


UNION OF INDIA Driving Licence
(RJ) (NT)

RJ14C20170051134

	जारी करने की तिथि Date of Issue 17/03/2017	वैधता / Validity 16/03/2037	
	जन्म तिथि Date of Birth 15/04/1992	Blood Group Unknown	



नाम / Name
KAVITA JANGID

पिता/पति का नाम / Son/Daughter/Wife of
BANWARI LAL JANGID

Kavita

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

RJ14C20170051134
D02094192M

	
LMV 17/03/2017	MCWG 17/03/2017

पता / Permanent Address
 94-A NEW COLONY MURLIPURA
 JAIPUR

Holder's Signature

जारीकर्ता / Issuing Authority Sign
 DTO Vidhyadhar Nagar

Form 7, Rule 16(2)



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(ASSOCIATES OF MAXCARE DIAGNOSTICS)

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General Physical Examination

Date of Examination: 13/04/24

Name: KAVITA JANCHID Age: 31 YRS DOB: 15/04/1993 Sex: Female

Referred By: DANU DEBARODA

Photo ID: DRIVING LIC ID #: R714220170051134

Ht: 160 (cm)

Wt: 50 (Kg)

Chest (Expiration): 80 (cm)

Abdomen Circumference: 73 (cm)

Blood Pressure: 90/59 mm Hg

PR: 78 / min

RR: 18 / min

Temp: Afebrile

BMI 20.3

Eye Examination: with glass
R I E - G I G, N I G, N C B

L I E - G I G, N I G, N C B

Other: _____

No

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

Signature Of Examinee: _____

Name of Examinee: KAVITA JANCHID

Dr. PIYUSH GOYAL

Signature Medical Examiner: _____
MBBS, DMRD (Radiologist)

Name Medical Examiner: DR. PIYUSH GOYAL

RMC No.-037041



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Patient ID 122473	Patient Mob No.9714105992	Registered On	13/04/2024 08:40:22
NAME Mrs. KAVITA JANGID		Collected On	13/04/2024 10:18:26
Age 31 Yrs 13 Mon 28 Days		Authorized On	13/04/2024 17:05:09
Ref. By BANK OF BARODA		Printed On	13/04/2024 17:05:16
Lab/Hosp Mr.MEDIWHEEL			

HAEMOGARAM

HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
FULL BODY HEALTH CHECKUP BELOW 40 FEMAL			
HAEMOGLOBIN (Hb)	9.8 L	g/dL	12.0 - 15.0
TOTAL LEUCOCYTE COUNT	4.00	/cumm	4.00 - 10.00
DIFFERENTIAL LEUCOCYTE COUNT			
NEUTROPHIL	57.0	%	40.0 - 80.0
LYMPHOCYTE	37.0	%	20.0 - 40.0
EOSINOPHIL	2.0	%	1.0 - 6.0
MONOCYTE	4.0	%	2.0 - 10.0
BASOPHIL	0.0	%	0.0 - 2.0
TOTAL RED BLOOD CELL COUNT (RBC)	4.46	$\times 10^6/\mu\text{L}$	3.80 - 4.80
HEMATOCRIT (HCT)	32.70 L	%	36.00 - 46.00
MEAN CORP VOLUME (MCV)	74.0 L	fL	83.0 - 101.0
MEAN CORP HB (MCH)	22.0 L	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	29.9 L	g/dL	31.5 - 34.5
PLATELET COUNT	409	$\times 10^3/\mu\text{L}$	150 - 410
RDW-CV	16.2 H	%	11.6 - 14.0

Technologist
Page No. 1 of 16

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



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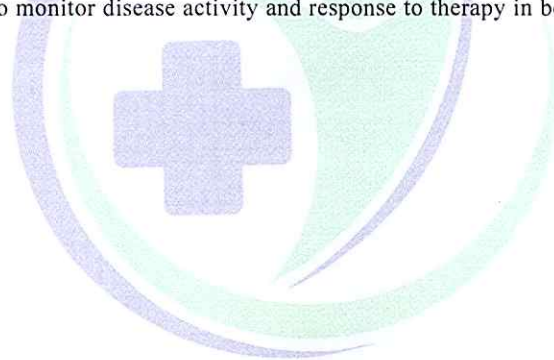
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HAEMATOTOLOGY

HAEMATOTOLOGY

Test Name	Value	Unit	Biological Ref Interval
Erythrocyte Sedimentation Rate (ESR) Method:- Westergreen	18	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

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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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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
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FASTING BLOOD SUGAR (Plasma) Method:- GLUCOSE OXIDASE/PEROXIDASE	72.1	mg/dl	70.0 - 115.0
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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 .

BLOOD SUGAR PP (Plasma) Method:- GLUCOSE OXIDASE/PEROXIDASE	78.9	mg/dl	70.0 - 140.0
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Instrument Name: HORIBA 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
GLYCOSYLATED HEMOGLOBIN (HbA1C) Method:- CAPILLARY with EDTA	5.2	mg%	Non-Diabetic < 6.0 Good Control 6.0-7.0 Weak Control 7.0-8.0 Poor control > 8.0
MEAN PLASMA GLUCOSE Method:- Calculated Parameter	104	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

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Technologist
Page No. 5 of 16



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HAEMATOLOGY

HAEMATOLOGY

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



Technologist
Page No. 5 of 16

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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIPID PROFILE			
SERUM TOTAL CHOLESTEROL Method:- CHOLESTEROL OXIDASE/PEROXIDASE	135.00	mg/dl	Desirable <200 Borderline 200-239 High > 240
InstrumentName: HORIBA Interpretation: Cholesterol measurements are used in the diagnosis and treatments of lipid lipoprotein metabolism disorders.			
SERUM TRIGLYCERIDES Method:- GLYCEROL PHOSPHATE OXIDASE/PREOXIDASE	78.60	mg/dl	Normal <150 Borderline high 150-199 High 200-499 Very high >500
InstrumentName: Radox Rx Imola Interpretation : 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	38.60	mg/dl	MALE- 30-70 FEMALE - 30-85
Instrument Name: Rx Daytona plus Interpretation: 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	83.30	mg/dl	Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190
VLDL CHOLESTEROL Method:- Calculated	15.72	mg/dl	0.00 - 80.00
T.CHOLESTEROL/HDL CHOLESTEROL RATIO Method:- Calculated	3.50		0.00 - 4.90
LDL / HDL CHOLESTEROL RATIO Method:- Calculated	2.16		0.00 - 3.50
TOTAL LIPID Method:- CALCULATED	402.49	mg/dl	400.00 - 1000.00

Technologist
Page No. 7 of 16

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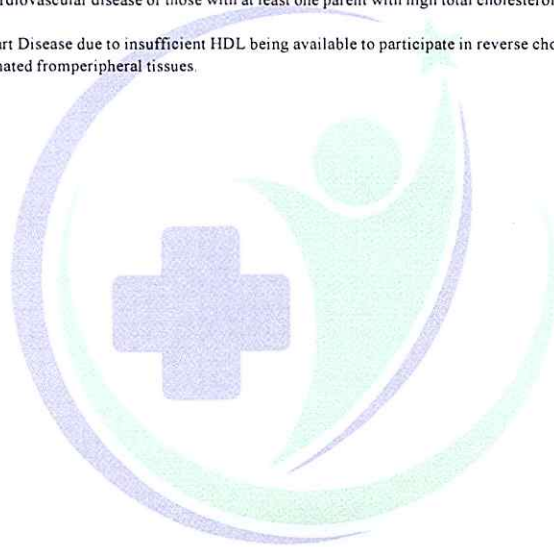
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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
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1. Measurements in the same patient can show physiological& analytical variations. Three serialsamples 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

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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIVER PROFILE WITH GGT			
SERUM BILIRUBIN (TOTAL) Method:- DIAZOTIZED SULFANILIC	0.52	mg/dL	Infants : 0.2-8.0 mg/dL Adult - Up to - 1.2 mg/dL
SERUM BILIRUBIN (DIRECT) Method:- DIAZOTIZED SULFANILIC	0.12	mg/dL	Up to 0.40 mg/dL
SERUM BILIRUBIN (INDIRECT) Method:- Calculated	0.40	mg/dl	0.30-0.70
SGOT Method:- IFCC	14.7	U/L	0.0 - 40.0
SGPT Method:- IFCC	16.5	U/L	0.0 - 35.0
SERUM ALKALINE PHOSPHATASE Method:- DGKC - SCE	74.20	U/L	64.00 - 306.00
InstrumentName: MISPA PLUS Interpretation: Measurements of alkaline phosphatase are of use in the diagnosis, treatment and investigation of hepatobiliary disease and in bone disease associated with increased osteoblastic activity. Alkaline phosphatase is also used in the diagnosis of parathyroid and intestinal disease.			
SERUM GAMMA GT Method:- Szasz methodology Instrument Name Randox Rx Imola	22.30	U/L	5.00 - 32.00
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.			
SERUM TOTAL PROTEIN Method:- BIURET	6.58	g/dl	6.00 - 8.40
SERUM ALBUMIN Method:- BROMOCRESOL GREEN	4.21	g/dl	3.50 - 5.50
SERUM GLOBULIN Method:- CALCULATION	2.37	gm/dl	2.20 - 3.50
A/G RATIO	1.78		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.

Technologist
Page No. 9 of 16

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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
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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., 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

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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
RFT / KFT WITH ELECTROLYTES			
SERUM UREA Method:- UREASE / GLUTAMATE DEHYDROGENASE	23.20	mg/dl	10.00 - 50.00
InstrumentName: HORIBA CA 60 Interpretation : Urea measurements are used in the diagnosis and treatment of certain renal and metabolic diseases.			
SERUM CREATININE Method:- JAFFE	0.90	mg/dl	Males : 0.6-1.50 mg/dl Females : 0.6 -1.40 mg/dl
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	5.21	mg/dl	2.40 - 7.00
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	141.7	mmol/L	135.0 - 150.0
POTASSIUM Method:- ISE	4.39	mmol/L	3.50 - 5.50
CHLORIDE Method:- ISE	1.0 L	mmol/L	94.0 - 110.0
SERUM CALCIUM Method:- Arsenazo III Method	9.25	mg/dL	8.80 - 10.20
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
SERUM ALBUMIN Method:- BRÖMOCRESOL GREEN	4.21	g/dl	3.50 - 5.50

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Technologist
 Page No: 1 of 16



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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
SERUM GLOBULIN Method:- CALCULATION	2.37	gm/dl	2.20 - 3.50
A/G RATIO	1.78		1.30 - 2.50

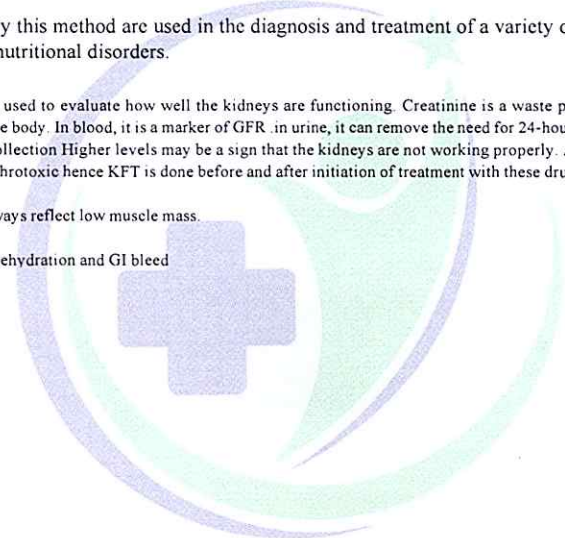
Interpretation : Measurements obtained by this method are used in the diagnosis and treatment of a variety of diseases involving the liver, kidney and bone marrow as well as other metabolic or nutritional disorders.

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
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Patient ID 122473	Patient Mob No.9714105992	Registered On	13/04/2024 08:40:22
NAME Mrs. KAVITA JANGID		Collected On	13/04/2024 10:18:26
Age 31 Yrs 13 Mon 28 Days		Authorized On	13/04/2024 17:05:09
Ref. By BANK OF BARODA		Printed On	13/04/2024 17:05:16
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

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IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
TOTAL THYROID PROFILE			
THYROID-TRIIODOTHYRONINE T3 Method:- ECLIA	0.97	ng/mL	0.70 - 2.04
THYROID - THYROXINE (T4) Method:- ECLIA	9.04	ug/dl	5.10 - 14.10
TSH Method:- ECLIA	1.771	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 Teitz 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 ***

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Page No. 16 of 16



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Lab/Hosp	Mr.MEDIWHEEL		

CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
Urine Routine			
<u>PHYSICAL EXAMINATION</u>			
COLOUR	PALE YELLOW		PALE YELLOW
APPEARANCE	Clear		Clear
<u>CHEMICAL EXAMINATION</u>			
REACTION(PH)	5.0		5.0 - 7.5
SPECIFIC GRAVITY	1.030		1.010 - 1.030
PROTEIN	NIL		NIL
SUGAR	NIL		NIL
BILIRUBIN	NEGATIVE		NEGATIVE
UROBILINOGEN	NORMAL		NORMAL
KETONES	NEGATIVE		NEGATIVE
NITRITE	NEGATIVE		NEGATIVE
<u>MICROSCOPY EXAMINATION</u>			
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

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Page No. 13 of 16

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NAME:	MRS. KAVITA JANGID	AGE	31 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.

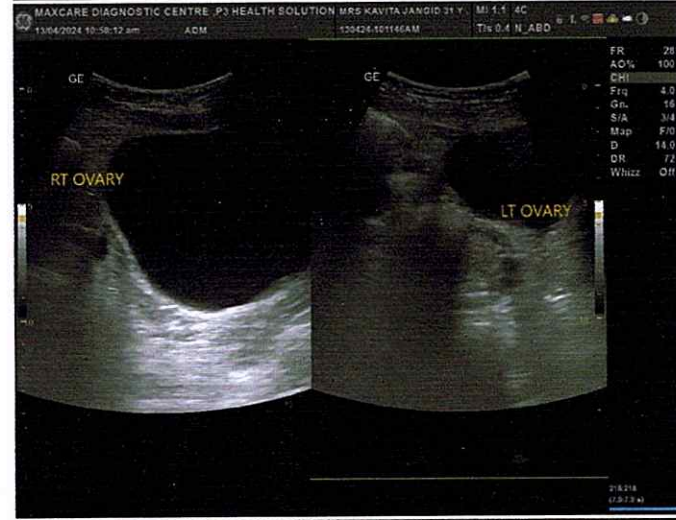
Soft tissue shadows appear normal.

IMPRESSION: *No significant abnormality is detected*

DR.SHALINI GOEL

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

RMC No.: 21954





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MRS. KAVITA JANGID	31 Y/F
Registration Date: 13/04/2024	Ref. by: BANK OF BARODA

ULTRASOUND OF WHOLE ABDOMEN

Liver is of normal size (13.0 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 (9.4 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. 9.0 x 3.4 cm.

Left kidney is measuring approx. 10.5 x 3.6 cm.

Urinary bladder does not show any calculus or mass lesion.

Uterus is anteverted and normal in size (measuring approx. 8.0 x 3.7 x 3.8 cm).

Myometrium shows normal echo -pattern. No focal space occupying lesion is seen. Endometrial echo is normal. Endometrial thickness is 6.0 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.

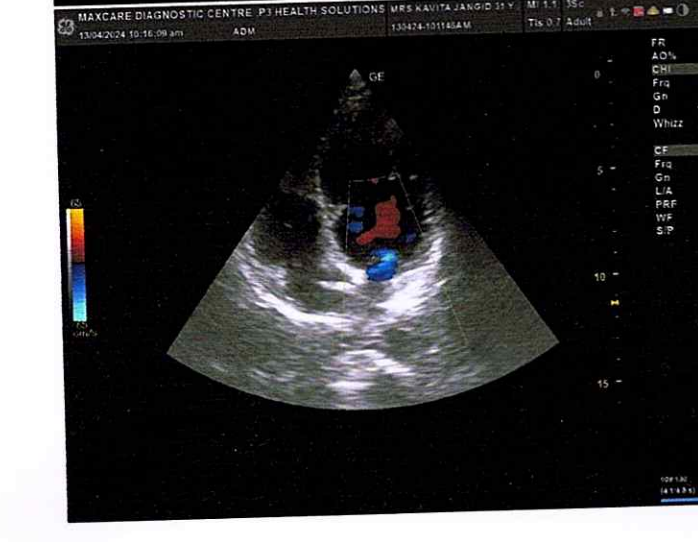
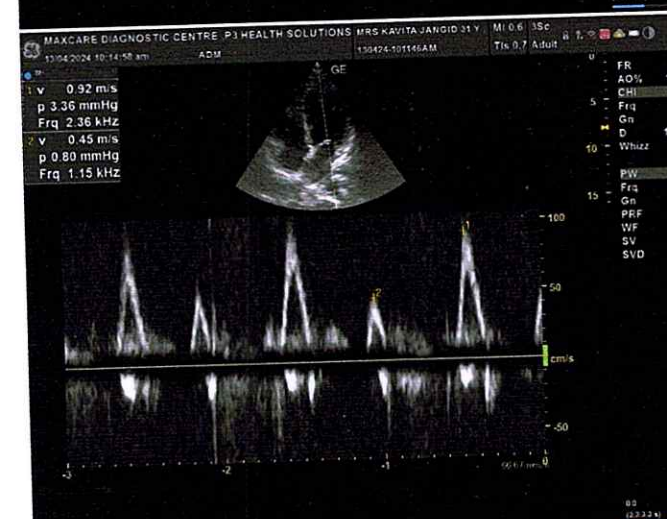
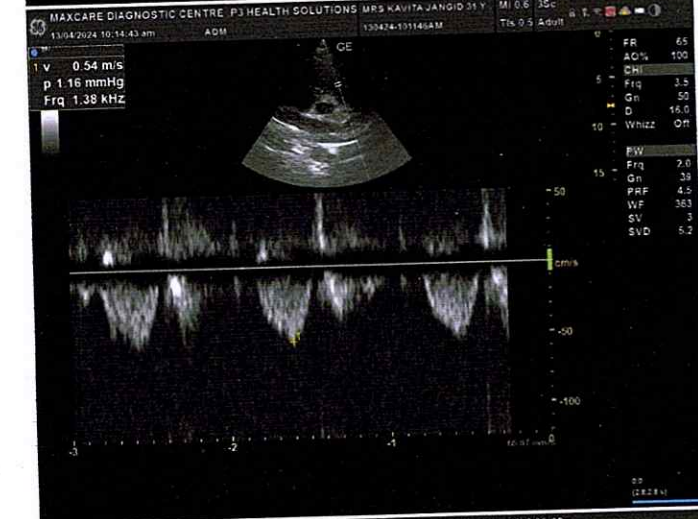
IMPRESSION: *No significant abnormality is detected*

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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MRS. KAVITA JANGID

31 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	2.3	Cm	LA	2.4	cm	IVS-D	0.9	cm
IVS-S	1.2	cm	LVID	4.5	cm	LVSD	2.7	cm
LVPW-D	0.9	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	0.92	m/sec	PEAK GRADIENT	Mm/hg
A VELOCITY	0.45	m/sec	MEAN GRADIENT	Mm/hg
MVA BY PHT		Cm2	MVA BY PLANIMETRY	Cm2
MITRAL REGURGITATION	TRACE			
AORTIC VALVE				
PEAK VELOCITY	1.31	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	MILD			
PULMONARY VALVE				
PEAK VELOCITY	0.54	M/sec.	PEAK GRADIENT	Mm/hg
MEAN VELOCITY			MEAN GRADIENT	Mm/hg
PULMONARY REGURGITATION	ABSENT			

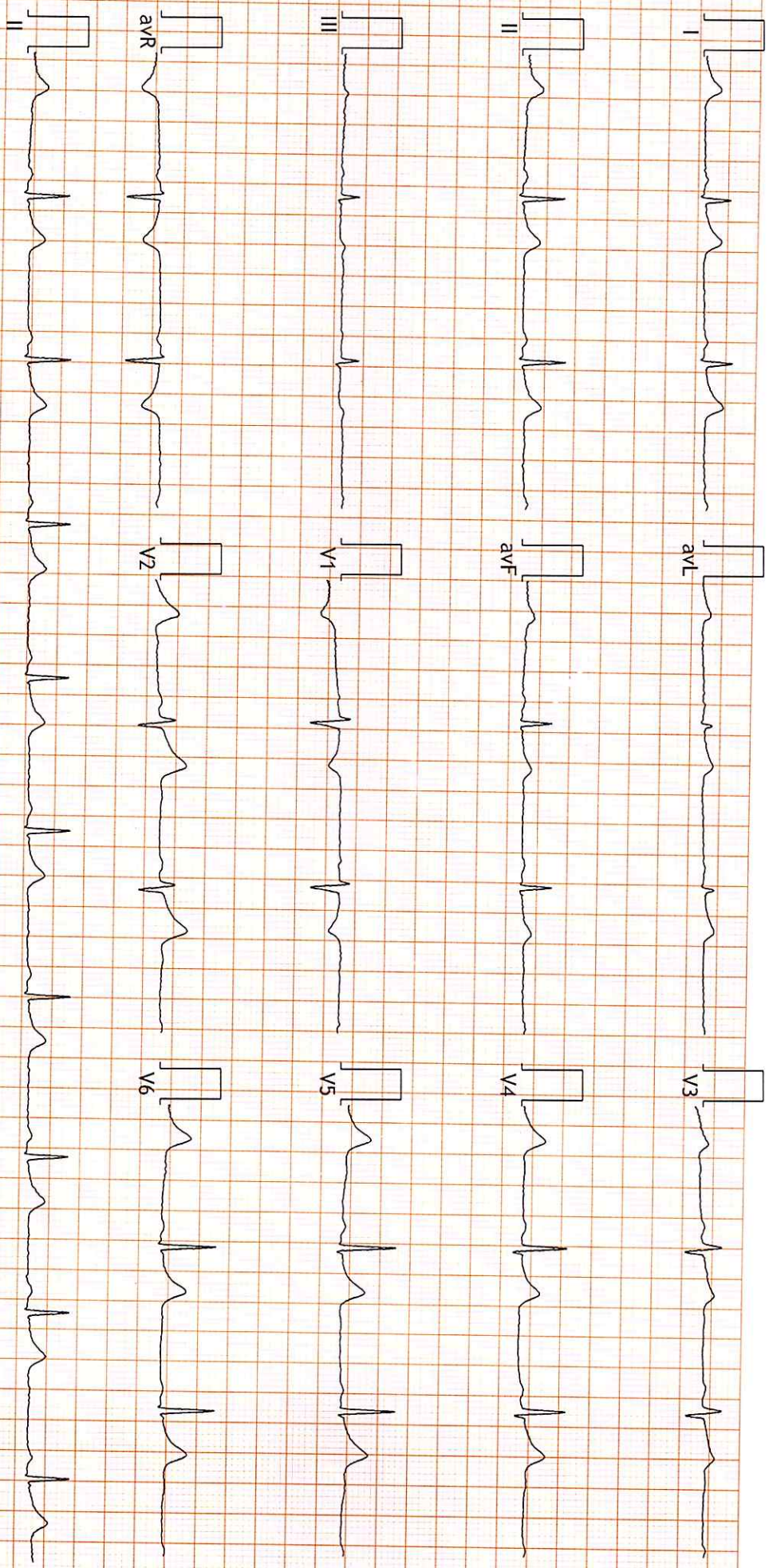
Impression—

- NORMAL LV SIZE & CONTRACTILITY.
- NO RWMA, LVEF 55-60%.
- MILD TR/ PAH (RVSP 31 MMHG+ RAP), TRACE MR.
- NORMAL DIASTOLIC FUNCTION.
- NO CLOT, NO VEGETATION, NO PERICARDIAL EFFUSION.

(Cardiologist)
DR. JYOTI AGARWAL

M.B.B.S, PGDCC (Cardiologist)

RMC No.- 27255



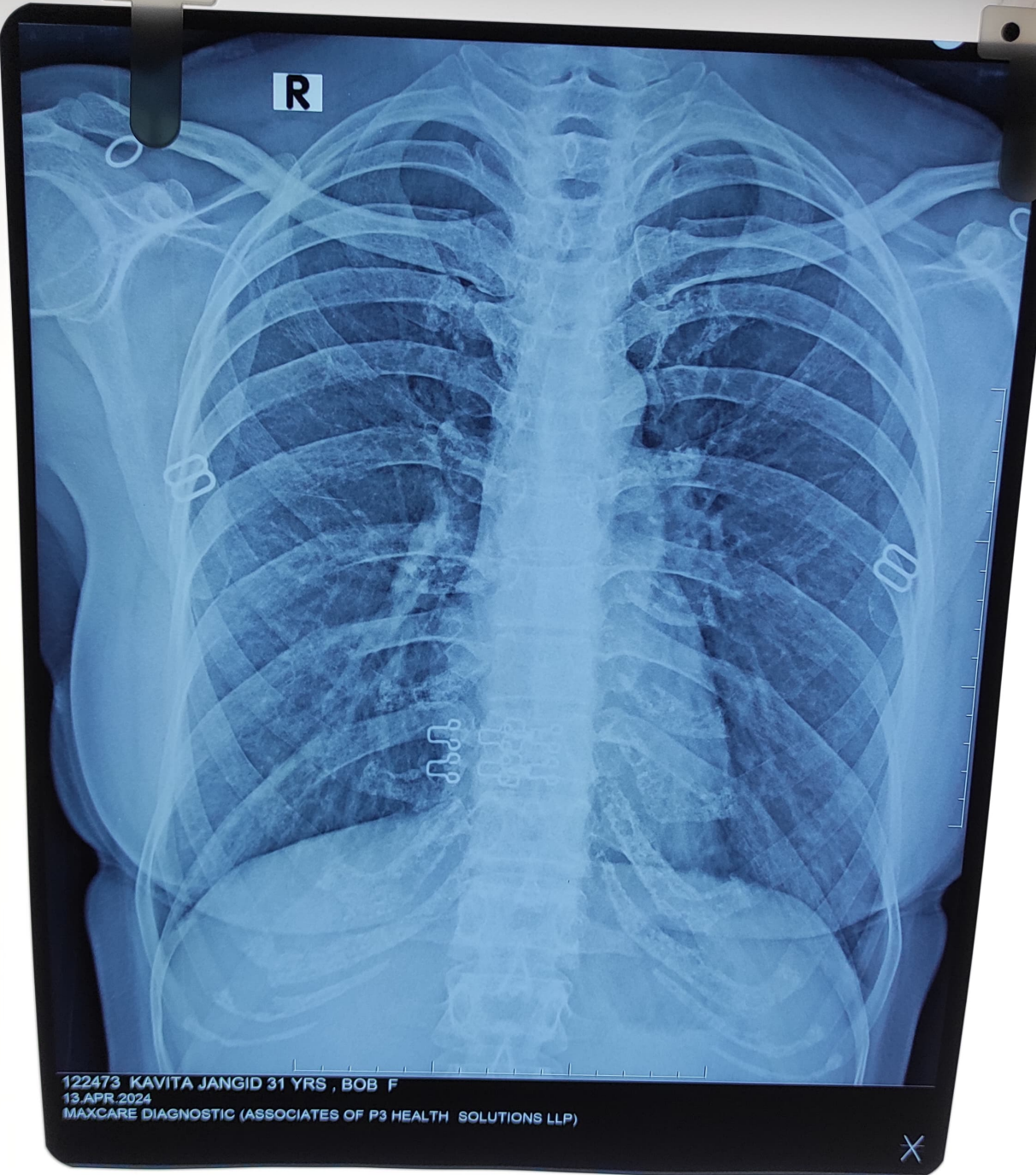
FINDINGS: Abnormal ECG with Indication of Sinus Bradycardia
Vent Rate : 56 bpm; PR Interval : 176 ms; QRS Duration: 94 ms; QT/QTc Int : 399/387 ms
P-QRS-T axis: 40 • 40 • 24 • (Deg)
Comments :

SINUS BRADYCARDIA

[Signature]

[Signature]

Dr. Naresh Kumar Mohanka
RMC No.: 35703
MBBS, DIP. CARDIO (ESCORTS)
D.E.M. (RCGP-UK)



122473 KAVITA JANGID 31 YRS , BOB F
13 APR 2024
MAXCARE DIAGNOSTIC (ASSOCIATES OF P3 HEALTH SOLUTIONS LLP)

X