

भारत सरकार
Government of India

आधार



Jitendra Thakur
Jitendra Thakur
जन्म तिथि/DOB: 27/09/1969
पुरुष/ MALE

~~9183 3045 5623~~ 9236
VID: 9183 3045 5623 9450

मेरा आधार, मेरी पहचान

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

भारतीय विशिष्ट पहचान प्राधिकरण
Unique Identification Authority of India

आधार

पता:
S/O: Bhogendra Thakur, A 40, satya nagar,
Jhotwara, Jaipur, Jaipur,
Rajasthan - 302012

Address:
S/O: Bhogendra Thakur, A 40, satya nagar,
Jhotwara, Jaipur, Jaipur,
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(ASSOCIATES OF MAXCARE DIAGNOSTICS)

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Central Spine, Vidhyadhar Nagar, Jaipur - 302023
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General Physical Examination

Date of Examination: 13/04/2024

Name: JITENDRA THAKUR Age: 54 yrs DOB: 07/09/1969 Sex: Male

Referred By: DANIK OF BARODA

Photo ID: AADHAR CARD ID #: 9036

Ht: 170 (cm)

Wt: 83 (Kg)

Chest (Expiration): 104 (cm)

Abdomen Circumference: 109 (cm)

Blood Pressure: 103/81 mm Hg PR: 79/min RR: 18/min Temp: Afebrile

BMI 28.7

Eye Examination: RIE - GIG, NIG, NCB
LEI - GIG, NIG, NCB

Other: No

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

Signature Of Examinee: [Signature]

Name of Examinee: JITENDRA THAKUR

Signature Medical Examiner: Dr. PIYUSH GOYAL
MBBS, DMRD (Radiologist)
RMC No.-037041

Name Medical Examiner: DR. PIYUSH GOYAL



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Patient ID 122475	Patient Mob No.9462962318	Registered On	13/04/2024 08:46:17
NAME Mr. JITENDRA THAKUR		Collected On	13/04/2024 10:18:34
Age 54 Yrs	Sex M	Authorized On	13/04/2024 17:13:30
Ref. By BANK OF BARODA		Printed On	13/04/2024 17:13:38
Lab/Hosp Mr.MEDIWHEEL			

HAEMOGARAM

HAEMATOTOLOGY

Test Name	Value	Unit	Biological Ref Interval
FULL BODY HEALTH CHECKUP ABOVE 40 MALE			
HAEMOGLOBIN (Hb)	15.3	g/dL	13.0 - 17.0
TOTAL LEUCOCYTE COUNT	7.10	/cumm	4.00 - 10.00
DIFFERENTIAL LEUCOCYTE COUNT			
NEUTROPHIL	54.0	%	40.0 - 80.0
LYMPHOCYTE	39.0	%	20.0 - 40.0
EOSINOPHIL	3.0	%	1.0 - 6.0
MONOCYTE	4.0	%	2.0 - 10.0
BASOPHIL	0.0	%	0.0 - 2.0
TOTAL RED BLOOD CELL COUNT (RBC)	5.44	$\times 10^6/\mu\text{L}$	4.50 - 5.50
HEMATOCRIT (HCT)	48.10	%	40.00 - 50.00
MEAN CORP VOLUME (MCV)	88.0	fL	83.0 - 101.0
MEAN CORP HB (MCH)	28.1	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	31.8	g/dL	31.5 - 34.5
PLATELET COUNT	324	$\times 10^3/\mu\text{L}$	150 - 410
RDW-CV	14.0	%	11.6 - 14.0

Technologist

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DR.TANU RUNGTA
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RMC No. 17226



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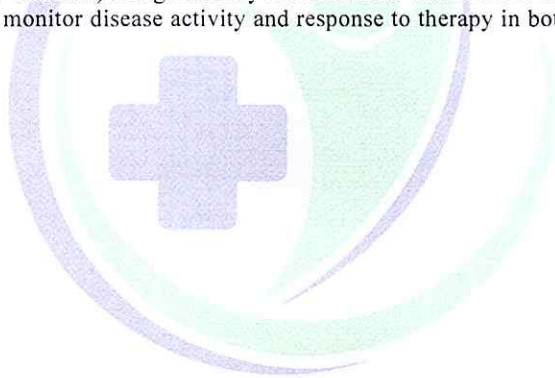
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HAEMATOLOGY

HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
Erythrocyte Sedimentation Rate (ESR) <small>Method:- Westergreen</small>	10	mm in 1st hr	00 - 15

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



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Lab/Hosp Mr.MEDIWHEEL

(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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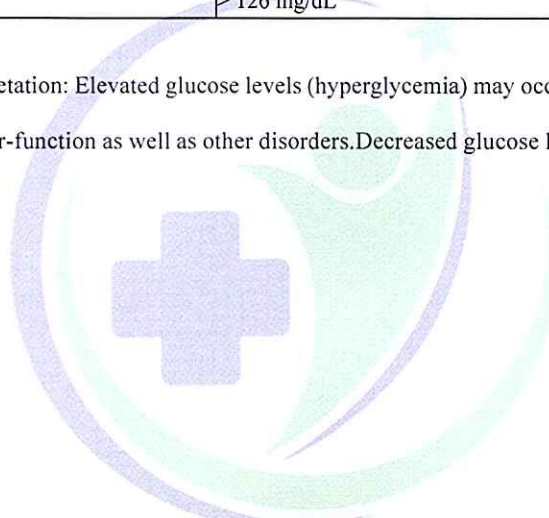


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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
FASTING BLOOD SUGAR (Plasma) Method:- GLUCOSE OXIDASE/PEROXIDASE	76.8	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 .



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HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
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GLYCOSYLATED HEMOGLOBIN (HbA1C)

Method:- CAPILLARY with EDTA

5.6 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

110 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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HAEMATOLOGY

HAEMATOLOGY

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

"O" POSITIVE



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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
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LIPID PROFILE

SERUM TOTAL CHOLESTEROL 167.00 mg/dl
 Method:- CHOLESTEROL OXIDASE/PEROXIDASE
 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 139.00 mg/dl
 Method:- GLYCEROL PHOSPHATE OXIDASE/PREOXIDASE
 Normal <150
 Borderline high 150-199
 High 200-499
 Very high >500

InstrumentName:Randox 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 42.30 mg/dl
 Method:- Direct clearance Method
 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 101.53 mg/dl
 Method:- Calculated Method
 Optimal <100
 Near Optimal/above optimal 100-129
 Borderline High 130-159
 High 160-189
 Very High > 190

VLDL CHOLESTEROL 27.80 mg/dl
 Method:- Calculated
 0.00 - 80.00

T.CHOLESTEROL/HDL CHOLESTEROL RATIO 3.95
 Method:- Calculated
 0.00 - 4.90

LDL / HDL CHOLESTEROL RATIO 2.40
 Method:- Calculated
 0.00 - 3.50

TOTAL LIPID 535.53 mg/dl
 Method:- CALCULATED
 400.00 - 1000.00

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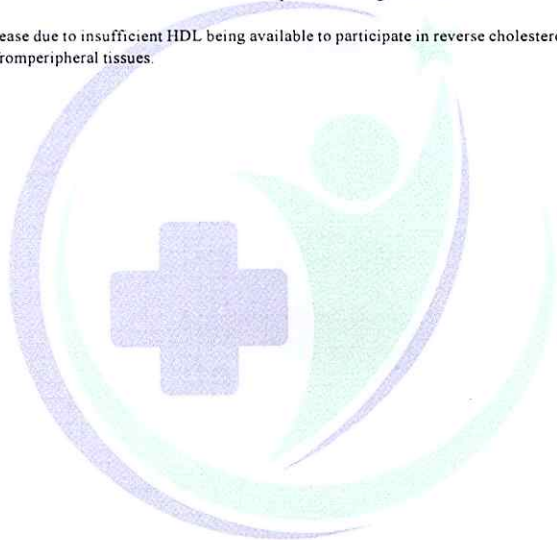
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BIOCHEMISTRY

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 fromperipheral tissues.



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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIVER PROFILE WITH GGT			
SERUM BILIRUBIN (TOTAL) Method:- DIAZOTIZED SULFANILIC	0.78	mg/dL	Infants : 0.2-8.0 mg/dL Adult - Up to - 1.2 mg/dL
SERUM BILIRUBIN (DIRECT) Method:- DIAZOTIZED SULFANILIC	0.23	mg/dL	Up to 0.40 mg/dL
SERUM BILIRUBIN (INDIRECT) Method:- Calculated	0.55	mg/dl	0.30-0.70
SGOT Method:- IFCC	26.3	U/L	0.0 - 40.0
SGPT Method:- IFCC	29.6	U/L	0.0 - 40.0
SERUM ALKALINE PHOSPHATASE Method:- DGKC - SCE	87.60	U/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.	32.30	U/L	10.00 - 45.00
SERUM TOTAL PROTEIN Method:- BIURET	6.88	g/dl	6.00 - 8.40
SERUM ALBUMIN Method:- BROMOCRESOL GREEN	4.25	g/dl	3.50 - 5.50
SERUM GLOBULIN Method:- CALCULATION	2.63	gm/dl	2.20 - 3.50
A/G RATIO	1.62		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.,

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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.



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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	33.20	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 diseases.

SERUM CREATININE Method:- JAFFE	1.17	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
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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	144.4	mmol/L	135.0 - 150.0
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POTASSIUM Method:- ISE	4.93	mmol/L	3.50 - 5.50
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CHLORIDE Method:- ISE	104.9	mmol/L	94.0 - 110.0
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SERUM CALCIUM Method:- Arsenazo III Method	8.99	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	6.88	g/dl	6.00 - 8.40
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SERUM ALBUMIN Method:- BROMOCRESOL GREEN	4.25	g/dl	3.50 - 5.50
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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
SERUM GLOBULIN Method:- CALCULATION	2.63	gm/dl	2.20 - 3.50
A/G RATIO	1.62		1.30 - 2.50

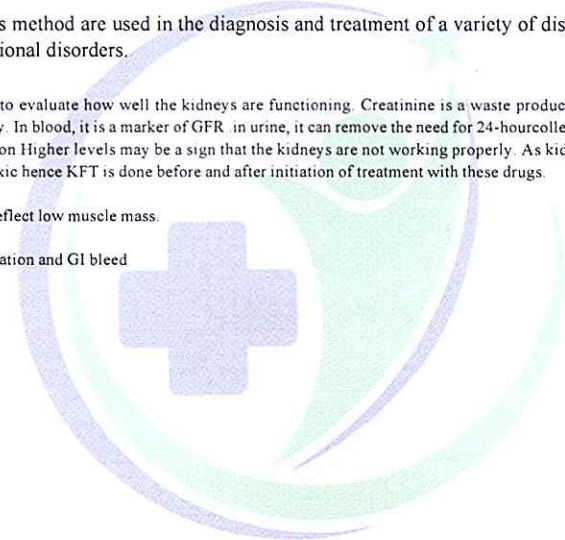
Interpretation : Measurements obtained by this method are used in the diagnosis and treatment of a variety of diseases involving the liver, kidney and bone marrow as well as other metabolic or nutritional disorders.

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
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Tanu
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 MD (Pathology)
 RMC No. 17226



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Patient ID 122475	Patient Mob No.9462962318	Registered On	13/04/2024 08:46:17
NAME Mr. JITENDRA THAKUR		Collected On	13/04/2024 10:18:34
Age	54 Yrs Sex M on 18/04/2024	Authorized On	13/04/2024 17:13:30
Ref. By	BANK OF BARODA	Printed On	13/04/2024 17:13:38
Lab/Hosp	Mr.MEDIWHEEL		

CLINICAL PATHOLOGY

CLINICAL PATHOLOGY

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



Technologist
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IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
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PSA (PROSTATE SPECIFIC ANTIGEN) -TOTAL 0.318 ng/mL 0.00-4.00
Method:- Methodology: CLIA

CLINICAL NOTES:- Prostate-specific antigen (PSA) is a 34-kD glycoprotein produced almost exclusively by the prostate gland.

PSA is normally present in the blood at very low levels. Increased levels of PSA may suggest the presence of prostate cancer.

1. Immediate PSA testing following digital rectal examination, ejaculation, prostatic massage, indwelling catheterization, ultrasonography and needle biopsy of prostate is not recommended as they falsely elevate levels

2. PSA values regardless of levels should not be interpreted as absolute evidence of the presence or absence of disease. All values should be correlated with clinical findings and other investigations

3. Physiological decrease in PSA level by 18% has been observed in sedentary patients either due to supine position or suspended sexual activity

Clinical Use

- An aid in the early detection of Prostate cancer when used in conjunction with Digital rectal examination in males more than 50 years of age and in those with two or more affected first degree relatives.
- Follow up and management of Prostate cancer patients
- Detect metastatic or persistent disease in patients following surgical or medical treatment of Prostate cancer

NOTE

PSA levels can be also increased by prostatitis, irritation, benign prostatic hyperplasia (BPH), and recent ejaculation, producing a false positive result. Digital rectal examination (DRE) has been shown in several studies to produce an increase in PSA. However, the effect is clinically insignificant, since DRE causes the most substantial increases in patients with PSA levels already elevated over 4.0 ng/mL.

Obesity has been reported to reduce serum PSA levels. Delayed early detection may partially explain worse outcomes in obese men with early prostate cancer. After treatment, higher BMI also correlates to higher risk of recurrence.

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IMMUNOASSAY

IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
-----------	-------	------	-------------------------

TOTAL THYROID PROFILE

THYROID-TRIIODOTHYRONINE T3

Method:- ECLIA

1.13 ng/mL

0.70 - 2.04

THYROID - THYROXINE (T4)

Method:- ECLIA

8.18 ug/dl

5.10 - 14.10

TSH

Method:- ECLIA

3.564 μ IU/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 \uparrow serum T3 & T4 values along with \downarrow TSH level.
- 2.Primary hypothyroidism is accompanied by \downarrow serum T3 and T4 values & \uparrow serum TSH levels
- 3.Normal T4 levels accompanied by \uparrow T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis
- 4.Normal or \downarrow T3 & \uparrow T4 levels indicate T4 Thyrotoxicosis (problem is conversion of T4 to T3)
- 5.Normal T3 & T4 along with \downarrow 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 ***

Tanu

Technologist
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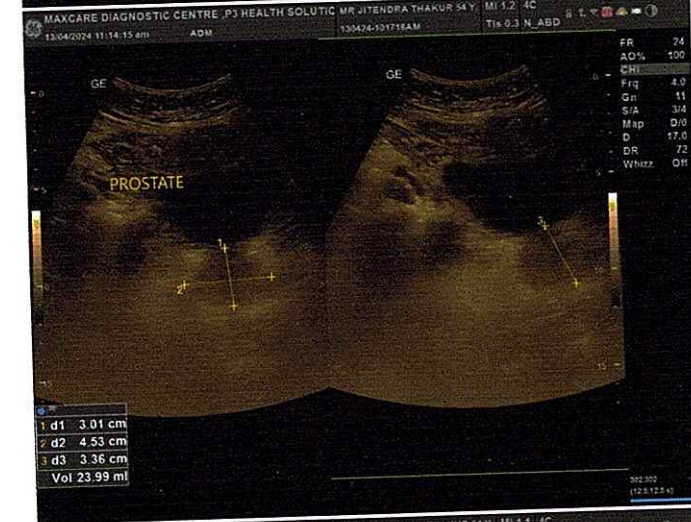
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Lab/Hosp Mr.MEDIWHEEL			

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	1.015		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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MR. JITENDRA THAKUR	54 Y/M
Registration Date: 13/04/2024	Ref. by: BANK OF BARODA

ULTRASOUND OF WHOLE ABDOMEN

Liver is of normal size (13.9 cm) **with increased echotexture**. No focal space occupying lesion is seen within liver parenchyma. Intrahepatic biliary channels are not dilated. Portal vein diameter is normal.

Gall bladder is well distended. Wall is not thickened. No calculus or mass lesion is seen in gall bladder. 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. Collecting system does not show any calculus or dilatation.

Right kidney is measuring approx. 10.4 x 4.3 cm.

Left kidney is measuring approx. 9.4 x 5.1 cm.

Urinary bladder is well distended and does not show any calculus or mass lesion.

Prostate is normal in size (measuring approx. 3.0 x 4.5 x 3.3 cm, volume 23-24 cc) with normal echotexture and outline.

No enlarged nodes are visualized. No retro-peritoneal lesion is identified.
No significant free fluid is seen in pelvis.

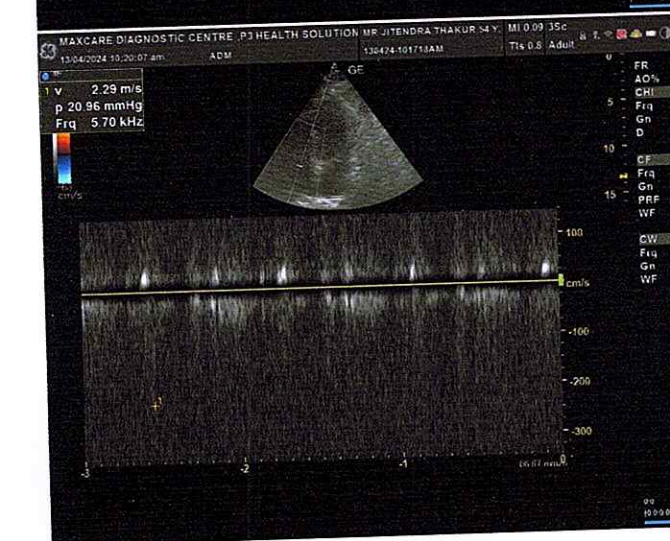
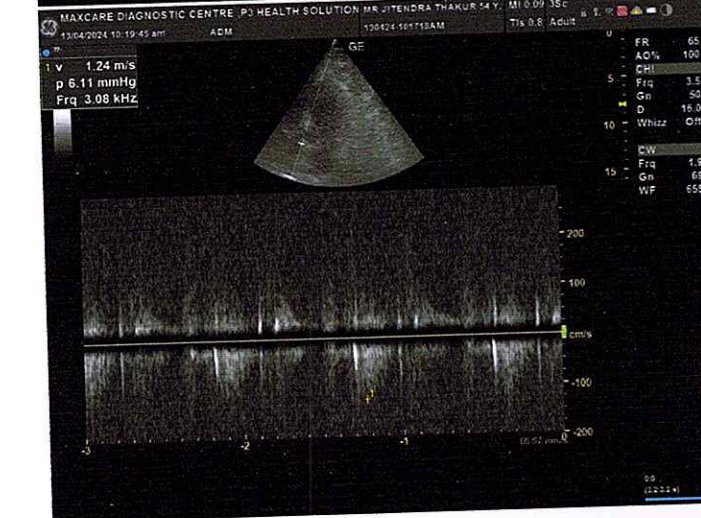
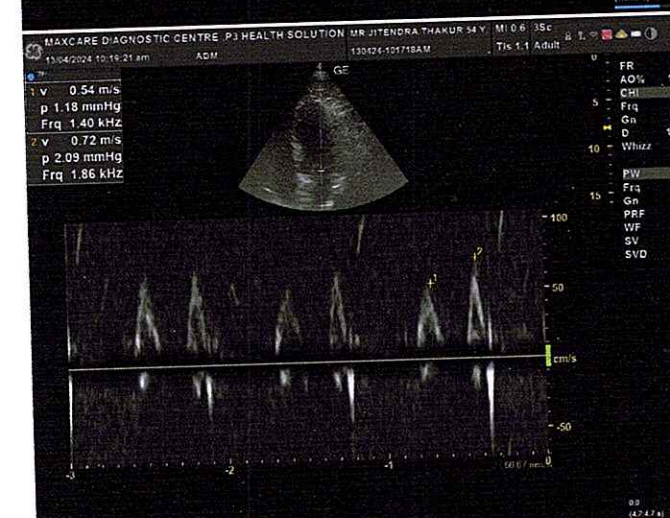
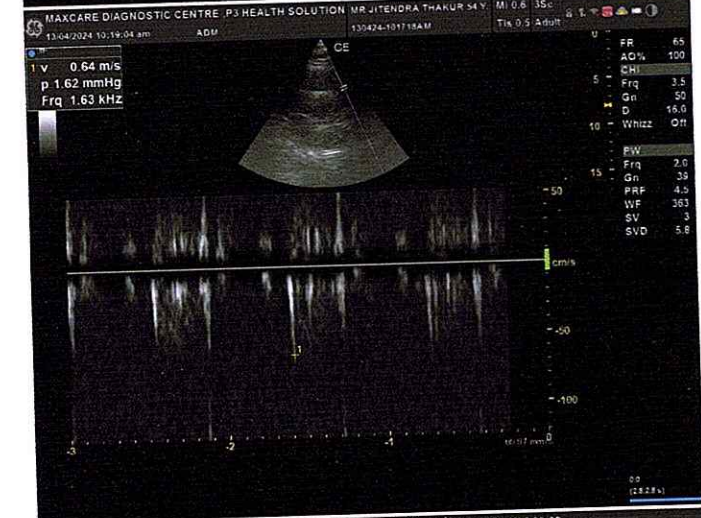
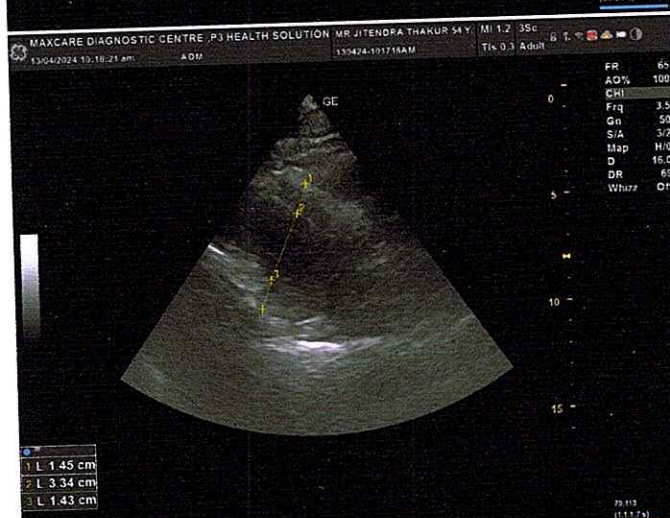
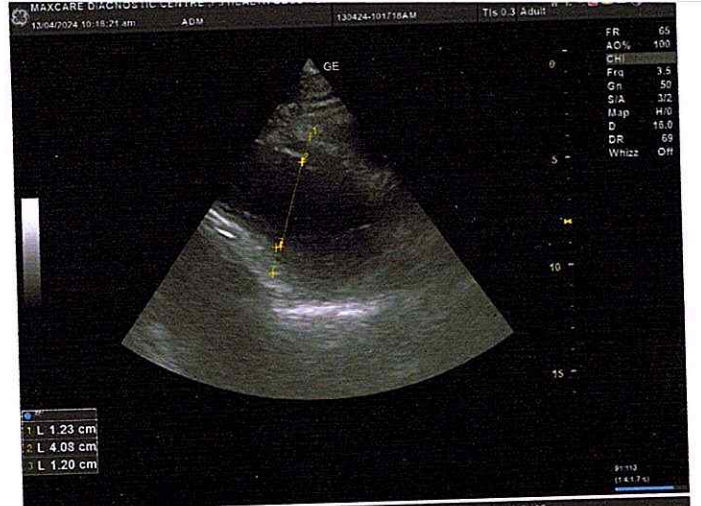
IMPRESSION:

- Grade I fatty liver.
- Rest no significant abnormality is detected.

Shalini

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

Dr. SHALINI GOEL
MBBS, DNB (Radiologist)
RMC No. 21954
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MR. JITENDRA THAKUR	54 Y/M
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	3.6	Cm	LA	3.2	cm	IVS-D	1.2	cm
IVS-S	1.4	cm	LVID	4.1	cm	LVSD	3.3	cm
LVPW-D	1.2	cm	LVPW-S	1.4	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	0.54	m/sec	PEAK GRADIENT	Mm/hg
A VELOCITY	0.72	m/sec	MEAN GRADIENT	Mm/hg
MVA BY PHT		Cm2	MVA BY PLANIMETRY	Cm2
MITRAL REGURGITATION		ABSENT		
AORTIC VALVE				
PEAK VELOCITY	1.24	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	0.64	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.
- CONCENTRIC LVH
- GRADE I DIASTOLIC DYSFUNCTION.
- NO CLOT, NO VEGETATION, NO PERICARDIAL EFFUSION.

Dr. IYOTI AGARWAL
(Cardiologist)
M.B.B.S, PGDCC (Cardiologist)
RMC No.- 27255



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NAME:	MR. JITENDRA THAKUR	AGE	54 YRS/M
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.

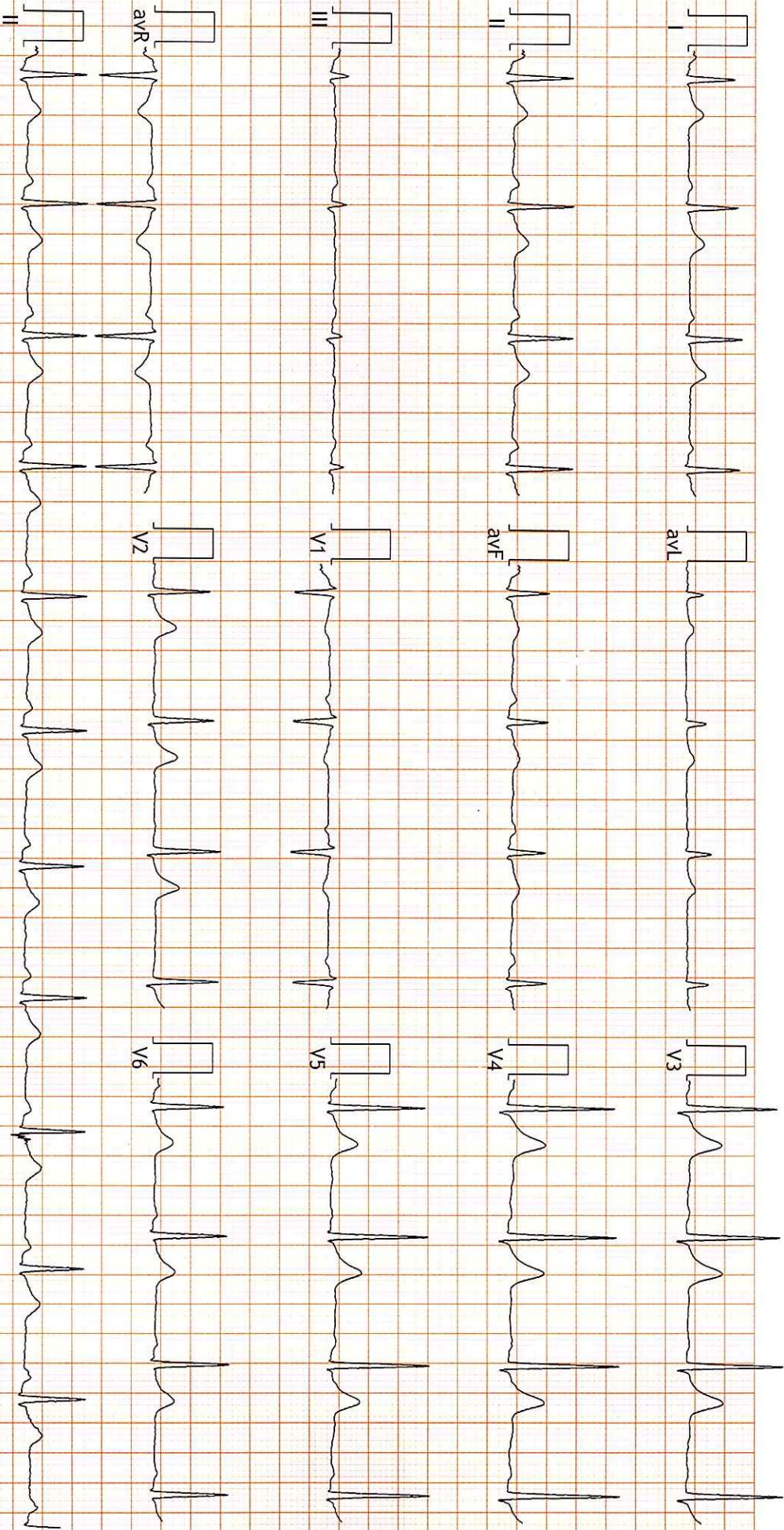
Soft tissue shadows appear normal.

IMPRESSION: *No significant abnormality is detected*

DR.SHALINI GOEL

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

RMC No.: 21954



FINDINGS: Normal Sinus Rhythm

Vent Rate : 67 bpm; PR Interval : 176 ms; QRS Duration: 100 ms; QT/QTc Int : 364/385 ms

P-QRS-T axis: 57•33•36• (Deg)

Comments :

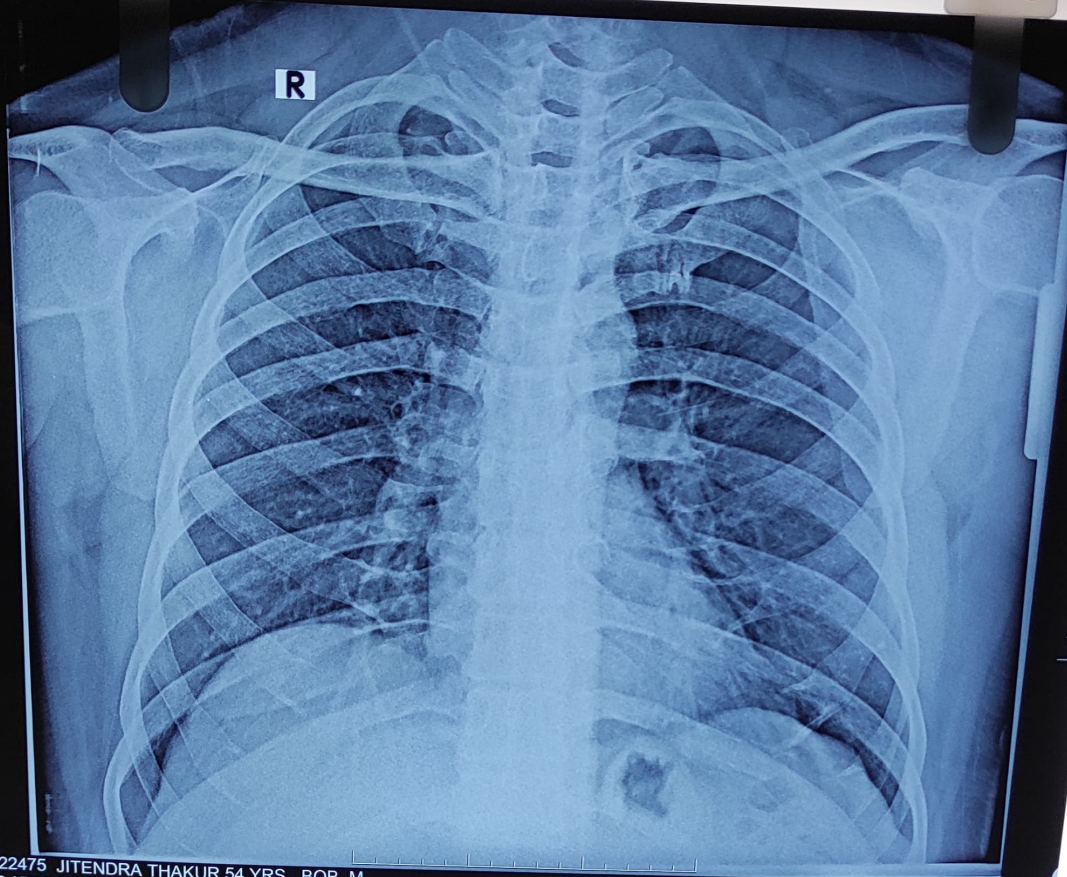
TUN

Dr. Naresh Kumar Mohanka

RMC No.: 35703

MBBS, DIP. CARDIO (ESCORTS)

D.E.M. (RCGP-UK)



122475 JITENDRA THAKUR 54 YRS , BOB M
13 APR 2024
MAXCARE DIAGNOSTIC (ASSOCIATES OF P3 HEALTH SOLUTIONS LLP)