



बैंक ऑफ बड़ोदा
Bank of Baroda
भारत का अंतर्राष्ट्रीय बैंक



नाम : मनीष टांक

Name : Manish Tank

कर्मचारी कूट क्र. : 110289


E.C. No. : 110289

सुरेन्द्र शर्मा
उप महाप्रबंधक
क्षेत्रीय प्रमुख

MANISH TANK
धारक के हस्ताक्षर
Signature of Holder

This is informed to
every staff member.



 GPS Map Camera

Jaipur, Rajasthan, India

XQ7M+R38, Central Spine Rd, Sector 2, Central Spine, Vidyadhar
Nagar, Jaipur, Rajasthan 302039, India

Lat 26.964526°

Long 75.782677°

25/11/23 01:52 PM GMT +05:30





भारत सरकार
GOVERNMENT OF INDIA



मनीष टांक

Manish Tank

जन्म तिथि/ DOB: 15/05/1989

पुरुष / MALE

Dr. PIYUSH
MBBS, D. V. RD (F)
RMC No.-0



~~0505 2121~~ 3298

आधार-आम आदमी का अधिकार

मनीष टांक



भारतीय विशिष्ट पहचान प्राधिकरण
UNIQUE IDENTIFICATION AUTHORITY OF INDIA

पता:

S/O: महावीर प्रसाद टेलर,
448, प्रताप नगर एक्षट.,
मुरलीपूरा, जयपुर, जयपुर,
राजस्थान - 302013

Address:

S/O: Mahaveer Prasad Tailor, 448,
pratap nagar ext., Murliपुरा, Jaipur,
Jaipur,
Rajasthan - 302013

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

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Aadhaar-Aam Admi ka Adhikar

मनीष टैलर



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General Physical Examination

Date of Examination: 05/11/23

Name: MANISH TANK Age: 34 YRS DOB: 15/05/1989 Sex: Male

Referred By: BANK OF BARODA

Photo ID: AADHARCAB ID #: 3098

Ht: 169 (cm)

Wt: 67 (Kg)

Chest (Expiration): 51 (cm)

Abdomen Circumference: 88 (cm)

Blood Pressure: 100/80 mm Hg PR: 78 / min RR: 18 / min Temp: Afebrile

BMI 23.5

Eye Examination: R I E 7 G I G, N I G, N C B
L I E 7 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: Manish Tank Name of Examinee: MANISH TANK

Signature Medical Examiner: Dr. Piyush Goyal Name Medical Examiner: DR. PIYUSH CHOYAL
MBBS, DM (D (Radiologist))
RMC No.-037041



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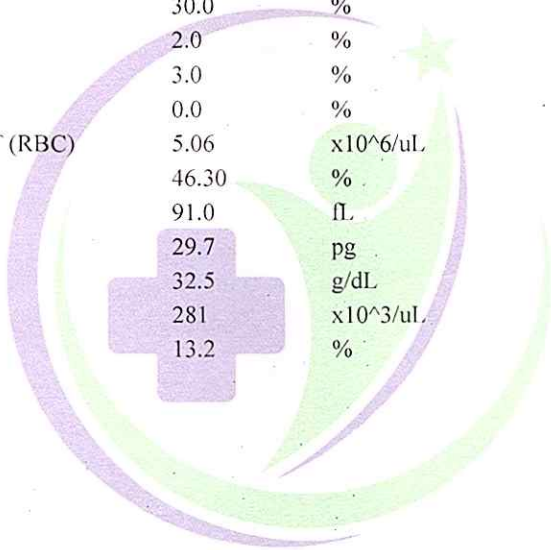
NAME :- Mr. MANISH TANK	Patient ID :-12233995	Date :- 25/11/2023	11:41:28
Age :- 34 Yrs 6 Mon 13 Days	Ref. By Doctor:-BANK OF BARODA		
Sex :- Male	Lab/Hosp :-		
	Company :-	Mr.MEDIWHEEL	

Final Authentication : 25/11/2023 17:29:52

HAEMOGARAM

HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
FULL BODY HEALTH CHECKUP BELOW 40 MALE			
HAEMOGLOBIN (Hb)	15.1	g/dL	13.0 - 17.0
TOTAL LEUCOCYTE COUNT	4.90	/cumm	4.00 - 10.00
DIFFERENTIAL LEUCOCYTE COUNT			
NEUTROPHIL	65.0	%	40.0 - 80.0
LYMPHOCYTE	30.0	%	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)	5.06	$\times 10^6/\mu\text{L}$	4.50 - 5.50
HEMATOCRIT (HCT)	46.30	%	40.00 - 50.00
MEAN CORP VOLUME (MCV)	91.0	fL	83.0 - 101.0
MEAN CORP HB (MCH)	29.7	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	32.5	g/dL	31.5 - 34.5
PLATELET COUNT	281	$\times 10^3/\mu\text{L}$	150 - 410
RDW-CV	13.2	%	11.6 - 14.0



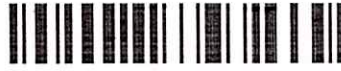
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NAME :- Mr. MANISH TANK	Patient ID :-42233995	Date :- 25/11/2023	11:41:28
Age :- 34 Yrs 6 Mon 13 Days	Ref. By Doctor:-BANK OF BARODA		
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	Company :-	Mr.MEDIWHEEL	

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HAEMATOLOGY

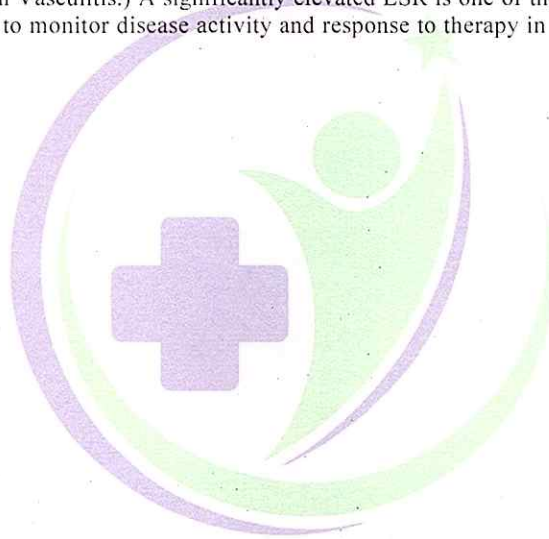
Erythrocyte Sedimentation Rate (ESR)
Method:- Westergreen

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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(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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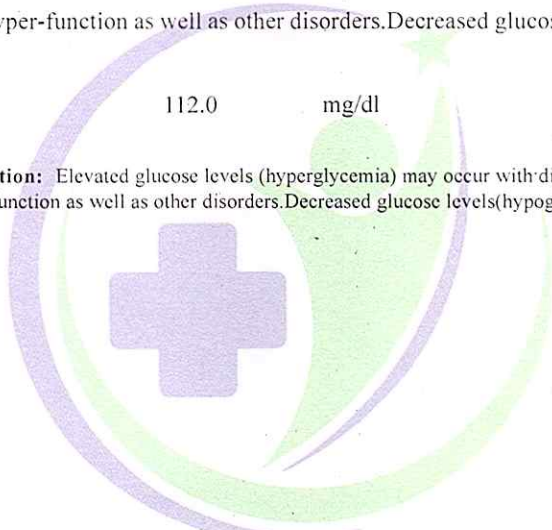
BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
FASTING BLOOD SUGAR (Plasma) Method:- GOD POD	103.0	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 .

BLOOD SUGAR PP (Plasma) Method:- GOD PAP	112.0	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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HAEMATOTOLOGY

Test Name	Value	Unit	Biological Ref Interval
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]

- Erythropoiesis**
 - Increased HbA1c: iron, vitamin B12 deficiency, decreased erythropoiesis.
 - Decreased HbA1c: administration of erythropoietin, iron, vitamin B12, reticulocytosis, chronic liver disease.
- Altered Haemoglobin-Genetic or chemical alterations in hemoglobin: hemoglobinopathies, HbF, methemoglobin, may increase or decrease HbA1c**
- Glycation**
 - Increased HbA1c: alcoholism, chronic renal failure, decreased intraerythrocytic pH.
 - Decreased HbA1c: certain hemoglobinopathies, increased intra-erythrocyte pH
- 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.
- 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

Note:

- Shortened RBC life span -HbA1c test will not be accurate when a person has a condition that affects the average lifespan of red blood cells (RBCs), such as hemolytic anemia or blood loss. When the lifespan of RBCs in circulation is shortened, the A1c result is falsely low and is an unreliable measurement of a person's average glucose over time.
- Abnormal forms of hemoglobin - The presence of some hemoglobin variants, such as hemoglobin S in sickle cell anemia, may affect certain methods for measuring A1c. In these cases, fructosamine can be used to monitor glucose control.

Advised:

- To follow patient for glycemic control test like fructosamine or glycated albumin may be performed instead.
- Hemoglobin HPLC screen to analyze abnormal hemoglobin variant.

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HAEMATOLOGY

BLOOD GROUP ABO
Method:- Haemagglutination reaction

"A" POSITIVE



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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIPID PROFILE			
TOTAL CHOLESTEROL Method:- CHOD-PAP methodology	137.00	mg/dl	Desirable <200 Borderline 200-239 High > 240
InstrumentName: MISPA PLUS Interpretation: Cholesterol measurements are used in the diagnosis and treatments of lipid lipoprotein metabolism disorders.			
TRIGLYCERIDES Method:- GPO-PAP	118.00	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	42.20	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 gives improved accuracy and reproducibility when compared to precipitation methods.			
LDL CHOLESTEROL Method:- Calculated Method	75.13	mg/dl	Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190
VLDL CHOLESTEROL Method:- Calculated	23.60	mg/dl	0.00 - 80.00
T.CHOLESTEROL/HDL CHOLESTEROL RATIO Method:- Calculated	3.25		0.00 - 4.90
LDL / HDL CHOLESTEROL RATIO Method:- Calculated	1.78		0.00 - 3.50
TOTAL LIPID Method:- CALCULATED	446.43	mg/dl	400.00 - 1000.00
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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BIOCHEMISTRY

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

Comments: 1- ATP III suggested the addition of Non HDL Cholesterol (Total Cholesterol – HDL Cholesterol) as an indicator of all atherogenic lipoproteins (mainly LDL & VLDL). The Non HDL Cholesterol is used as a secondary target of therapy in persons with triglycerides ≥ 200 mg/dL. The goal for Non HDL Cholesterol in those with increased triglyceride is 30 mg/dL above that set for LDL Cholesterol.

2 -For calculation of CHD risk, history of smoking, any medication for hypertension & current B.P. levels are required



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BIOCHEMISTRY

LIVER PROFILE WITH GGT

SERUM BILIRUBIN (TOTAL) Method:- DMSO/Diazo	0.63	mg/dL	Infants : 0.2-8.0 mg/dL Adult - Up to - 1.2 mg/dL
SERUM BILIRUBIN (DIRECT) Method:- DMSO/Diazo	0.25	mg/dL	Up to 0.40 mg/dL
SERUM BILIRUBIN (INDIRECT) Method:- Calculated	0.38	mg/dl	0.30-0.70
SGOT Method:- IFCC	26.5	U/L	0.0 - 40.0
SGPT Method:- IFCC	34.4	U/L	0.0 - 40.0
SERUM ALKALINE PHOSPHATASE Method:- DGKC - SCE	86.50	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.	25.60	U/L	10.00 - 45.00
SERUM TOTAL PROTEIN Method:- Direct Biuret Reagent	6.52	g/dl	6.00 - 8.40
SERUM ALBUMIN Method:- Bromocresol Green	4.25	g/dl	3.50 - 5.50
SERUM GLOBULIN Method:- CALCULATION	2.27	gm/dl	2.20 - 3.50
A/G RATIO	1.87		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 32.20 mg/dl 10.00 - 50.00
Method:- Urease/GLDH

InstrumentName: HORIBA CA 60 **Interpretation :** Urea measurements are used in the diagnosis and treatment of certain renal and metabolic diseases.

SERUM CREATININE 1.09 mg/dl Males : 0.6-1.50 mg/dl
Method:- Jaffe's Method 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 6.81 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 140.0 mmol/L 135.0 - 150.0
Method:- ISE

POTASSIUM 4.44 mmol/L 3.50 - 5.50
Method:- ISE

CHLORIDE 101.2 mmol/L 94.0 - 110.0
Method:- ISE

SERUM CALCIUM 9.52 mg/dL 8.80 - 10.20
Method:- Arsenazo III Method

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 6.52 g/dl 6.00 - 8.40
Method:- Direct Biuret Reagent

SERUM ALBUMIN 4.25 g/dl 3.50 - 5.50
Method:- Bromocresol Green

SERUM GLOBULIN 2.27 gm/dl 2.20 - 3.50
Method:- CALCULATION

A/G RATIO 1.87 1.30 - 2.50

Interpretation : Measurements obtained by this method are used in the diagnosis and treatment of a variety of disorders of liver, kidney and

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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-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.



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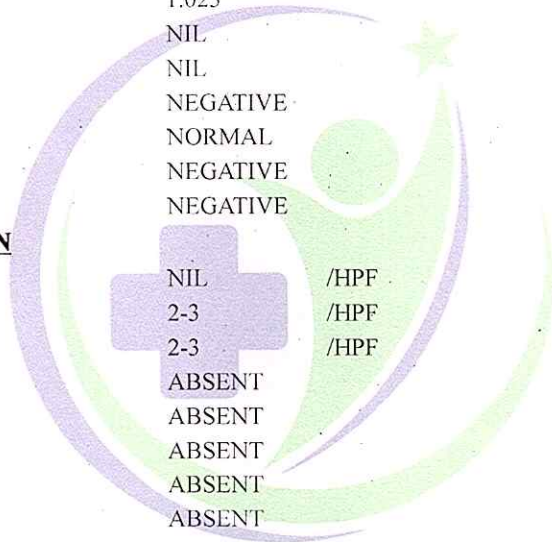
📍 B-14, Vidhyadhar Enclave-II, Near Axix Bank
Central Spine, Vidhyadhar Nagar, Jaipur - 302023
☎ +91 141 4824885 📧 maxcarediagnostics1@gmail.com

NAME :- Mr. MANISH TANK	Patient ID :-12233995	Date :- 25/11/2023	11:41:28
Age :- 34 Yrs 6 Mon 13 Days	Ref. By Doctor:-BANK OF BARODA		
Sex :- Male	Lab/Hosp :-		
	Company :- Mr.MEDIWHEEL		

Final Authentication : 25/11/2023 17:29:52

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	1.025		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



Technologist
VIKARANTSI
Page No: 12 of 16

Tanu

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



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	Company :- Mr.MEDIWHEEL		

Final Authentication : 25/11/2023 17:29:52

CLINICAL PATHOLOGY

URINE SUGAR (FASTING)
Collected Sample Received

Nil

Nil



Technologist
VIKARANTSI
Page No: 13 of 16

Tanu
DR.TANU RUNGTA
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Final Authentication : 25/11/2023 17:29:52

CLINICAL PATHOLOGY

STOOL ANALYSIS

PHYSICAL EXAMINATION

MUCUS

BLOOD

MICROSCOPIC EXAMINATION

RBC's

/HPF

WBC/HPF

/HPF

OVA

CYSTS

OTHERS

Collected Sample Received



Technologist
VIKARANTSI
Page No: 14 of 16

Tanu

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Final Authentication : 25/11/2023 17:29:52

TOTAL THYROID PROFILE

IMMUNOASSAY

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

THYROID-TRIIODOTHYRONINE T3 Method:- ECLIA	1.15	ng/mL	0.70 - 2.04
--	------	-------	-------------

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. Transient increase in TSH levels or abnormal TSH levels can be seen in some non thyroidal conditions, simultaneous measurement of TSH with free T4 is useful in evaluating differential diagnosis

INTERPRETATION-Ultra Sensitive 4th generation assay 1.Primary hyperthyroidism is accompanied by ↑serum T3 & T4 values along with ↑ TSH level 2.Low TSH, high FT4 and TSH receptor antibody (TRAb) +ve seen in patients with Graves disease 3.Low TSH, high FT4 and TSH receptor antibody (TRAb) -ve seen in patients with Toxic adenoma/Toxic Multinodular goiter 4.HighTSH, Low FT4 and Thyroid microsomal antibody increased seen in patients with Hashimotos thyroiditis 5.HighTSH, Low FT4 and Thyroid microsomal antibody normal seen in patients with Iodine deficiency/Congenital T4 synthesis deficiency 6.Low TSH, Low FT4 and TRH stimulation test -Delayed response seen in patients with Tertiary hypothyroidism 7.Primary hypothyroidism is accompanied by ↓ serum T3 and T4 values & ↑serum TSH levels 8.Normal T4 levels accompanied by ↑ T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis 9.Normal or ↑ T3 & ↑ T4 10.Normal T3 & T4 along with ↑ TSH indicate mild / Subclinical Hyperthyroidism 11.Normal T3 & ↑ T4 along with ↑ TSH is seen in Hypothyroidism 12.Normal T3 & T4 levels with ↑ TSH indicate Mild / Subclinical Hypo

DURING 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.

REMARK-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. Abnormal thyroid test findings often found in critically ill patients should be repeated after the critical nature of the condition is resolved. 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.

THYROID-THYROXINE (T4) Method:- ECLIA	0.89	uIU/mL	5.10 - 14.10
---	------	--------	--------------

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. Transient increase in TSH levels or abnormal TSH levels can be seen in some non thyroidal conditions, simultaneous measurement of TSH with free T4 is useful in evaluating differential diagnosis

INTERPRETATION-Ultra Sensitive 4th generation assay 1.Primary hyperthyroidism is accompanied by ↑serum T3 & T4 values along with ↑ TSH level. 2.Low TSH, high FT4 and TSH receptor antibody (TRAb) +ve seen in patients with Graves disease 3.Low TSH, high FT4 and TSH receptor antibody (TRAb) -ve seen in patients with Toxic adenoma/Toxic Multinodular goiter 4.HighTSH, Low FT4 and Thyroid microsomal antibody increased seen in patients with Hashimotos thyroiditis 5.HighTSH, Low FT4 and Thyroid microsomal antibody normal seen in patients with Iodine deficiency/Congenital T4 synthesis deficiency 6.Low TSH, Low FT4 and TRH stimulation test -Delayed response seen in patients with Tertiary hypothyroidism 7.Primary hypothyroidism is accompanied by ↓ serum T3 and T4 values & ↑serum TSH levels 8.Normal T4 levels accompanied by ↑ T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis 9.Normal or ↑ T3 & ↑ T4 10.Normal T3 & T4 along with ↑ TSH indicate mild / Subclinical Hyperthyroidism 11.Normal T3 & ↑ T4 along with ↑ TSH is seen in Hypothyroidism 12.Normal T3 & T4 levels with ↑ TSH indicate Mild / Subclinical Hypo

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REMARK-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. Abnormal thyroid test findings often found in critically ill patients should be repeated after the critical nature of the condition is resolved. 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.

TSH Method:- ECLIA	1.798	uIU/mL	0.350 - 5.500
------------------------------	-------	--------	---------------

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. Transient increase in TSH levels or abnormal TSH levels can be seen in some non thyroidal conditions, simultaneous measurement of TSH with free T4 is use:

Tanu

Technologist
VIKARANTSI
Page No: 15 of 16

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Sex :- Male	Lab/Hosp :-		
	Company :- Mr.MEDIWHEEL		

Final Authentication : 25/11/2023 17:29:52

IMMUNOASSAY

evaluating differential diagnosis

INTERPRETATION-Ultra Sensitive 4th generation assay

- 1.Primary hyperthyroidism is accompanied by ↑serum T3 & T4 values along with ↓ TSH level.
- 2.Low TSH,high FT4 and TSH receptor antibody(TRAb) +ve seen in patients with Graves disease
- 3.Low TSH,high FT4 and TSH receptor antibody(TRAb) -ve seen in patients with Toxic adenoma/Toxic Multinodular goiter
- 4.HighTSH,Low FT4 and Thyroid microsomal antibody increased seen in patients with Hashimoto's thyroiditis
- 5.HighTSH,Low FT4 and Thyroid microsomal antibody normal seen in patients with Iodine deficiency/Congenital T4 synthesis deficiency
- 6.Low TSH,Low FT4 and TRH stimulation test -Delayed response seen in patients with Tertiary hypothyroidism
- 7.Primary hypothyroidism is accompanied by ↓ serum T3 and T4 values & ↑serum TSH levels
- 8.Normal T4 levels accompanied by ↓ T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis
- 9.Normal or ↓ T3 & ↑T4 levels indicate T4 Thyrotoxicosis (problem is conversion of T4 to T3)
- 10.Normal T3 & T4 along with ↓ TSH indicate mild / Subclinical Hyperthyroidism .
- 11.Normal T3 & ↓ T4 along with ↑ TSH is seen in Hypothyroidism .
- 12.Normal T3 & T4 levels with ↑ TSH indicate Mild / Subclinical Hypothyroidism
- 13.Slightly ↑ T3 levels may be found in pregnancy and in estrogen therapy while ↓ levels may be encountered in severe illness , malnutrition , renal failure and during therapy with drugs like propranolol.
- 14.Although ↑ TSH levels are nearly always indicative of Primary Hypothyroidism ,rarely they can result from TSH secreting pituitary tumours.

DURING 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.

REMARK-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. Abnormal thyroid test findings often found in critically ill patients should be repeated after the critical nature of the condition is resolved.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.

*** End of Report ***

Technologist
VIKARANTSI
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Tanu
DR.TANU RUNGTA
MD (Pathology)
RMC No. 17226



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MR. MANISH TANK	34 Y/M
Registration Date: 25/11/2023	Ref. by: BANK OF BARODA

ULTRASOUND OF WHOLE ABDOMEN

Liver is of normal size (111 mm) with bright parenchymal 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. 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. 84 mm.

Left kidney is measuring approx. 89 mm.

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

Prostate is normal in size with normal echotexture and outline.

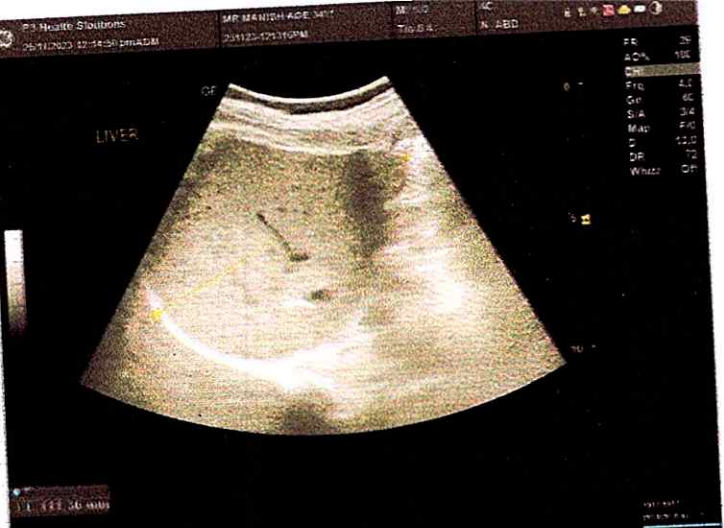
No enlarged nodes are visualized. No retro-peritoneal lesion is identified.

No significant free fluid is seen in pelvis.

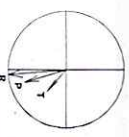
IMPRESSION:

- Grade I hepatic steatosis.
- No free fluid or lymphadenopathy.

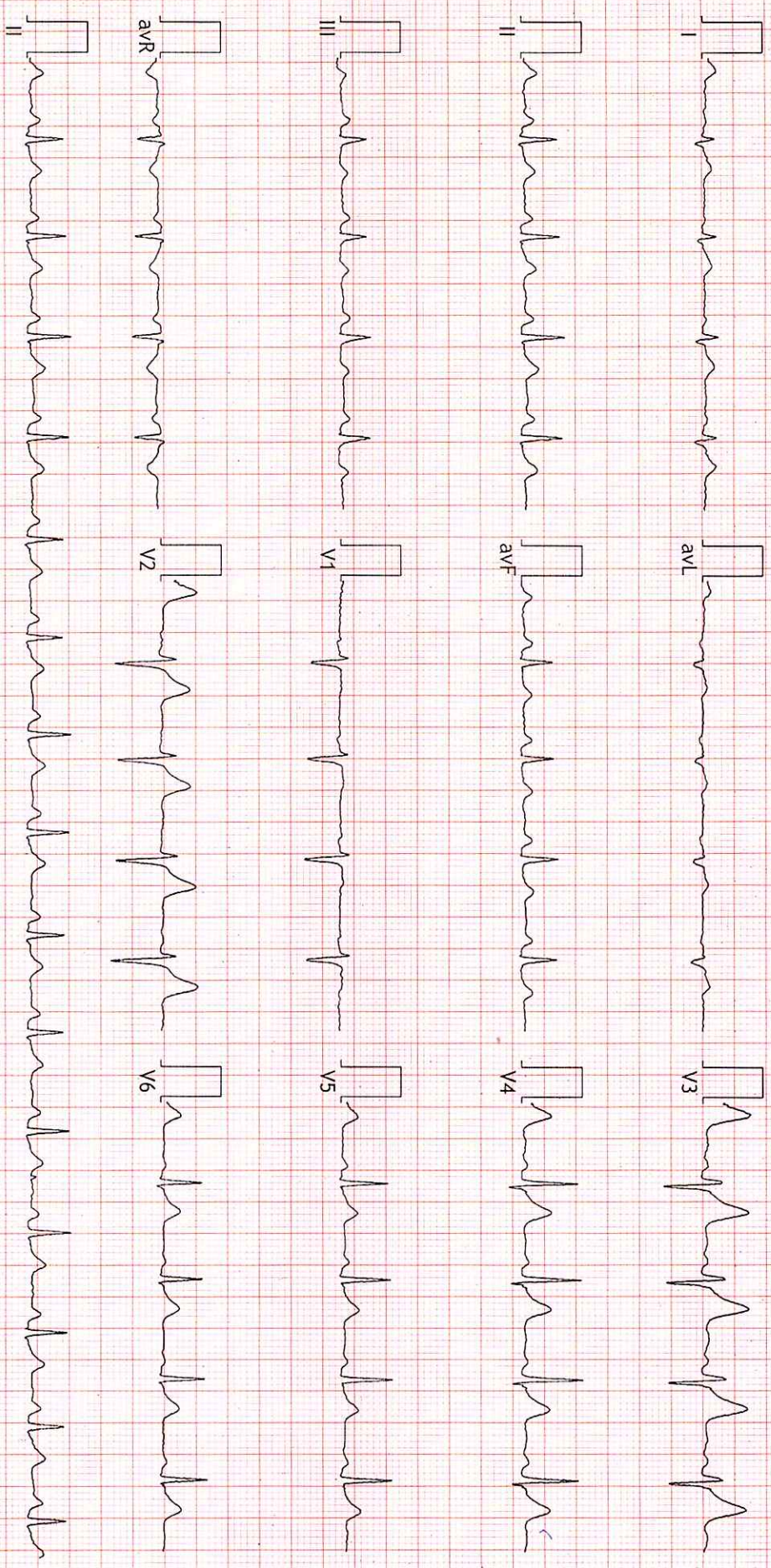
Dr. Mukesh Sharma
M.B.B.S; M.D. (Radiodiagnosis)
RMC No. 43418/17437



Tem's (P) Ltd
#P3 HEALTH SOLUTIONS LLP B-14, Vidhyadhar nahar , Jaipur
 12233995/Mr Manish Tank 34Yrs/Male Kgs/ Cms BP: ___/___ mmHg
 Ref.: BANK OF BARODA Test Date: 25-Nov-2023(1:20:27 P) Notch: 50Hz 0.05Hz - 35Hz 10mm/mV 25mm/Sec



PR Interval: 116 ms
 QRS Duration: 112 ms
 QT/QTc: 323/398ms
 P-QRS-T Axis: 73 - 85 - 41 (Deg)



FINDINGS: Normal Sinus Rhythm
 Vent Rate : 91 bpm; PR Interval : 116 ms; QRS Duration : 112 ms; QT/QTc Int : 323/398 ms
 P-QRS-T axis: 73 • 85 • 41 • (Deg)
 Comments :

TUNL

Handwritten signature

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

Dr. NARESH MOHINKA

P3 HEALTH SOLUTIONS LLP

B-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur
 12233995/W/ MANISH TANK 34 Yrs/Male 0 Kg/0 Cms
 Date: 25-Nov-2023 01:30:04 PM
 Ref By : BANK OF BARODA
 Medication : Nil

Summary

Protocol : BRUCE
 History : Nil

Stage	StageTime (min:sec)	PhaseTime (min:sec)	Speed (mph)	Grade (%)	METS	H.R. (bpm)	B.P. (mmHg)	R.P.P. (x100)	PVC	Comments
Supine					1.0	102	130/85	132	-	
Standing					1.0	102	130/85	132	-	
HV					1.0	107	130/85	139	-	
EXStart					1.0	103	130/85	133	-	
Stage 1	3:01	3:02	1.7	10.0	4.7	129	140/85	180	-	
Stage 2	3:01	6:02	2.5	12.0	7.1	150	150/90	225	-	
PeakEx	1:07	7:08	3.4	14.0	8.3	168	160/90	268	-	
Recovery	1:00		0.0	0.0	1.2	129	160/90	206	-	
Recovery	2:00		0.0	0.0	1.0	119	160/90	190	-	
Recovery	3:00		0.0	0.0	1.0	113	150/90	169	-	
Recovery	4:00		0.0	0.0	1.0	112	140/85	156	-	

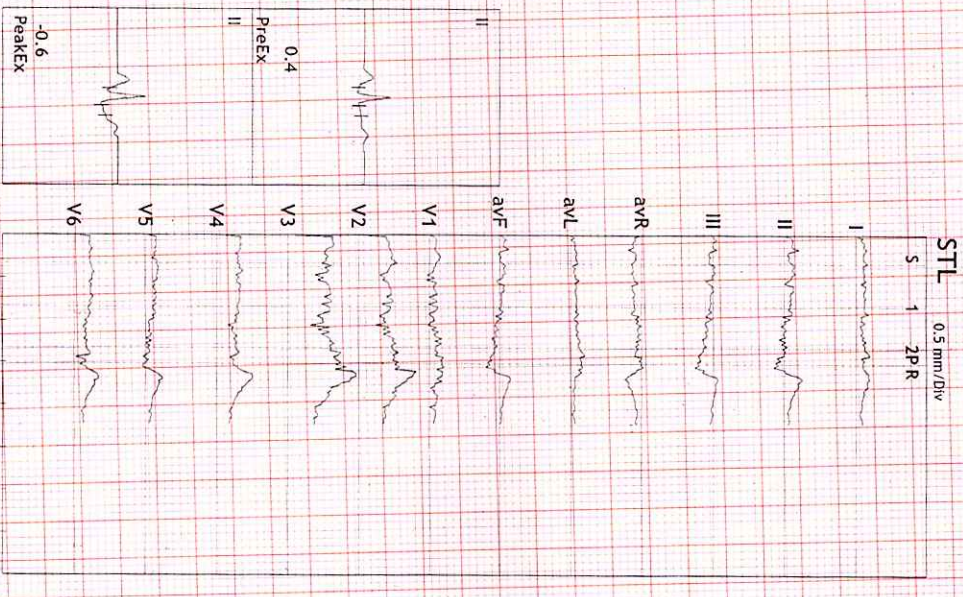
Findings :

Exercise Time : 07:07
 Max HR Attained : 168 bpm 90% of Max Predictable HR 186
 Max BP : 160/90(mmHg)
 Max Workload attained : 8.3(Fair Effort Tolerance)

Handwritten signature

TTT B Negative 408 RM I.

Advice/Comments:



Dr. Nareesh Kumar Mohanka
 MBBS, DIP. CARDIO (ESCORTS)
 D.E.M (RCGP-UK)



DR. NARESH MOHINKA

B-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur
 12233995/Mr MANISH TANK
 34 Yrs/Male
 0 Kg/0 Cms
 Date: 25-Nov-2023 01:30:04 PM
 4X 60 ms Post J

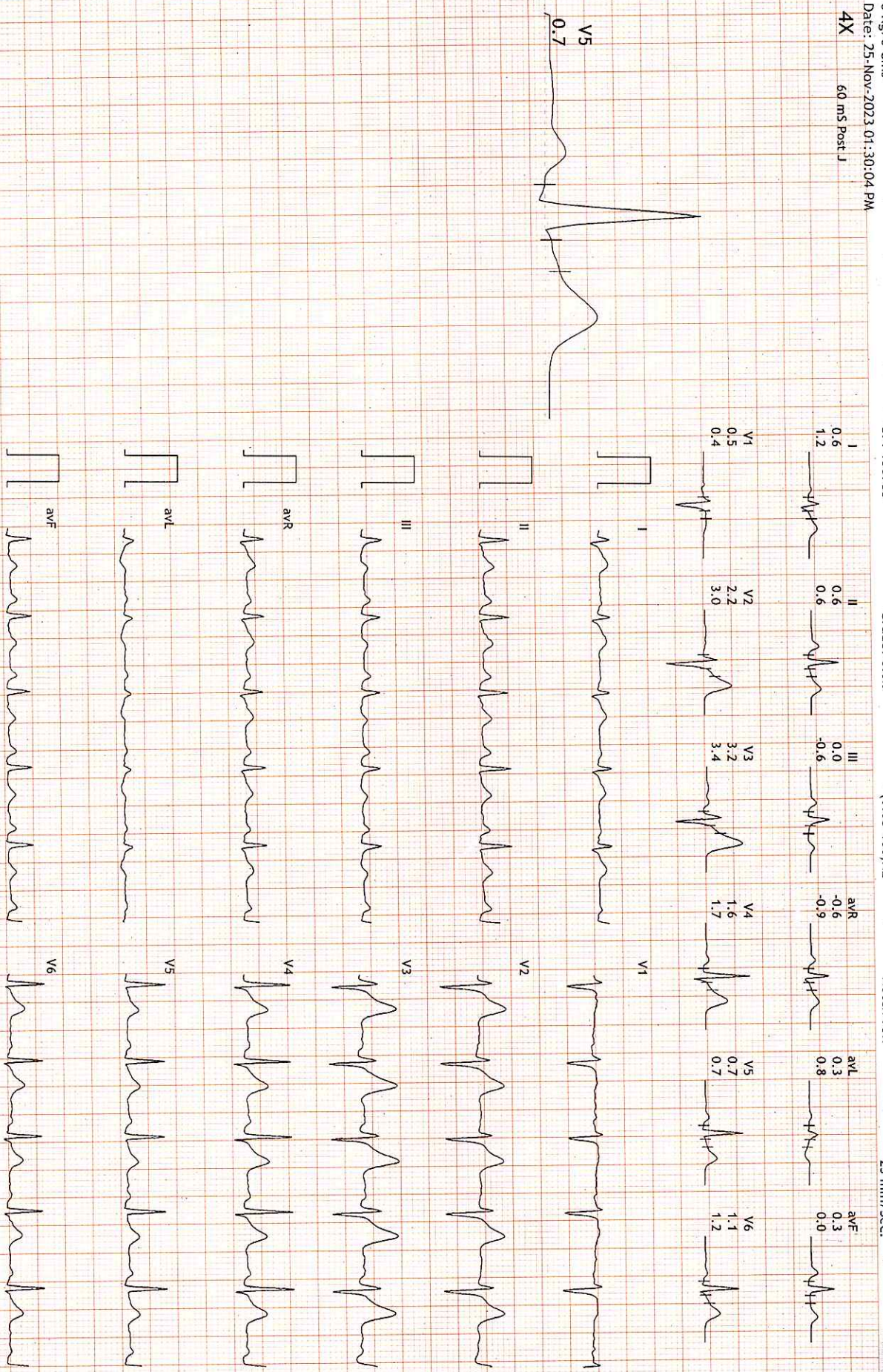
HR: 102 bpm
 METS: 1.0
 BP: 130/85

MPPR: 54% of 186
 Speed: 0.0 mph
 Grade: 0.0%

Raw ECG
 BRUCE
 (0.05-100)Hz

Ex Time 00:31
 BLC :On
 Notch :On

Supine
 10.0 mm/mV
 25 mm/Sec.



4X

60 ms Post J

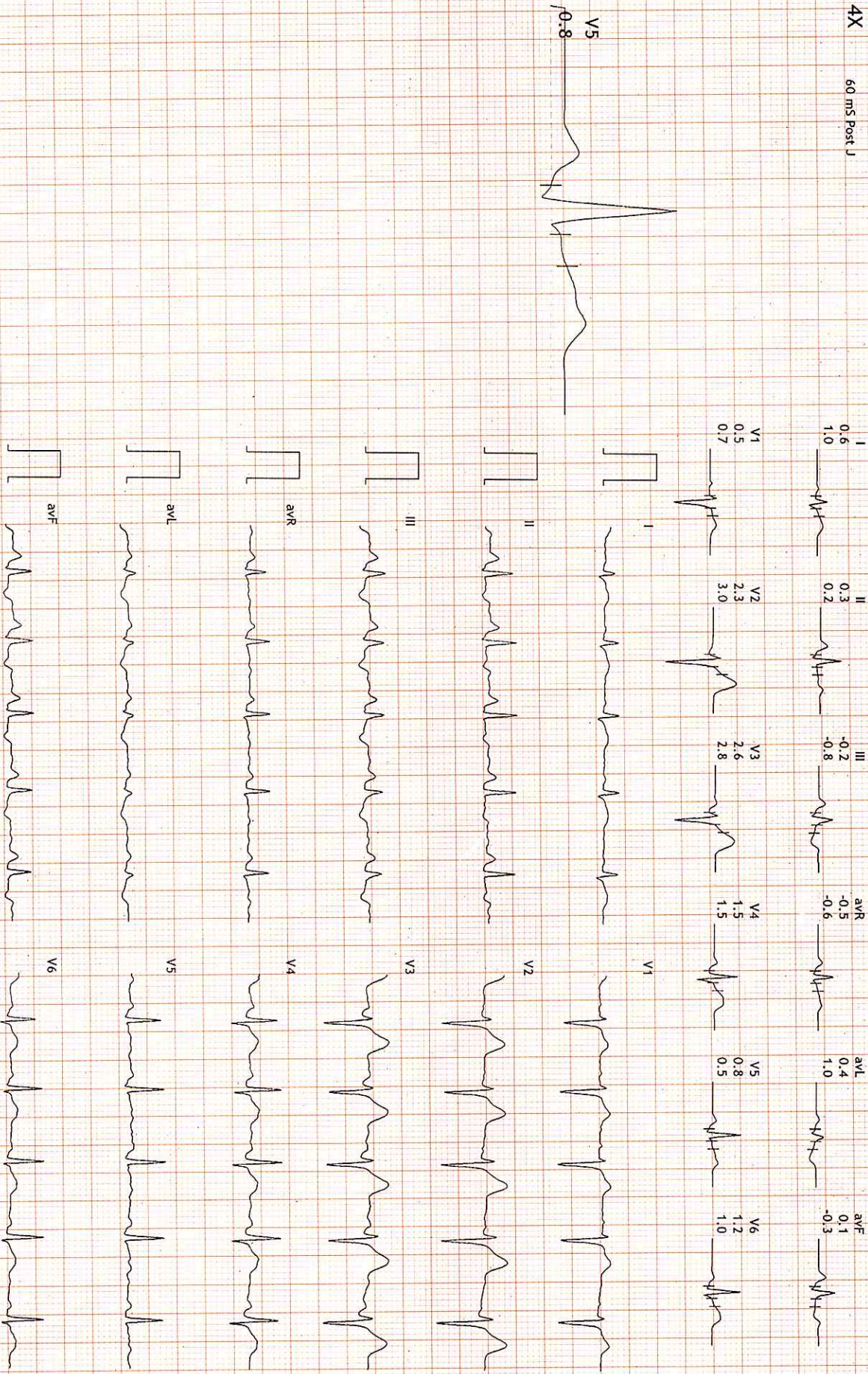
HR: 108 bpm
METs: 1.0
BP: 130/85

MPHR: 58% of 186
Speed: 0.0 mph
Grade: 0.0%

Raw ECG
BRUCE
(0.05-100)Hz

EX Time 01:14
BLC: On
Notch: On

Standing
10.0 mm/mv
25 mm/Sec.



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B-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur

12233995/MR MANISH TANK

34 Yrs/Male

0 Kg/0 Cms

Date: 25-Nov-2023 01:30:04 PM

12 Lead + Median

HR: 107 bpm
METs: 1.0
BP: 130/85

MPHR: 57% of 186
Speed: 0.0 mph
Grade: 0.0%

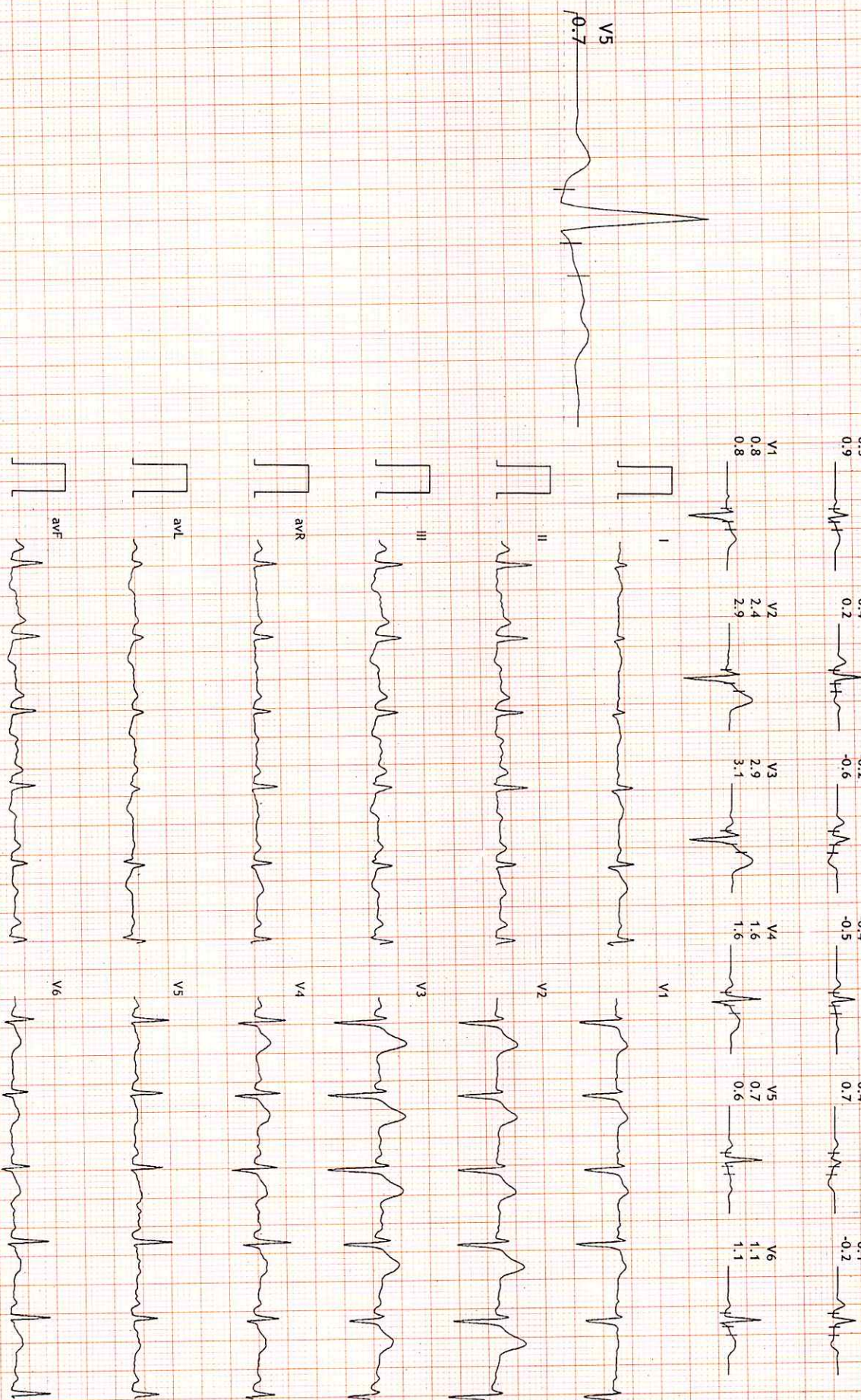
Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 01:34
BLC : On
Notch : On

HV
10.0 mm/mV
25 mm/Sec.



4X 60 mS Post J



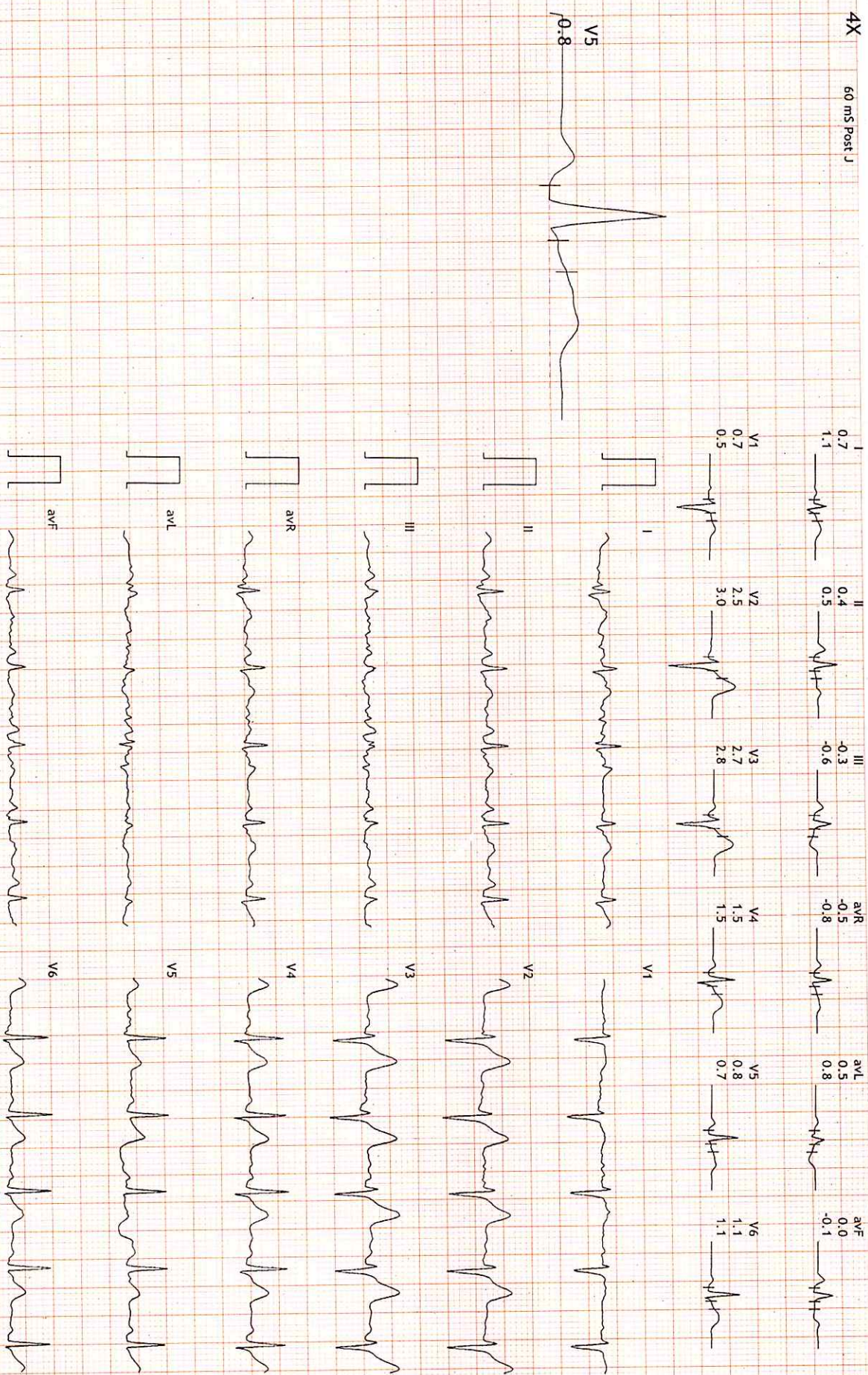
HR: 103 bpm
METs: 1.0
BP: 130/85

MPHR: 55% of 186
Speed: 0.0 mph
Grade: 0.0%

Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 01:57
BLC : On
Notch : On

ExStart
10.0 mm/mV
25 mm/Sec.



HR: 130 bpm
METS: 4.7
BP: 140/85

MPHR: 69% of 186
Speed: 1.7 mph
Grade: 10.0%

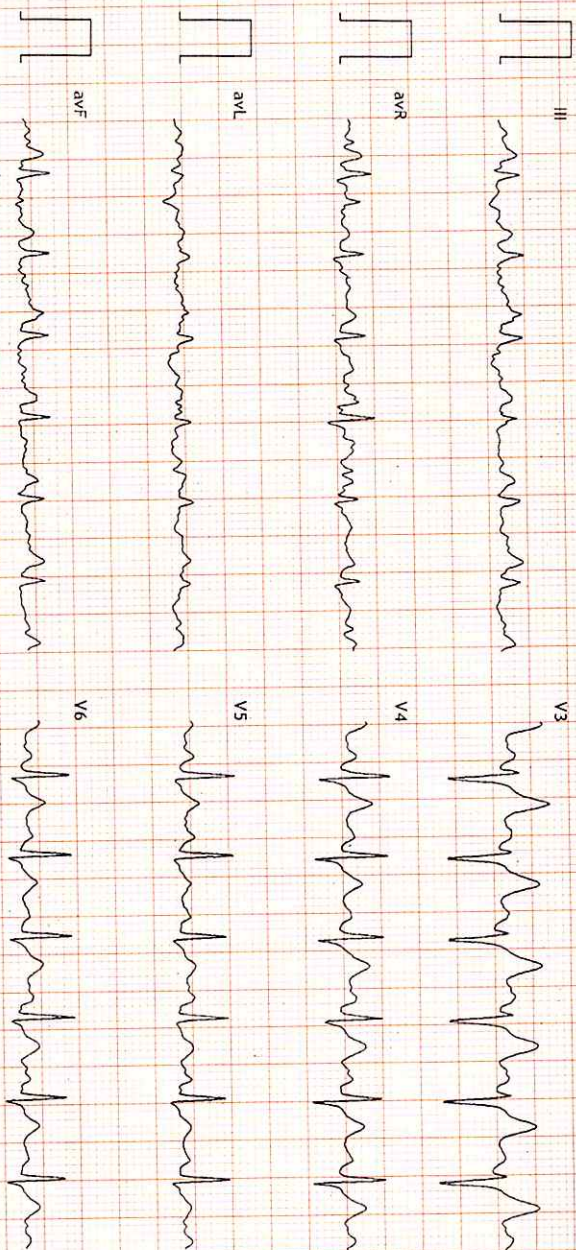
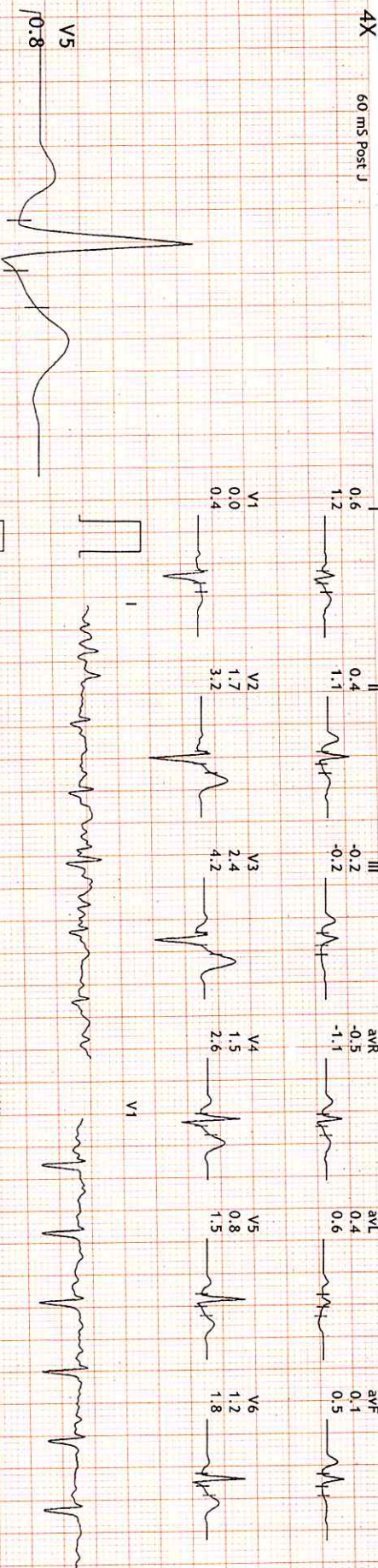
Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 02:59
BLC : On
Notch : On

BRUCE: Stage 1(3:00)
10.0 mm/mV
25 mm/Sec.



4X 60 ms Post J



HR: 150 bpm
METs: 7.1
BP: 150/90

MPHR: 80% of 186
Speed: 2.5 mph
Grade: 12.0%

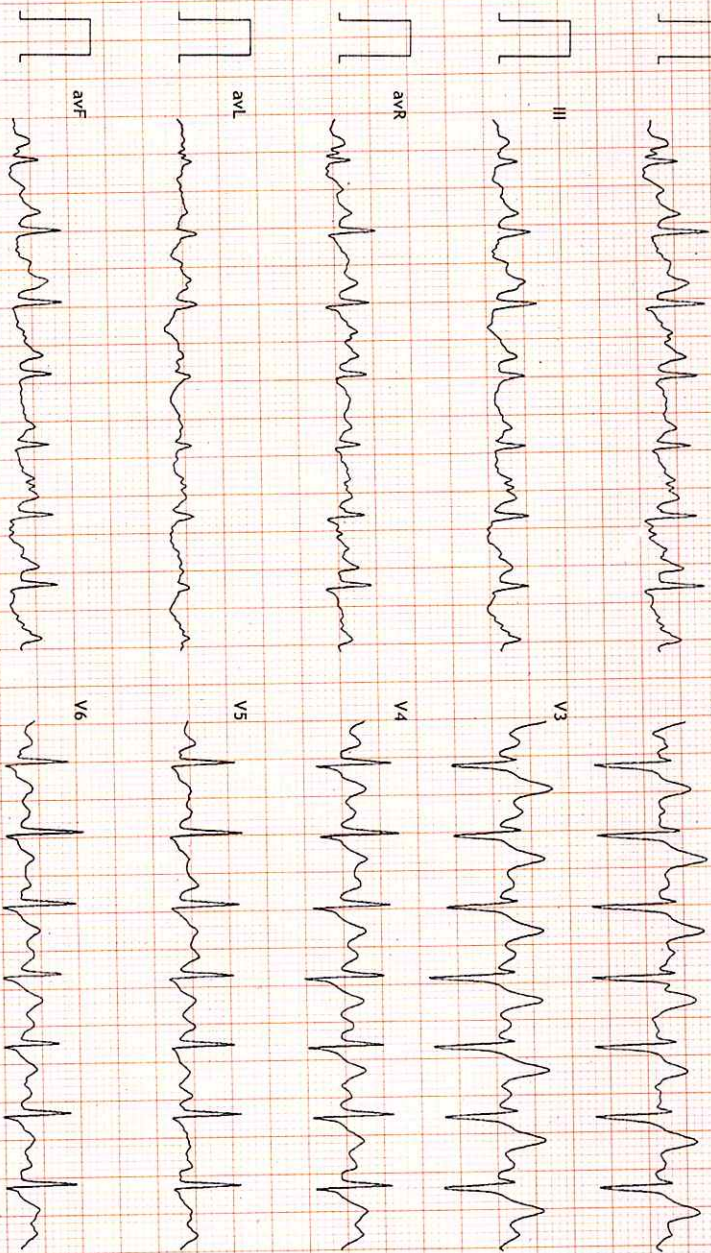
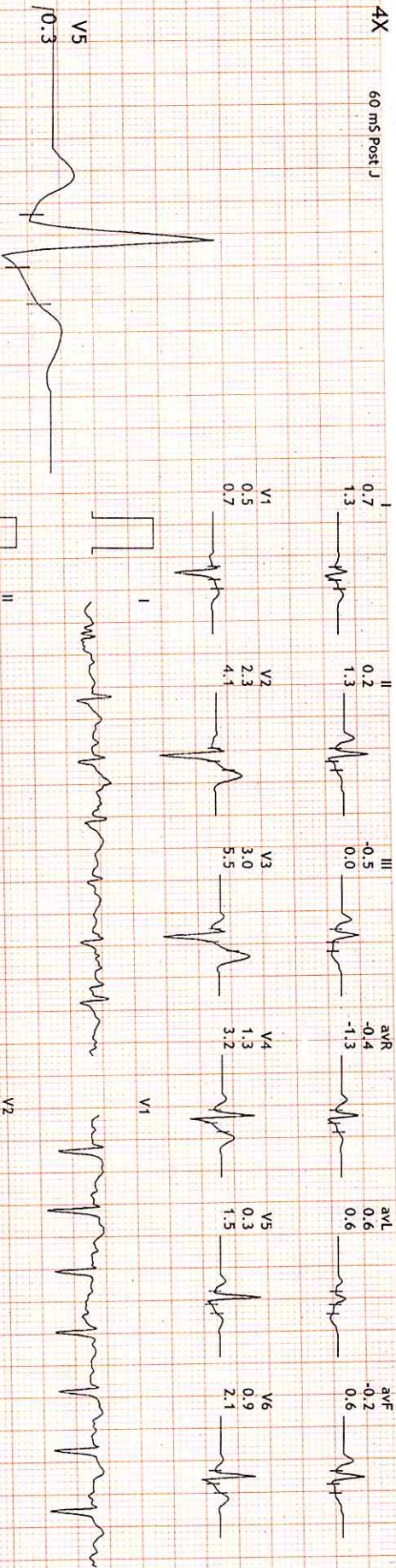
Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 05:59
BLC : On
Notch : On

BRUCE: Stage 2(3:00)
10.0 mm/mv
25 mm/Sec.



4X 60 ms Post J



P3 HEALTH SOLUTIONS LLP
 B-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur
 12233995/Mr MAANISH TANK
 34 Yrs/Male
 0 Kg/0 Cms
 Date: 25-NOV-2023 01:30:04 PM

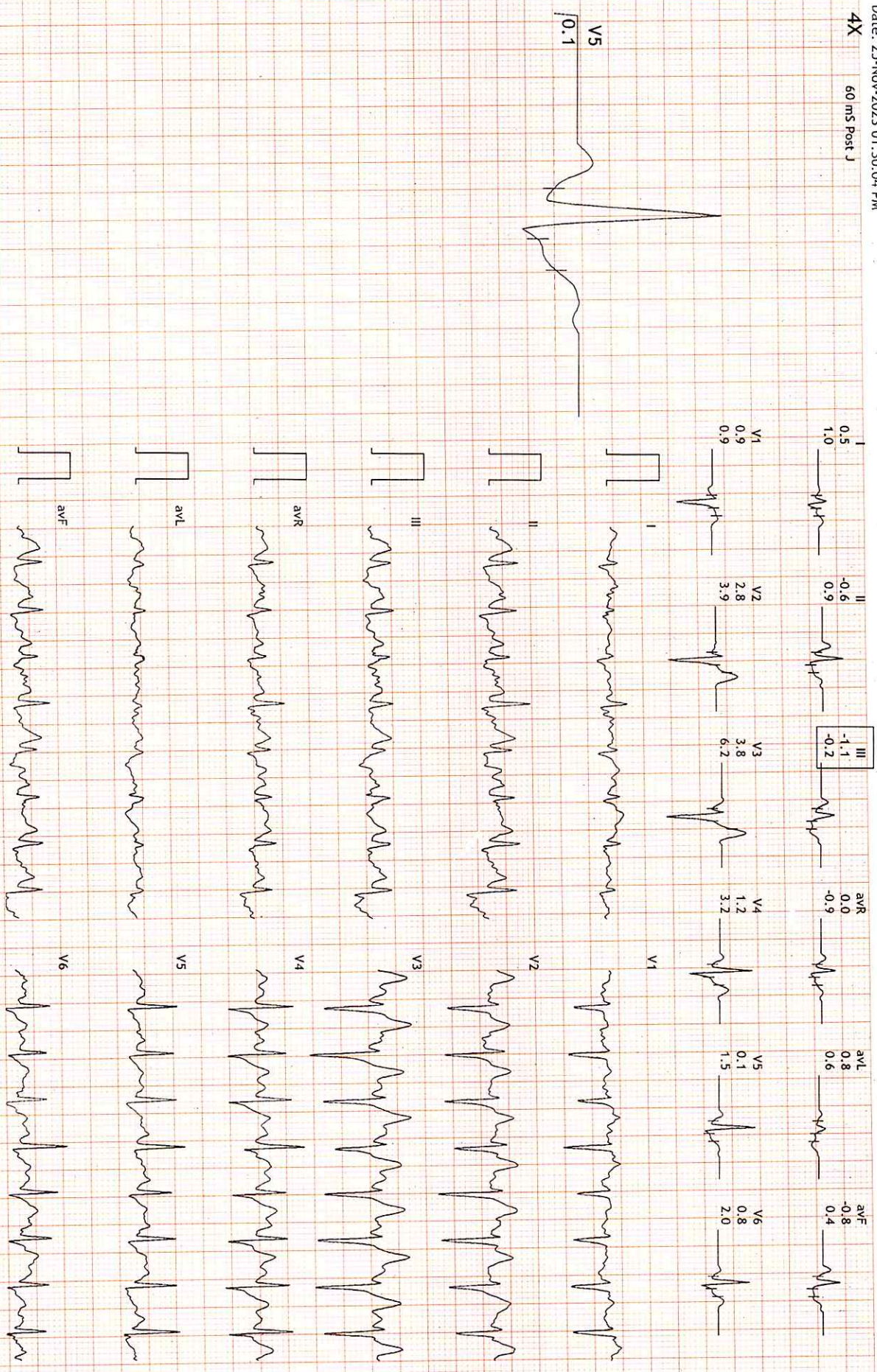
12 Lead + Median

4X 60 ms Post J

HR: 168 bpm
 METS: 8.3
 BP: 160/90
 MPRR: 90% of 186
 Speed: 3.4 mph
 Grade: 14.0%

Raw ECG
 BRUCE
 (0.05-100)Hz
 Ex Time 07:05
 BLC : On
 Notch : On

BRUCE:PeakX(1:05)
 10.0 mm/mV
 25 mm/Sec.



HR: 130 bpm
METs: 1.3
BP: 160/90

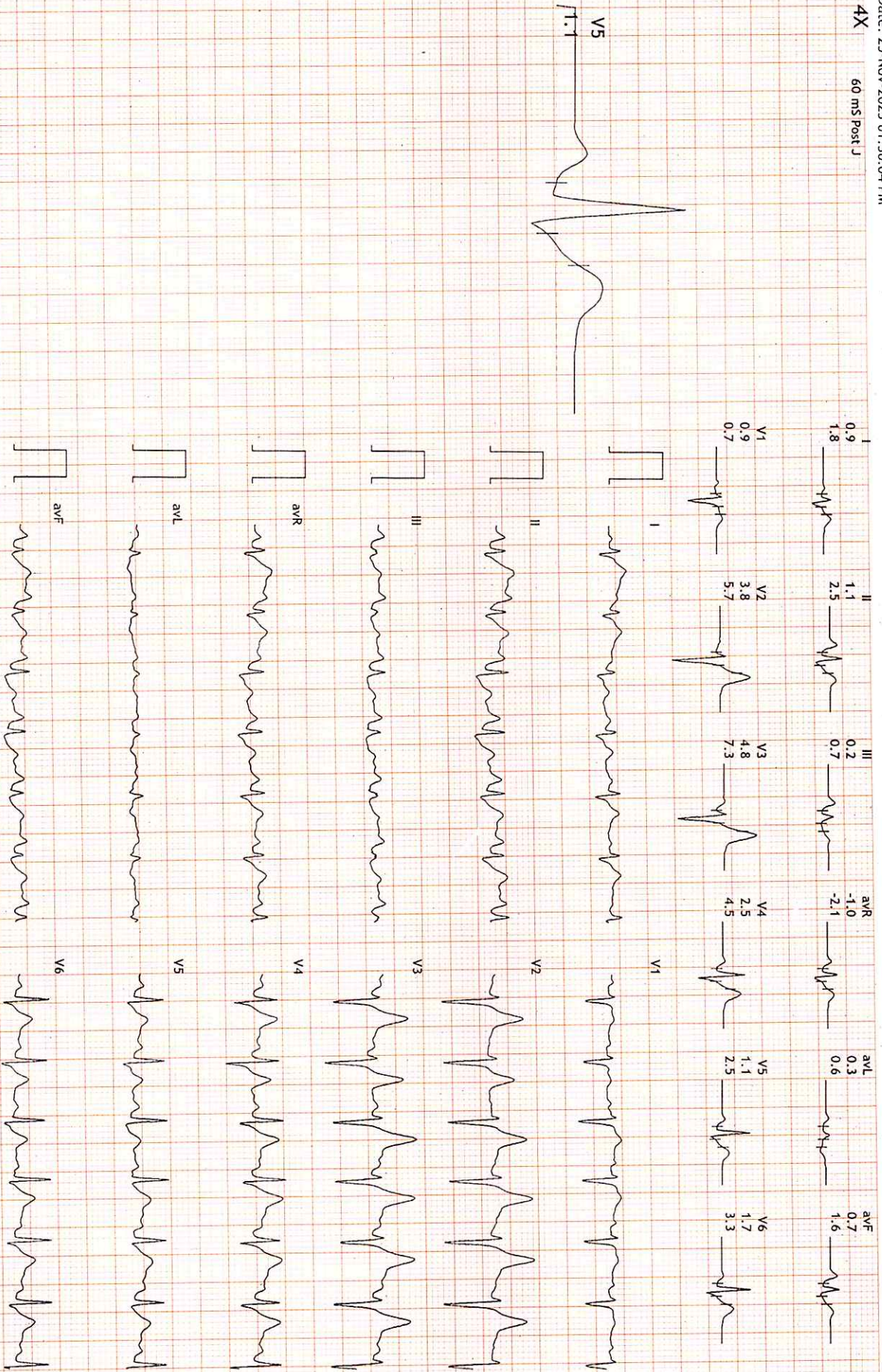
MPHR: 69% of 186
Speed: 0.0 mph
Grade: 0.0%

Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 07:07
BLC : On
Notch : On

Recovery(1:00)
10.0 mm/mV
25 mm/Sec.

4X 60 mS Post J



HR: 119 bpm
METs: 1.0
BP: 160/90

MPHR: 63% of 186
Speed: 0.0 mph
Grade: 0.0%

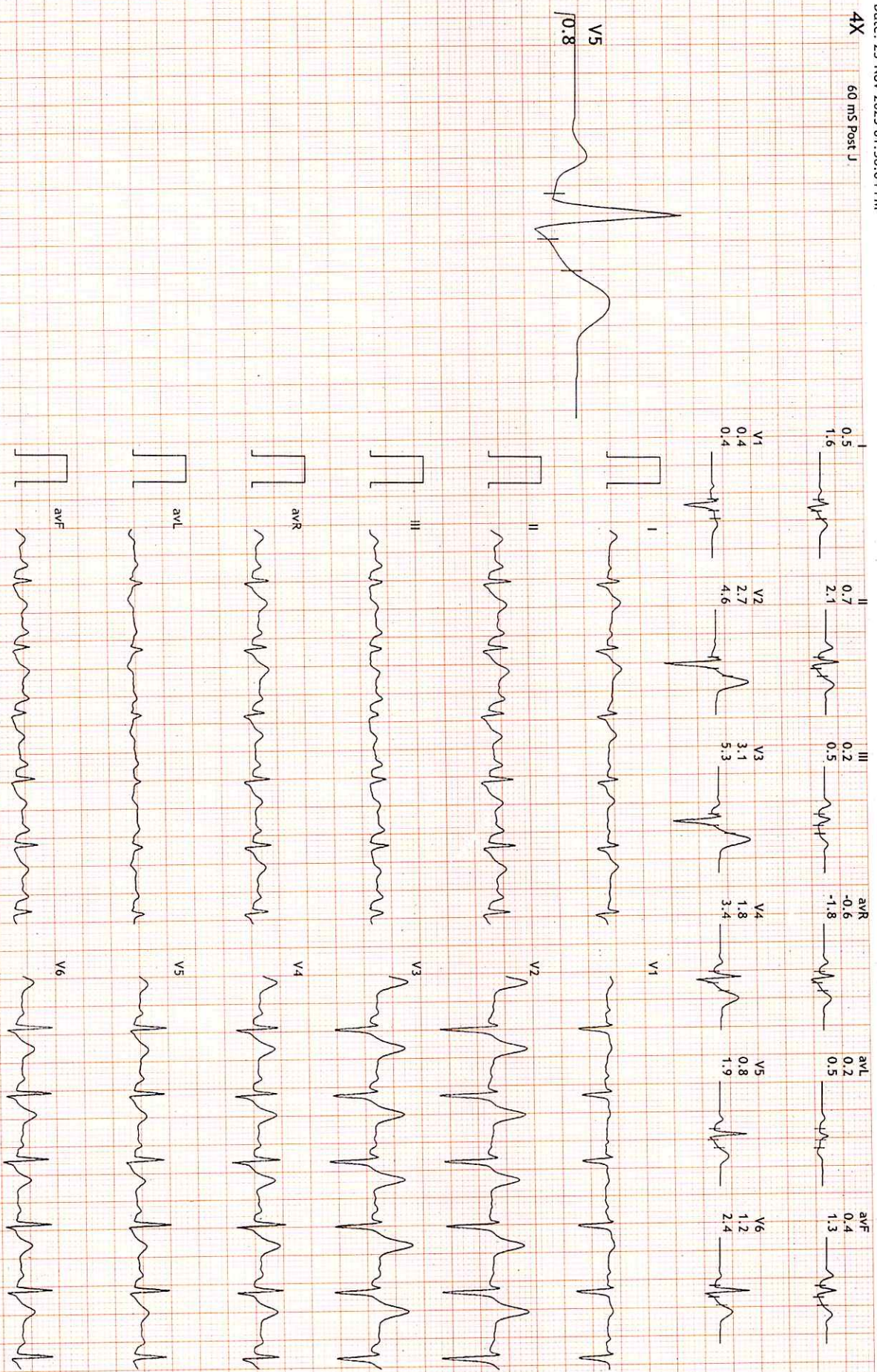
Raw ECG
BRUCE
(0.05-100)Hz

EX Time 07:07
BLC : On
Notch : On

Recovery(2:00)
10.0 mm/mv
25 mm/Sec.



4X 60 ms Post J



HR: 113 bpm

MEFS: 1.0

BP: 150/90

MPHR: 60% of 186

Speed: 0.0 mph

Grade: 0.0%

Raw ECG

BRUCE

(0.05-100)Hz

Ex Time 07:07

BLC :On

Notch :On

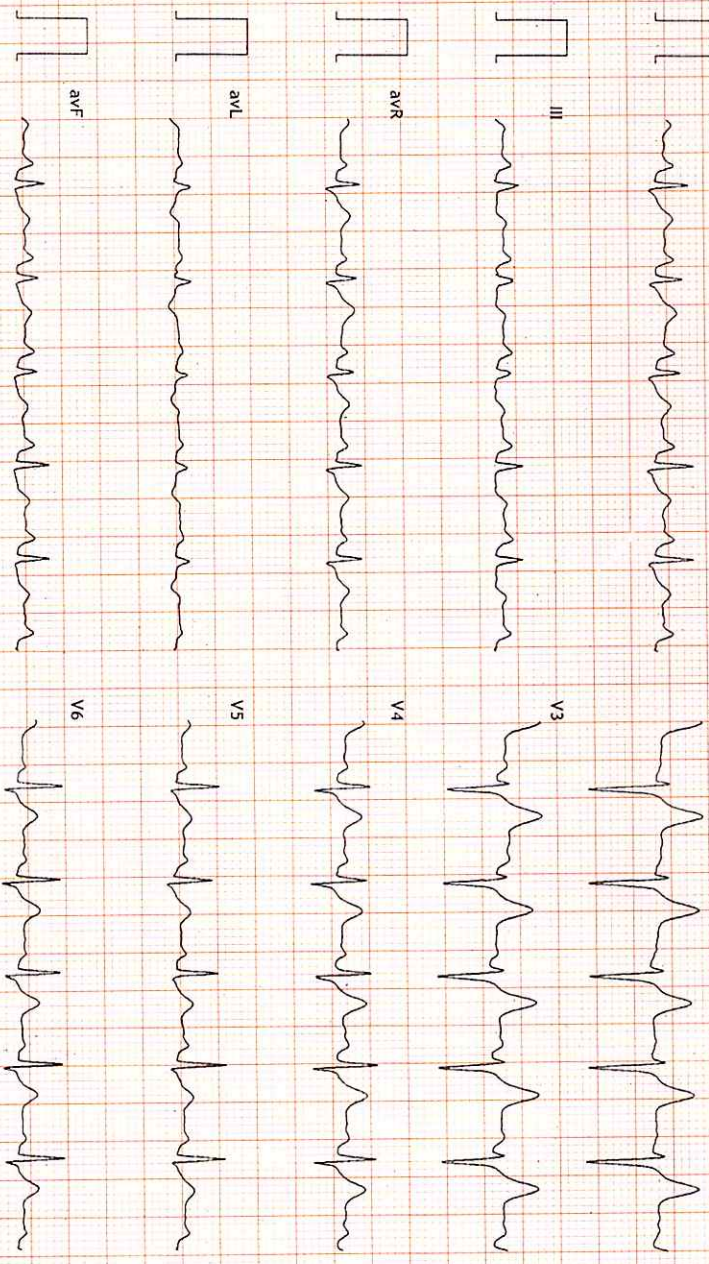
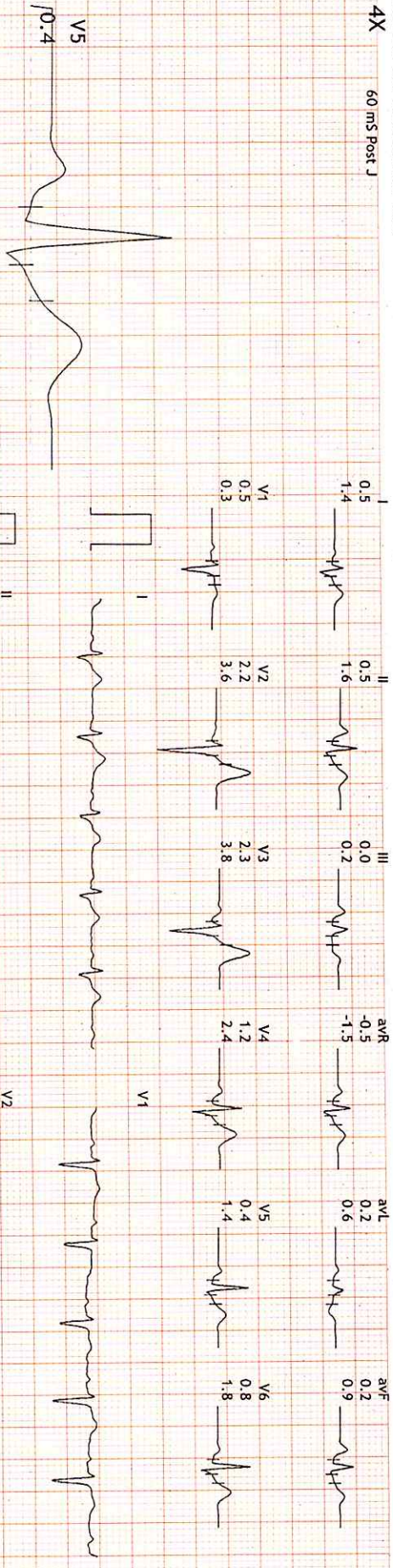
Recovery(3:00)

10.0 mm/mV

25 mm/Sec.



4X 60 ms Post J



HR: 113 bpm
METs: 1.0
BP: 140/85

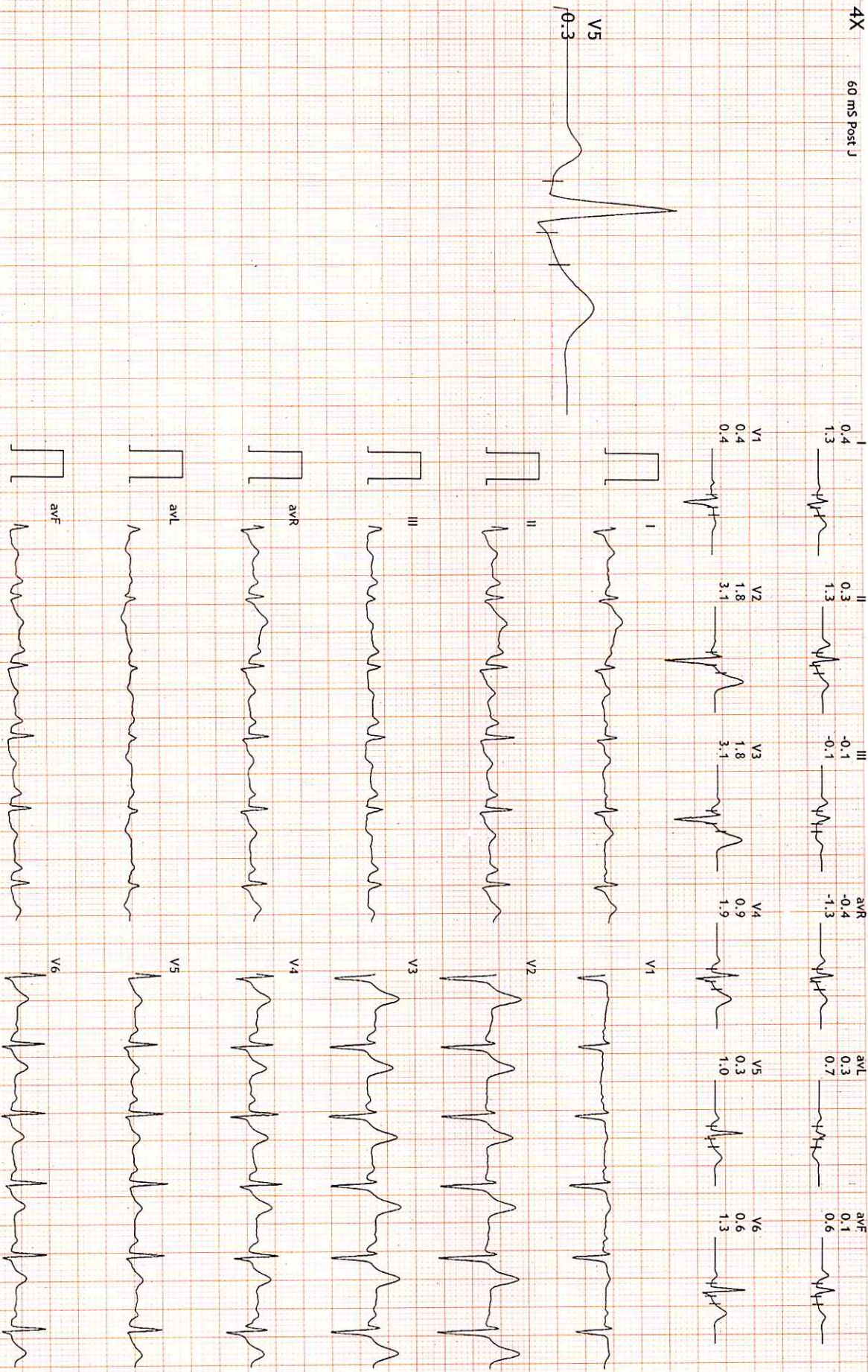
MPHR: 60% of 186
Speed: 0.0 mph
Grade: 0.0%

Raw ECG
BRUCE
(0.05-100)Hz

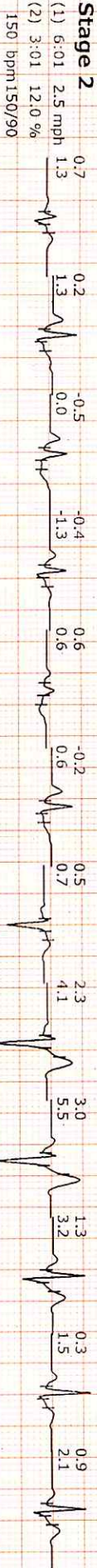
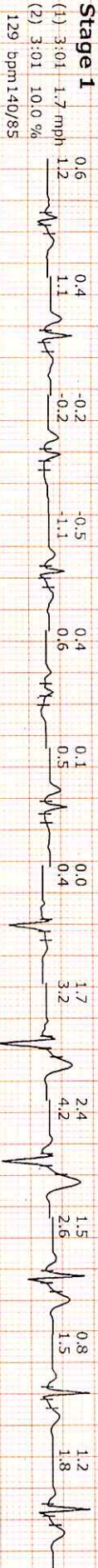
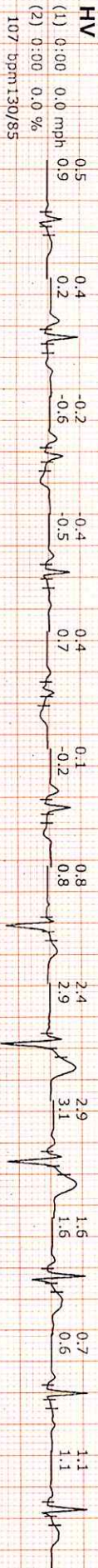
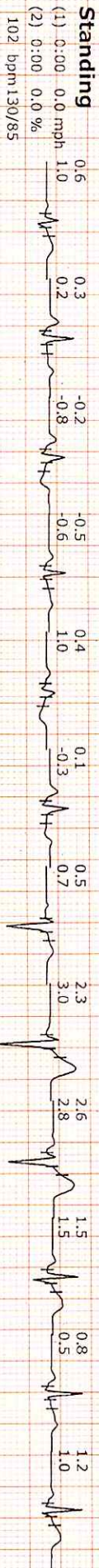
Ex Time 07:07
BLC :On
Notch :On

Recovery(4:00)
10.0 mm/mV
25 mm/Sec.

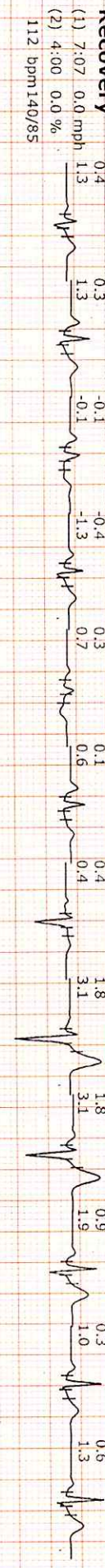
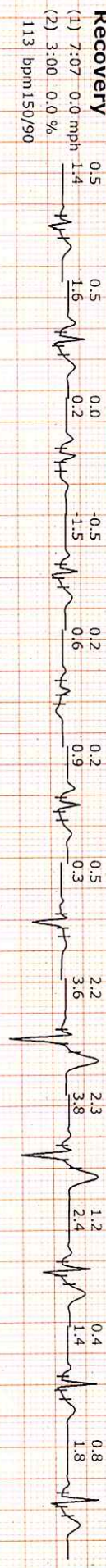
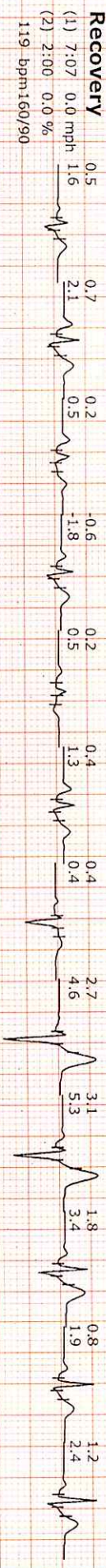
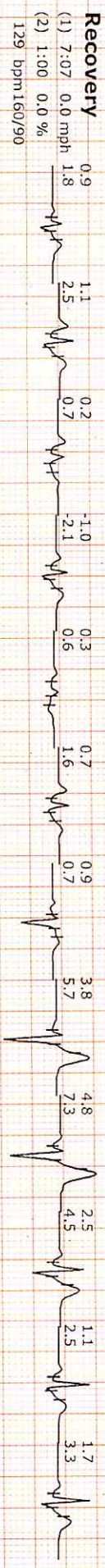
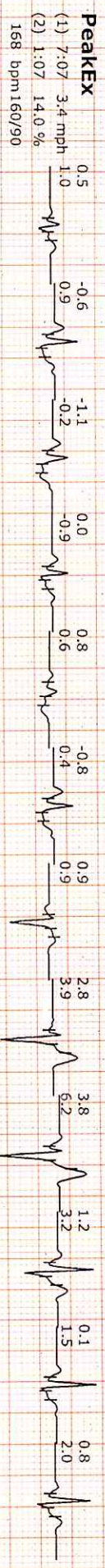
4X 60 ms Post J



I II III aVR aVL aVF V1 V2 V3 V4 V5 V6



I II III aVR aVL aVF V1 V2 V3 V4 V5 V6





P3 HEALTH SOLUTIONS LLP
(ASSOCIATES OF MAXCARE DIAGNOSTICS)



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NAME:	MRS. MANISH TANK	AGE	34 YRS/M
REF.BY	BANK OF BARODA	DATE	25/11/2023

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 in lung parenchyma.

Dr. Mukesh Sharma
M.B.B.S; M.D. (Radiodiagnosis)
RMC No. 43418/17437

R

12233995 MANISH TANK 34 BOB M
25.NOV.2023
MAXCARE DIAGNOSTIC (ASSOCIATES OF P3 HEALTH SOLUTIONS LLP)

