



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



(ASSOCIATES OF MAXCARE DIAGNOSTICS)

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



General Physical Examination

Date of Examination: 26/03/25	
Name: DINESH KUMAR SHARMA	Age: 54 DOB: 01/09/1069 Sex: MAC
Referred By: RANK OF BARODA	
Photo ID: AADHINA ID#: 4690	
Ht: <u>163</u> (cm)	Wt: <u>¬</u> (Kg)
Chest (Expiration):\overline{\lambda} (cm)	Abdomen Circumference: 108 (cm)
Blood Pressure: 120/80 mm Hg PR: 79	/min RR: 18 /min Temp: Afebyle
BMI 25 Eye Examination: RE-1616 Al6	JCB
Eye Examination:	
Other:	
On examination he/she appears physically and me	entally fit: Yes / No
Signature Of Examine :	Name of Examinee: -DIVESH KUMAR SHARMA-
Signature Medical Examiner: - DMRD (Radiolo MBBS, DMRD) - 037041	Name Medical Examiner - DR. PINUSH LAND



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NAME :- Mr. DINESH KUMAR SHARMA

Age:- 54 Yrs 6 Mon 26 Days

Sex :- Male

Patient ID: -12234988

Date :- 26/03/2024

10:38:56

Ref. By Doctor:-BANK OF BARODA

Lab/Hosp :-

Company :-

Mr.MEDIWHEEL

Final Authentication: 26/03/2024 17:58:09

HAEMOGARAM

HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
FULL BODY HEALTH CHECKUP BELOW 40	FEMAL		
HAEMOGLOBIN (Hb)	14.1	g/dL	13.0 - 17.0
TOTAL LEUCOCYTE COUNT	6.60	/cumm	4.00 - 10.00
DIFFERENTIAL LEUCOCYTE COUNT			
NEUTROPHIL	55.0	%	40.0 - 80.0
LYMPHOCYTE	40.0	%	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)	4.90	x10^6/uL	4.50 - 5.50
HEMATOCRIT (HCT)	44.60	%	40.00 - 50.00
MEAN CORP VOLUME (MCV)	91.0	fL	83.0 - 101.0
MEAN CORP HB (MCH)	28.8	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	31.6	g/dL	31.5 - 34.5
PLATELET COUNT	164	x10^3/uL	150 - 410
RDW-CV	14.3 H	%	11.6 - 14.0

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HAEMATOLOGY

Erythrocyte Sedimentation Rate (ESR)

00

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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(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

WAS CALDINATED AND					
Test Name	Value	Unit	Biological Ref Interval		
FASTING BLOOD SUGAR (Plasma) Methord:- GOD POD	84.2	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

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)

Methord: - GOD PAP

86.9

mg/dl

70.0 - 140.0

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.

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HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
GLYCOSYLATED HEMOGLOBIN (HbA1C) Methord:- CAPILLARY with EDTA	5.7	mg%	Non-Diabetic < 6.0 Good Control 6.0-7.0 Weak Control 7.0-8.0 Poor control > 8.0
MEAN PLASMA GLUCOSE Methord:- Calculated Parameter	112	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 erythropolesis.
- Decreased HbA1c: administration of erythropoletin, 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.

- 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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DR.TANU RUNGTA MD (Pathology)

RMC No. 17226



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HAEMATOLOGY

BLOOD GROUP ABO Methord:- Haemagglutination reaction "O" POSITIVE



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BIOCHEMISTRY			
Test Name	Value	Unit	Biological Ref Interval
LIPID PROFILE			
TOTAL CHOLESTEROL Methord:- CHOD-PAP methodology	159.00	mg/dl	Desirable <200 Borderline 200-239 High> 240
InstrumentName:MISPA PLUS Interpreta disorders.	tion: Cholesterol measurements	are used in the diagnosis a	and treatments of lipid lipoprotein metabolism
TRIGLYCERIDES Methord:- GPO-PAP	159.00 H	mg/dl	Normal <150 Borderline high 150-199
	100000000000000000000000000000000000000		High 200-499 Very high >500
InstrumentName:Randox Rx Imola Interp	retation : Triglyceride measurer	ments are used in the diagr	nosis and treatment of diseases involving lipid

metabolism and various endocrine disorders e.g. diabetes mellitus, nephrosis and liver obstruction.

DIRECT HDL CHOLESTEROL

Methord:- Direct clearance Method

43.20

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 improved accuracy and reproducibility when compared to precipitation methods

LDL CHOLESTÉROL Methord:- Calculated Method

89.30

mg/dl

Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190

VLDL CHOLESTEROL

31.80

mg/dl

0.00 - 80.00

Methord:- Calculated

3.68

T.CHOLESTEROL/HDL CHOLESTEROL RATIO Methord: - Calculated

0.00 - 4.90

LDL / HDL CHOLESTEROL RATIO Methord:- Calculated

2.07

0.00 - 3.50

TOTAL LIPID

537.37

mg/dl

400.00 - 1000.00

Methord:- CALCULATED 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

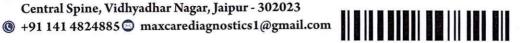
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DR.TANU RUNGTA MD (Pathology)

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BIOCHEMISTRY

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

LIVER PROFILE WITH GGT			
SERUM BILIRUBIN (TOTAL) Methord:- DMSO/Diazo	0.72	mg/dL	Infants : 0.2-8.0 mg/dL Adult - Up to - 1.2 mg/dL
SERUM BILIRUBIN (DIRECT) Methord:- DMSO/Diazo	0.22	mg/dL	Up to 0.40 mg/dL
SERUM BILIRUBIN (INDIRECT) Methord:- Calculated	0.50	mg/dl	0.30-0.70
SGOT Methord:- IFCC	22.7	U/L	0.0 - 40.0
SGPT Methord:- IFCC	26.5	U/L	0.0 - 40.0
SERUM ALKALINE PHOSPHATASE Methord:- DGKC - SCE	87.20	U/L	53.00 - 141.00
SERUM GAMMA GT Methord:- Szasz methodology Instrument Name Randox Rx Imola Interpretation: Elevations in GGT levels are seen earlier and more pronounced than thos	20.20 e with other liver enzymes	U/L s in cases of obstructive jaundice and	10.00 - 45.00
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 r	ormal)are observed with i	nfectious hepatitis.	
SERUM TOTAL PROTEIN Methord:- Direct Biuret Reagent	6.74	g/dl	6.00 - 8.40
SERUM ALBUMIN Methord:- Bromocresol Green	4.20	g/dl	3.50 - 5.50
SERUM GLOBULIN Methord:- CALCULATION	2.54	gm/dl	2.20 - 3.50
A/G RATIO	1.65		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., 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

RFT / KFT WITH ELECTROLYTES

SERUM UREA Methord:- Urease/GLDH 30.20

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

Methord:- Jaffe's Method

1.04

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

Methord:- Arsenazo III Method

5.21

mg/dl

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 135.0 - 150.0 139.0 mmol/L Methord:- ISE 3.50 - 5.50 **POTASSIUM** 4.36 mmol/L Methord:- ISE 94.0 - 110.0 **CHLORIDE** 103.0 mmol/L Methord:- ISE mg/dL 8.80 - 10.20 SERUM CALCIUM 9.65

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 Methord:- Direct Biuret Reagent	6.74	g/dl	6.00 - 8.40
SERUM ALBUMIN Methord:- Bromocresol Green	4.20	g/dl	3.50 - 5.50
SERUM GLOBULIN Methord:- CALCULATION	2.54	gm/dl	2.20 - 3.50
A/G RATIO	1.65		1.30 - 2.50

Interpretation: Measurements obtained by this method are used in the diagnosis and treatment of a variety of dis

'iver, kidney and

DR.TANU RUNGTA

MD (Pathology) RMC No. 17226

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BIOCHEMISTRY

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-hourcollections 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 bloodincreases. 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



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CLINICAL PATHOLOGY

URINE SUGAR (FASTING)
Collected Sample Received

Nil

Nil



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TOTAL THYROID PROFILE

IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
THYROID-TRIIODOTHYRONINE T3 Methord:- ECLIA	1.13	ng/mL	0.70 - 2.04
THYROID - THYROXINE (T4) Methord:- ECLIA	7.64	ug/dl	5.10 - 14.10
TSH Methord:- ECLIA	1.232	μIU/mL	0.350 - 5.500

4th Generation Assay, Reference ranges vary between laboratories

PREGNANCY - REFERENCE RANGE for TSH IN ulU/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 circardian 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 1 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 \(\text{TSH indicate mild / Subclinical Hyperthyroidism} \)

. 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 clinical data for interpretation.

*** End of Report ***

Lechnologist MGR Page No: 15 of 15 DR.TANU RUNGTA

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CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
Urine Routine		8	
PHYSICAL EXAMINATION			
COLOUR	PALE YELLO	OW	PALE YELLOW
APPEARANCE	Clear		Clear
CHEMICAL EXAMINATION			
REACTION(PH)	5.5		5.0 - 7.5
SPECIFIC GRAVITY	1.025		1.010 - 1.030
PROTEIN	NIL		NIL
SUGAR	NIL		NIL
BILIRUBIN	NEGATIVE		NEGATIVE
UROBILINOGEN	NORMAL		NORMAL
KETONES	NEGATIVE		NEGATIVE
NITRITE	NEGATIVE		NEGATIVE
MICROSCOPY EXAMINATION			
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		

Technologist MGR Page No: 12 of 15



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



MR. DINESH KUMAR SHARMA	54 Y/M
Registration Date: 26/03/2024	Ref. by: BANK OF BARODA

ULTRASOUND OF WHOLE ABDOMEN

Liver is of normal size (14.4 cm) with increased echotexture. No focal space occupying lesion is seen within liver parenchyma. Intra hepatic 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 (10.1 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. 11.4 x 3.9 cm.

• A simple, well-defined cortical cyst measuring 15 x 16 mm is noted in lower pole. Left kidney is measuring approx. 11.2 x 4.5 cm.

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

Prostate is enlarged in size (measuring 4.5 x 5.5 x 4.4 cm, volume 59-60 cc) with indentation of base of urinary bladder by median lobe and intravesical component of 9-10 mm. No focal lesion is seen.

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

IMPRESSION:

- Grade 4 prostatomegaly.
- Grade I fatty liver.

DR.SHALINI GOEL

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

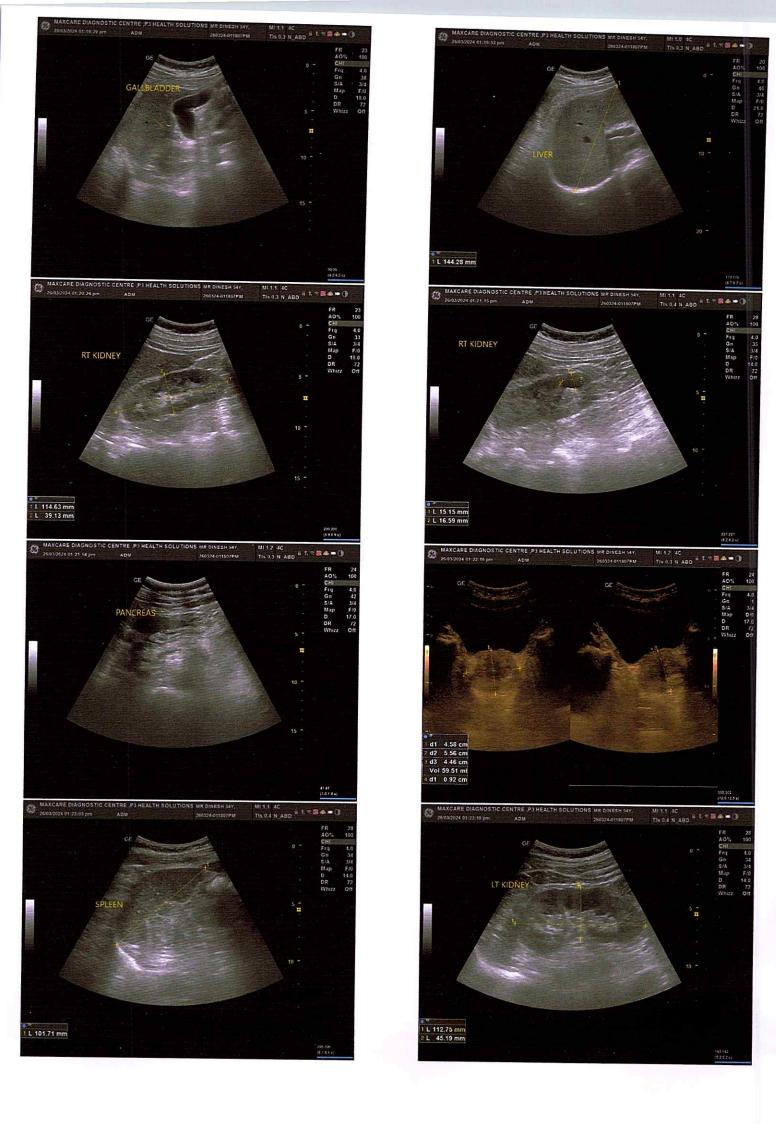
RMC no.: 21954

Dr. SHALINI GOEL

MBBS, DNB (Radiologist)

RMC No. 21954

P-3 Health Solutions LLP



\ef.: BANK OF BARODA Test Date: 26-Mar-2024(3:05:13 P) Notch: 50Hz 0.05Hz - 35Hz וems (א) בזם P3 HEALTH SOLUTIONS LLP B-14, Vidhyadhar nahar , Jaipur 128541925461288/Mr Dinesh Kumar Sharma 54Yrs-11Months/Male P-QRS-T axis: 65 · 67 · 56 · (Deg) Comments: Vent Rate: 74 bpm; PR Interval: 170 ms; QRS Duration: 88 ms; QT/QTc Int: 362/402 ms FINDINGS: Normal Sinus Rhythm avR 5 avL < Kgs/ 10mm/mV Cms 25mm/Sec BP: mmHg HR: PR Interval: 170 ms
QRS Duration: 88 ms
74 bpm QT/QTc: 362/402ms 8 5 **V**4 5 P-QRS-T Axis: 65 - 67 - 56 (Deg) MML MBBS, DIP, CARDIO (ESCORTS)
D.E.M. (RCGP-UK) Dr. Naresh Kumar Mohanka

B-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur 54 Yrs/Male 0 Kg/0 Cms

Date: 26-Mar-2024 03:07:21 PM

10238595/MR DINESH KUMAR SHARMA

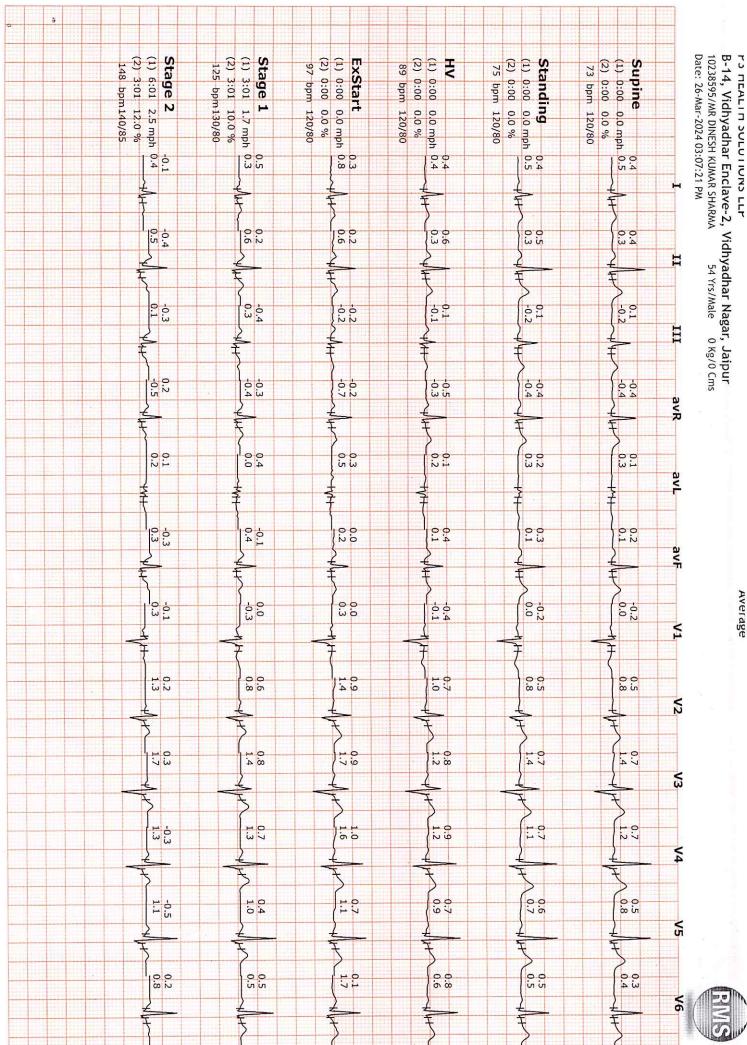
Protocol : BRUCE History: Nil

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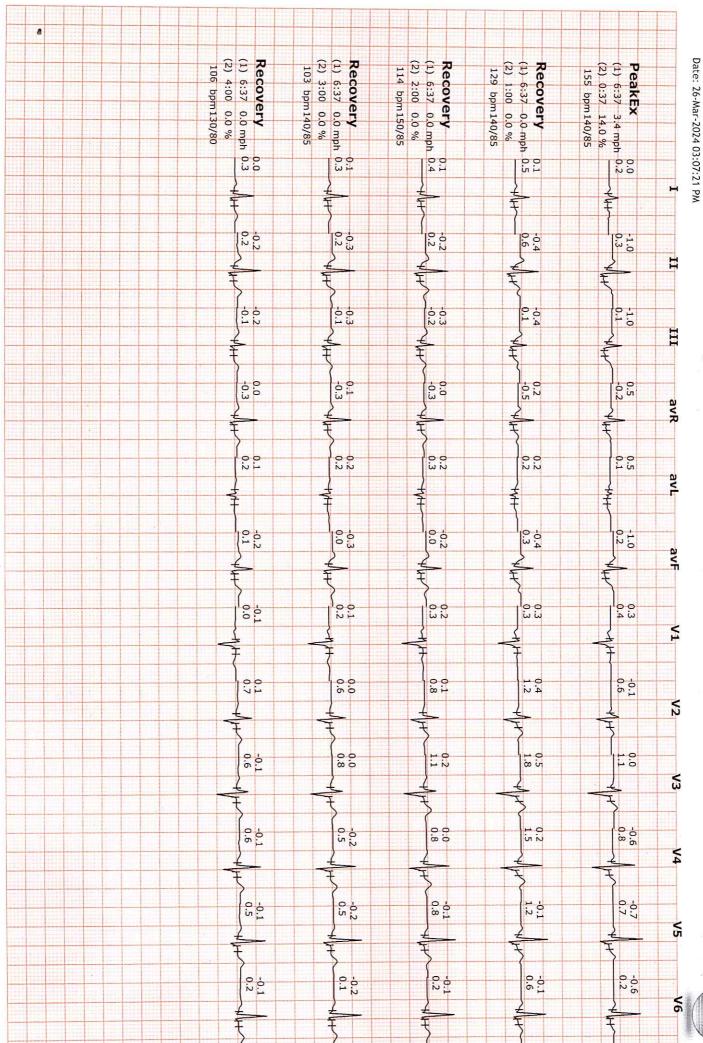
Advice/Comments: Stage Findings: PeakEx Stage 1 Supine Recovery Recovery Recovery Recovery Stage 2 ExStart YH Objective: Standing Medication: Nil Ref.By : BANK OF BARODA Max BP : 150/85(mmHg) Max HR Attained **Exercise Time** Max WorkLoad attained :7.7(Fair Effort Tolerance) StageTime PhaseTime Speed 3:00 2:00 1:00 0:37 4:00 3:01 3:01 6:38 6:02 3:02 :155 bpm 93% of Max Predictable HR 0.0 0.0 0.0 3.4 Grade 14.0 ~ 4 1.0 7.7 4.7 1.0 1.0 1.0 1.2 METS .0 .0 106 103 129 155 125 H.R. 148 (bpm) 97 73 89 75 166 140/85 130/80 140/85 120/80 120/80 130/80 120/80 150/85 140/85 140/85 120/80 Negative В. Р. R.P.P. ×100 87 217 137 14 207 116 7 180 62 106 90 PVC Comments PreEx V3 -0.2 PeakEx 3 0.9 avR ave V2 av MBBS, DIP. CARDIO (ESCORTS)
D.E.M. (RCGP-UK) **V3** < 46 **Y**5 V4 STL Naresh Kumar Mohanka 6 0.5 mm/Div 3 12 5 18 21 Min.



Average



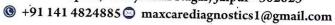
rs педет п эоготтом егг B-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur 10238595/mr DINESH KUMAR SHARMA 54 Yrs/Male 0 Kg/0 Cms Date: 26-Mar-2024 03:07:21 PM





(ASSOCIATES OF MAXCARE DIAGNOSTICS)

B-14, Vidhyadhar Enclave-II, Near Axix Bank Central Spine, Vidhyadhar Nagar, Jaipur - 302023





NAME:	MR. DINESH KUMAR SHARMA	AGE	54 YRS/M
REF.BY	BANK OF BARODA	DATE	26/03/2024

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.

Note is made of degenerative changes in visualized bones and spine

IMPRESSION: No significant abnormality is detected.

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DR.SHALINÍ GOEL

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

RMC No.: 21954

