABSENT			
TRICUSPID VALVE				
PEAK VELOCITY		m/sec	PEAK GRADIENT	mm/hg
MEAN VELOCITY		m/sec	MEAN GRADIENT	mm/hg
VMax VELOCITY				
TRICUSPID REGURGITATION	ABSENT			
PULMONARY VALVE				
PEAK VELOCITY	0.71	M/sec.	PEAK GRADIENT	Mm/hg
MEAN VELOCITY			MEAN GRADIENT	Mm/hg
PULMONARY REGURGITATION	ABSENT			

**Impression—**

- NORMAL LV SIZE & CONTRACTILITY.
- NO RWMA, LVEF 55-60%.
- ALL CARDIAC VALVES ARE NORMAL.
- NORMAL DIASTOLIC FUNCTION.
- NO CLOT, NO VEGETATION, NO PERICARDIAL EFFUSION.

**Dr. JYOTI AGARWAL**  
M.B.B.S, (Cardiologist)  
RMC No.- 27255

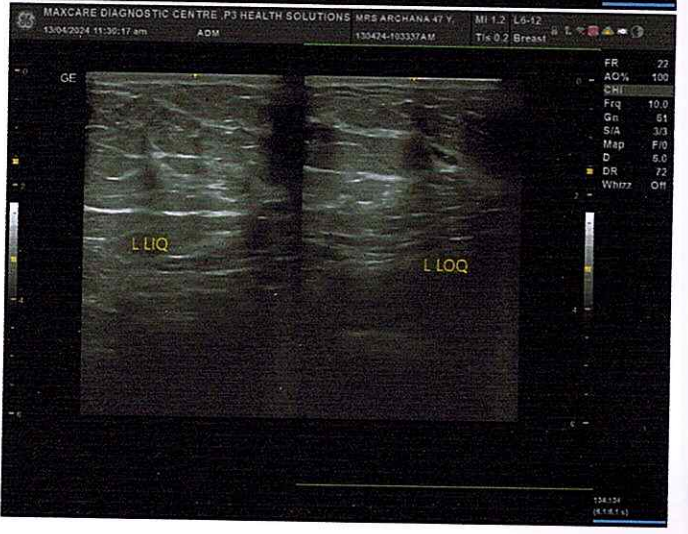
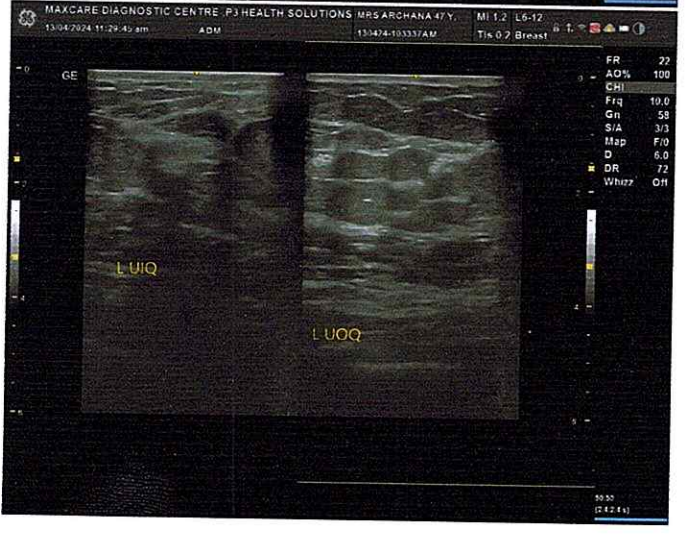
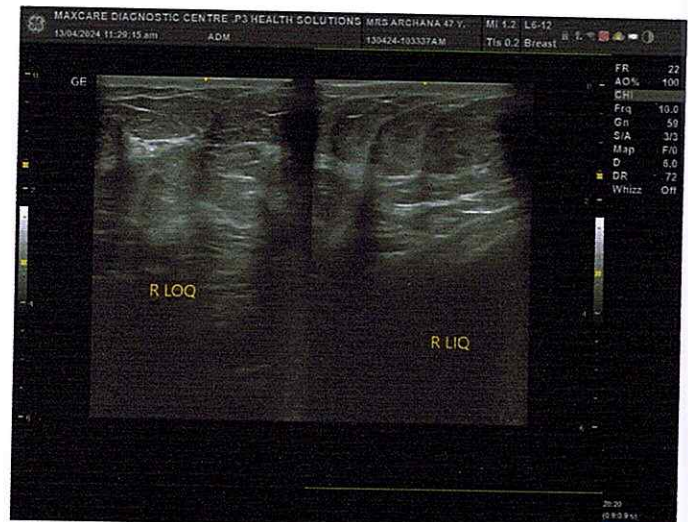
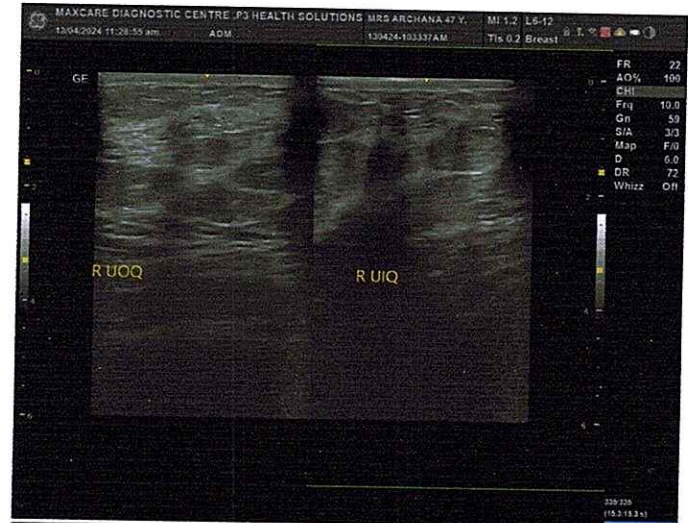














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Central Spine, Vidhyadhar Nagar, Jaipur - 302023  
☎ +91 141 4824885 📧 maxcarediagnostics1@gmail.com



NAME:	MRS. ARCHANA	AGE	47 YRS/F
REF.BY	BANK OF BARODA	DATE	13/04/2024

## CHEST X-RAY (PA VIEW)

Bilateral lung fields appear clear.

Bilateral costo-phrenic angles appear clear.

Cardiothoracic ratio is normal.

Thoracic soft tissue and skeletal system appear unremarkable.

Soft tissue shadows appear normal.

**IMPRESSION:** *No significant abnormality is detected*

*Shalini*

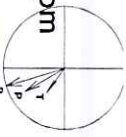
DR.SHALINI GOEL

M.B.B.S, D.N.B (Radiodiagnosis)

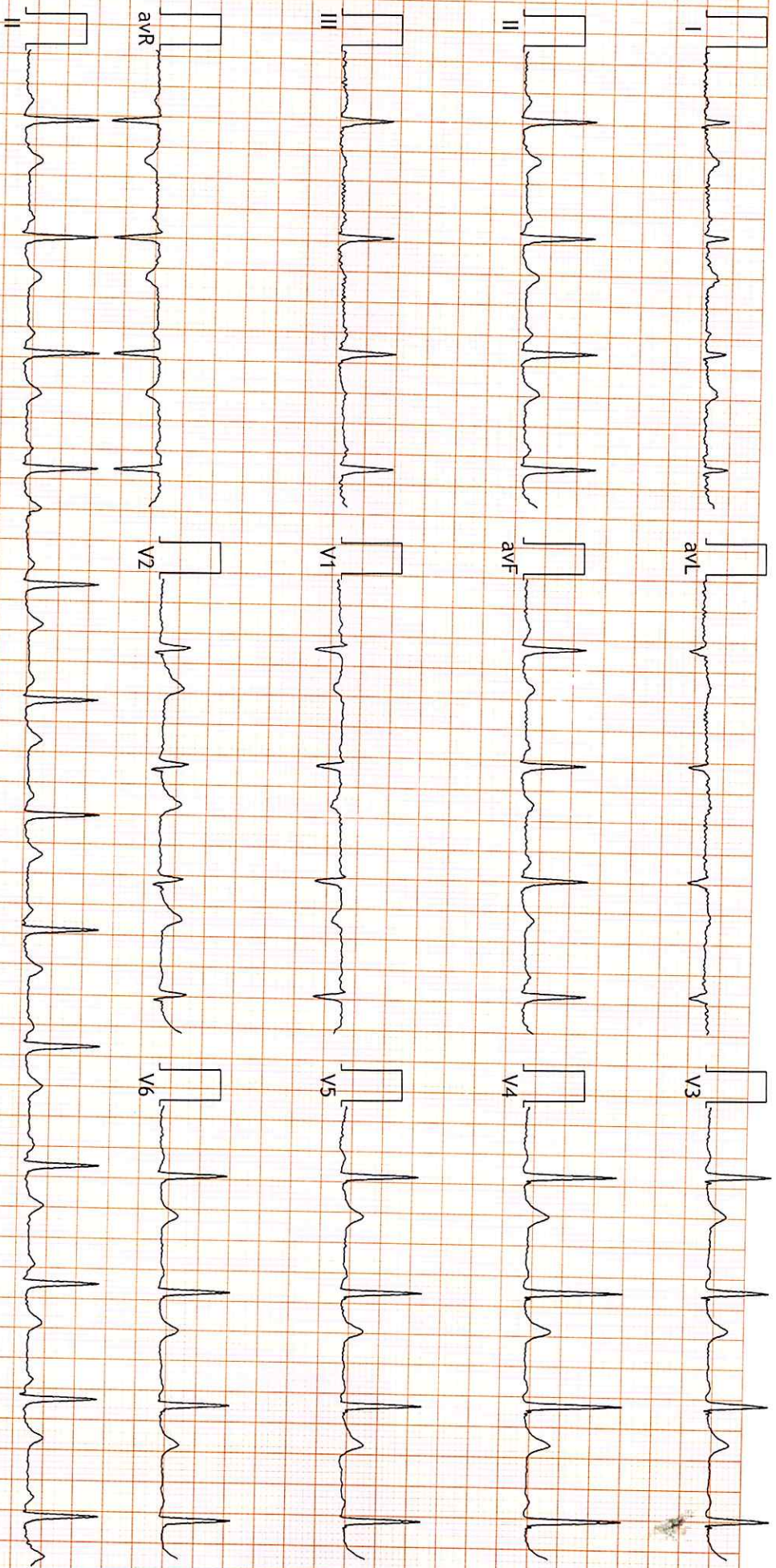
RMC No.: 21954



HR: 78 bpm



PR Interval: 142 ms  
QRS Duration: 86 ms  
QT/QTc: 371/423ms  
P-QRS-T Axis: 60 - 73 - 36 (Deg)



FINDINGS: Normal Sinus Rhythm  
Vent Rate : 78 bpm; PR Interval : 142 ms; QRS Duration: 86 ms; QT/QTc Int : 371/423 ms  
P-QRS-T axis: 60 • 73 • 36 • (Deg)  
Comments :

35/4/24

TUSA

Dr. Naresh Kumar Mohanka

MBBS, DIP. CARDIO (ESCORTS)  
RMC No., 35703

D.E.M. (BCCP)



