भारत सरकार

Government of India

आधार

बिमान साहू

Biman Sahoo

जन्म तिथि/DOB: 10/12/1983

पुरुष/ MALE

Issue Date: 04/05/2017

7889 6568 9642

VID : 9112 4322 2525 8282

मेरा आधार, मेरी पहचान

भारतीय विशिष्ट पहचान प्राधिकरण

Unique Identification Authority of India

AADHAAR

पता:
द्वारा: गौरांग प्रसाद साहू, फ्लैट न - एफ़ - 8/84 ,
एडब्ल्यूएचओ कॉलोनी, सुगन विहार, सेक्टर 1, विद्याधर
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राजस्थान - 302039

Address:
C/O: Gouranga Prasad Sahoo, Flat No - F -
8/84 , AWHO Colony, Sugan Vihar, Sector 1,
Vidyadhar nagar, Jaipur,
Rajasthan - 302039

Download Date: 11/08/2023

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GPS Map Camera
Jaipur, Rajasthan, India
G-22 Vidhadher Enclave 14, near Cine Star, Sector 2, Central Spine,
Vidyadhar Nagar, Jaipur, Rajasthan 302039, India
Lat 26.964526°
Long 75.782503°
24/12/23 02:03 PM GMT +05:30
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GPS Map Camera
Jaipur, Rajasthan, India
Vidhyadhar Enclave II, b 14, Sector 2 Rd, Sector 2, Central Spine,
Vidyadhar Nagar, Jaipur, Rajasthan 302039, India
Lat 26.964464°
Long 75.782608°
24/12/23 02:05 PM GMT +05:30
Google


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Dr. PIYUSH
MBBS, MD (Radio
MMC No.-037041

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General Physical Examination

Date of Examination: 24/12/1993

Name: BIMAN SAHOO Age: 46YRS DOB: 10/12/1983 Sex: Male

Referred By: BANK OF BARODA

Photo ID: AADHAR CARD ID #: 9649

Ht: 166 (cm) Wt: 84 (Kg)

Chest (Expiration): 98 (cm) Abdomen Circumference: 86 (cm)

Blood Pressure: 130/80 mm Hg PR: 89/ min RR: 17/ min Temp: Afebrile

BMI 29.9

Eye Examination: RE- EG, NG, NCB
LE- EG, NG, NCB

Other: No

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

Signature Of Examine : Name of Examinee: BIMAN SAHOO

Signature Medical Examiner : Name Medical Examiner DR. PIYUSH GOYAL

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

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NAME :- Mr. BIMAN SAHOO	Patient ID :42234236	Date :- 24/12/2023	10:27:03
Age :- 40 Yrs 14 Days	Ref. By Doctor:-BANK OF BARODA		
Sex :- Male	Lab/Hosp :-		
	Company :- Mr.MEDIWHEEL		

Final Authentication : 24/12/2023 16:38:39

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	7.30	/cumm	4.00 - 10.00
DIFFERENTIAL LEUCOCYTE COUNT			
NEUTROPHIL	71.0	%	40.0 - 80.0
LYMPHOCYTE	23.0	%	20.0 - 40.0
EOSINOPHIL	2.0	%	1.0 - 6.0
MONOCYTE	4.0	%	2.0 - 10.0
BASOPHIL	0.0	%	0.0 - 2.0
TOTAL RED BLOOD CELL COUNT (RBC)	5.23	x10^6/uL	4.50 - 5.50
HEMATOCRIT (HCT)	46.90	%	40.00 - 50.00
MEAN CORP VOLUME (MCV)	90.0	fL	83.0 - 101.0
MEAN CORP HB (MCH)	28.9	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	32.2	g/dL	31.5 - 34.5
PLATELET COUNT	188	x10^3/uL	150 - 410
RDW-CV	13.9	%	11.6 - 14.0

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HAEMATOLOGY

Erythrocyte Sedimentation Rate (ESR)	13	mm in 1st hr	00 - 15
Method:- Westergreen			

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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		Company :-	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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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
FASTING BLOOD SUGAR (Plasma) Method - GOD POD	84.9	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	89.2	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) 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 4-8 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 azidothymidine, 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

BLOOD GROUP ABO Method - Haemagglutination reaction	"O" POSITIVE

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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIPID PROFILE			
TOTAL CHOLESTEROL Method:- CHOD-PAP methodology	193.00	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	**154.00** H	mg/dl	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 Method:- Direct clearance Method	51.20	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	116.13	mg/dl	Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190
VLDL CHOLESTEROL Method:- Calculated	30.80	mg/dl	0.00 - 80.00
T.CHOLESTEROL/HDL CHOLESTEROL RATIO Method:- Calculated	3.77		0.00 - 4.90
LDL / HDL CHOLESTEROL RATIO Method:- Calculated	2.27		0.00 - 3.50
TOTAL LIPID Method:- CALCULATED	609.55	mg/dl	400.00 - 1000.00

1 Measurements in the same patient can show physiological& analytical variations. Three serial samples 1 week apart are recommended for Total Cholesterol, Triglycerides, HDL& LDL Cholesterol.

2 As per NCEP guidelines, all adults above the age of 20 years should be screened for lipid status.Selective screening of children above the age of 2 years with a family history of premature cardiovascular disease or those with at least one parent with high total cholesterol is

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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 fromperipheral tissues

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BIOCHEMISTRY

LIVER PROFILE WITH GGT

SERUM BILIRUBIN (TOTAL) Method:- DMSO/Diazo	0.62	mg/dL	Infants : 0.2-8.0 mg/dL Adult - Up to - 1.2 mg/dL
SERUM BILIRUBIN (DIRECT) Method:- DMSO/Diazo	0.20	mg/dL	Up to 0.40 mg/dL
SERUM BILIRUBIN (INDIRECT) Method:- Calculated	0.42	mg/dl	0.30-0.70
SGOT Method:- IFCC	24.5	U/L	0.0 - 40.0
SGPT Method:- IFCC	29.1	U/L	0.0 - 40.0
SERUM ALKALINE PHOSPHATASE Method:- DGKC - SCE	88.60	U/L	53.00 - 141.00
SERUM GAMMA GT Method:- Szasz methodology Instrument Name:Randox Rx Imola	32.50	U/L	10.00 - 45.00
SERUM TOTAL PROTEIN Method:- Direct Biuret Reagent	7.89	g/dl	6.00 - 8.40
SERUM ALBUMIN Method:- Bromocresol Green	4.46	g/dl	3.50 - 5.50
SERUM GLOBULIN Method:- CALCULATION	3.43	gm/dl	2.20 - 3.50
A/G RATIO	1.30		1.30 - 2.50

Interpretation:- Elevations in GGT levels occurs 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.

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 Method - Urease/GLDH	36.50	mg/dl	10.00 - 50.00

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

SERUM CREATININE Method - Jaffe's Method	1.26	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	5.83	mg/dl	2.40 - 7.00

InstrumentName:HORIBA YUMIZEN CA60 Daytona plus **Interpretation**: **Elevated Urate**:High purine diet,Alcohol• Renal insufficiency,Drugs , Polycythaemia vera, Malignancies,Hypothyroidism,Rare enzyme defects ,Downs syndrome,Metabolic syndrome, Pregnancy,Gout.

SODIUM Method - ISE	138.5	mmol/L	135.0 - 150.0
POTASSIUM Method - ISE	4.42	mmol/L	3.50 - 5.50
CHLORIDE Method - ISE	98.9	mmol/L	94.0 - 110.0
SERUM CALCIUM Method - Arsenazo III Method	10.00	mg/dl.	8.80 - 10.20

InstrumentName:MISPA PLUS **Interpretation**: Serum calcium levels are believed to be controlled by parathyroid hormone and vitamin D. Increases in serum PTH or vitamin D are usually associated with hypercalcemia .Hypocalcemia may be observed in hypoparathyroidism, nephrosis and pancreatitis.

SERUM TOTAL PROTEIN Method - Direct Biuret Reagent	7.89	g/dl	6.00 - 8.40
SERUM ALBUMIN Method - Bromocresol Green	4.46	g/dl	3.50 - 5.50
SERUM GLOBULIN Method - CALCULATION	3.43	gm/dl	2.20 - 3.50
A/G RATIO	1.30		1.30 - 2.50

Interpretation : Measurements obtained by this method are used in the diagnosis and treatment of a variety of dis [illegible] 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-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 Method - ECLIA	1.12	ng/mL	0.70 - 2.04

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. Transient increase in TSH levels or abnormal TSH levels can be seen in some non thyroidal conditions,simoultaneous 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.High TSH,Low FT4 and Thyroid microsomal antibody increased seen in patients with Hashimotos thyroiditis 5.High TSH,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 levels8.Normal T4 levels accompanied by ↑ T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis9.Normal or ↑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 Hypoth

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	9.11	µg/dl	5.10 - 14.10

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. Transient increase in TSH levels or abnormal TSH levels can be seen in some non thyroidal conditions,simoultaneous 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.High TSH,Low FT4 and Thyroid microsomal antibody increased seen in patients with Hashimotos thyroiditis 5.High TSH,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 levels8.Normal T4 levels accompanied by ↑ T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis9.Normal or ↑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 Hypoth

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.

TSH Method - ECLIA	2.977	µIU/mL	0.350 - 5.500

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

Technologist
MGR

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

P3 HEALTH SOLUTIONS LLP
(ASSOCIATES OF MAXCARE DIAGNOSTICS)

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

NAME :- Mr. BIMAN SAHOO		Patient ID :-12234236	Date :- 24/12/2023 10:27:03
Age :-	40 Yrs 14 Days	Ref. By Doctor:-BANK OF BARODA	
Sex :-	Male	Lab/Hosp :-	
		Company :-	Mr.MEDIWHEEL

Final Authentication : 24/12/2023 16:38:39

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.High TSH,Low FT4 and Thyroid microsomal antibody increased seen in patients with Hashimotos thyroiditis
5.High TSH,Low FT4 and Thyroid microsomal antibody normal seen in patients with Iodine deficiency,Congenital T4 synthesis deficiency
6.Low TSH,Low FT4 and TRH simulation 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 propanolol
14.Although ↑ TSH levels are nearly always indicative of Primary Hypothiroidism ,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 T-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 ***

MGR
Technologist

DR.TANU RUNGTA
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RMC No. 17226

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+91 141 4824885 maxcarediagnostics1@gmail.com

NAME :- Mr. BIMAN SAHOO	Patient ID :-12234236	Date :- 24/12/2023	10:27:03
Age :- 40 Yrs 14 Days	Ref. By Doctor:-BANK OF BARODA		
Sex :- Male	Lab/Hosp :-		
	Company :- Mr.MEDIWHEEL		

Final Authentication : 24/12/2023 16:36:39

CLINICAL PATHOLOGY

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

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


LIVER


LIVER


GALLBLADDER


KIDNEY


PROSTATE


RT KIDNEY


SPLEEN


LT KIDNEY

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MR. BIMAN SAHOO	40 Y/M
Registration Date: 24/12/2023	Ref. by: BANK OF BARODA

ULTRASOUND OF WHOLE ABDOMEN

Liver is of normal size (11.5 cm). Echo-texture is normal. A well-defined, anechoic cystic lesion of size 13 x 14 mm is seen in segment IV – suggestive of simple hepatic cyst. Intra hepatic biliary channels are not dilated. Portal vein diameter is normal.

Gall bladder shows a well-defined calculus of size 17-18 mm with posterior acoustic shadowing in body region; however, no evidence of pericholecystic free fluid is noted. No 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 (10.0 cm). Echotexture is normal. No focal lesion is seen.

Kidneys are normally sited and are of normal size and shape. Cortico-medullary echoes are normal. Collecting system does not show any calculus or dilatation.
Right kidney is measuring approx. 9.6 x 4.1 cm.
Left kidney is measuring approx. 10.1 x 4.5 cm.

Urinary bladder is partially 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: Cholelithiasis as described above.

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

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

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Central Spine, Vidhyadhar Nagar, Jaipur - 302023

+91 141 4824885 maxcarediagnostics1@gmail.com

NAME:	MR. BIMAN SAHOO	AGE	40 YRS/M
REF.BY	BANK OF BARODA	DATE	24/12/2023

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.

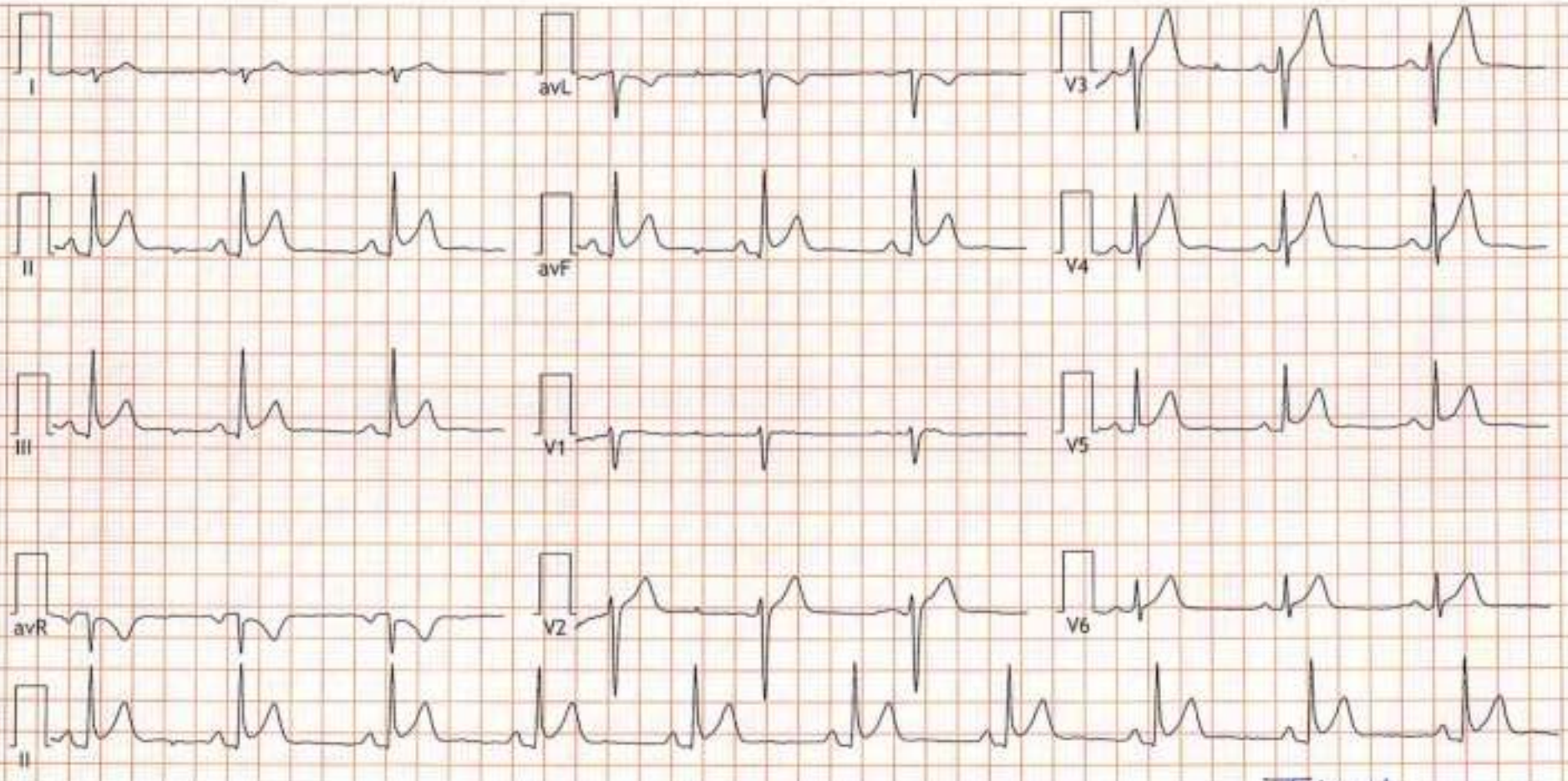
DR.SHALINI GOEL

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

RMC No.: 21954

Tems (P) Ltd
#P3 HEALTH SOLUTIONS LLP B-14, Vidhyadhar nahar , Jaipur
12234237/Mr Biman Sahoo 40Yrs/Male Kgs/ Cms BP: ___/___ mmHg HR: 59 bpm
Ref.: BANK OF BARODA Test Date: 24-Dec-2023(2:18:24 P) Notch: 50Hz 0.05Hz - 35Hz 10mm/mV 25mm/Sec

PR Interval: 162 ms
QRS Duration: 106 ms
QT/QTc: 356/353ms
P-QRS-T Axis: 74 - 91 - 74 (Deg)



TUNL

FINDINGS: Abnormal ECG with Indication of Sinus Bradycardia
Vent Rate : 59 bpm; **PR Interval :** 162 ms; **QRS Duration:** 106 ms; **QT/QTc Int :** 356/353 ms
P-QRS-T axis: 74• 91• 74• (Deg)
Comments :

Dr. NARESH MOHINKA

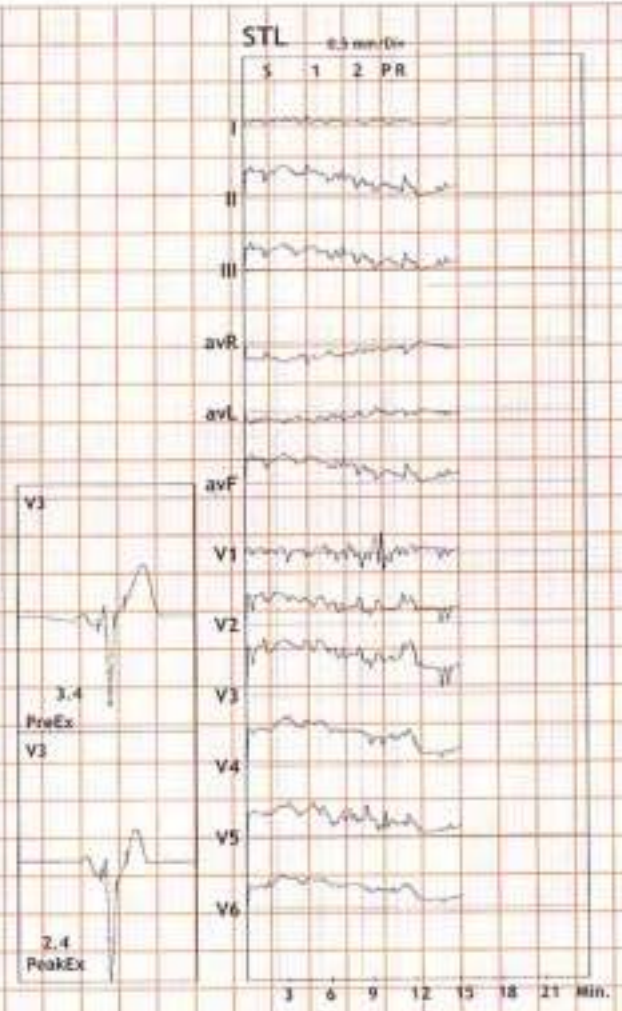
P3 HEALTH SOLUTIONS LLP
B-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur

Summary

12234237/Mr BIMAN SAHOO 40 Yrs/Male 0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM
Ref.By : BANK OF BARODA
Medication : Nil
Objective :

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	62	125/80	77	-	
Standing					1.0	60	125/80	75	-	
HV					1.0	89	125/80	111	-	
ExStart					1.0	76	125/80	95	-	
Stage 1	3:01	3:02	1.7	10.0	4.7	110	135/80	148	-	
Stage 2	3:01	6:02	2.5	12.0	7.1	123	145/85	178	-	
PeakEx	2:29	8:30	3.4	14.0	9.7	156	155/85	241	-	
Recovery	1:00		0.0	0.0	1.2	106	155/85	164	-	
Recovery	2:00		0.0	0.0	1.0	89	165/90	146	-	
Recovery	3:00		0.0	0.0	1.0	83	155/85	128	-	
Recovery	4:00		0.0	0.0	1.0	81	145/85	117	-	



Findings :

Exercise Time :08:29
Max HR Attained :156 bpm 87% of Max Predictable HR 180
Max BP : 165/90(mmHg)
Max WorkLoad attained :9.7(Good Effort Tolerance)

TMT is Negative for RMI.

Advice/Comments:

DR. NARESH MOHINKA

P3 HEALTH SOLUTIONS LLP

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

12234237/Mr BIMAN SAHOO
40 Yrs/Male
0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

HR: 62 bpm
METS: 1.0
BP: 125/80

MPHR: 34% of 180
Speed: 0.0 mph
Grade: 0.0%

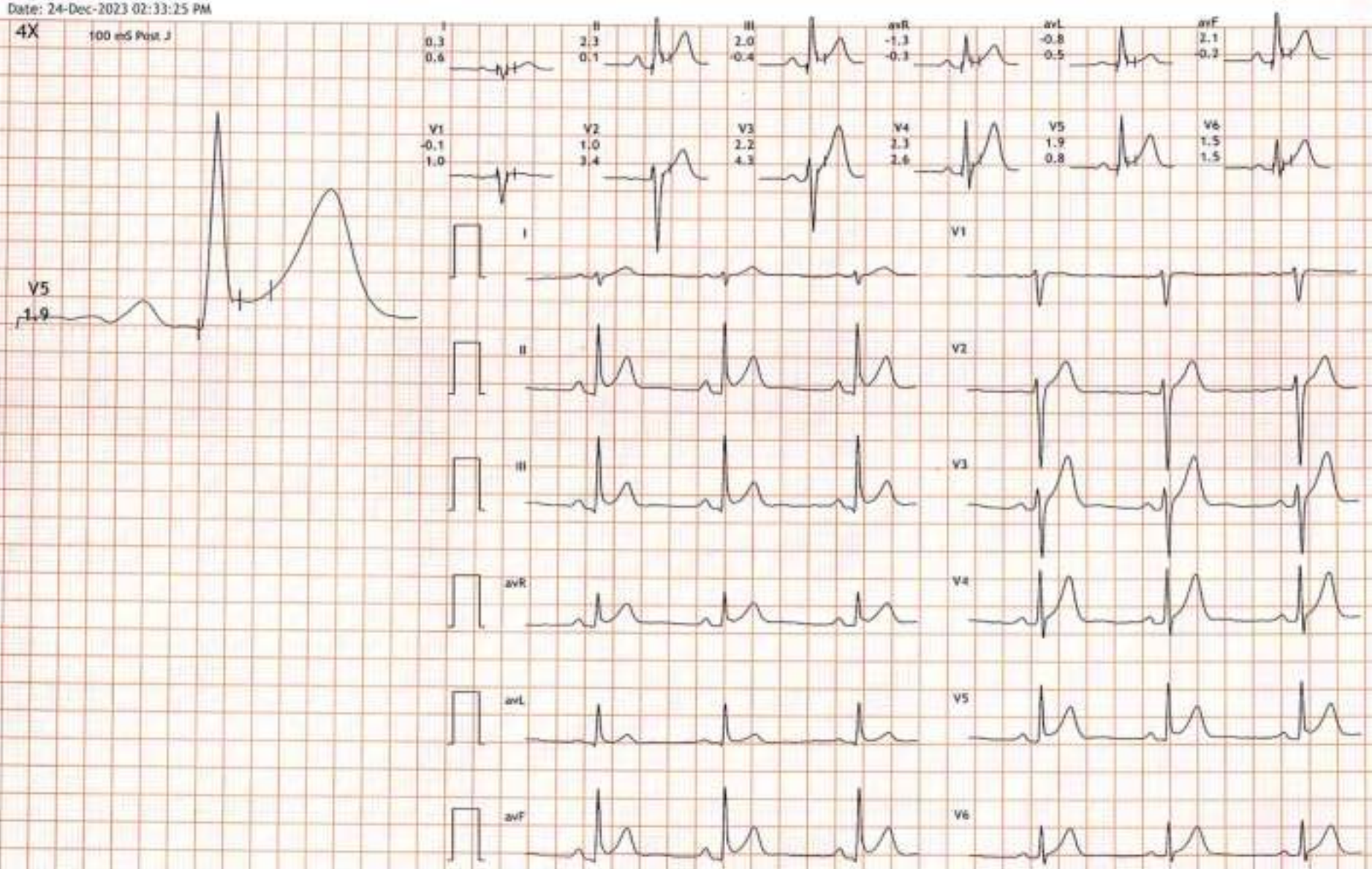
Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 00:31
BLC :On
Notch :On

Supine
10.0 mm/mV
25 mm/Sec.

12 Lead + Median

RMS



DR. NARESH MOHINKA

P3 HEALTH SOLUTIONS LLP
B-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur

12 Lead + Median

12234237/Mr BIMAN SAHOO
40 Yrs/Male
0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

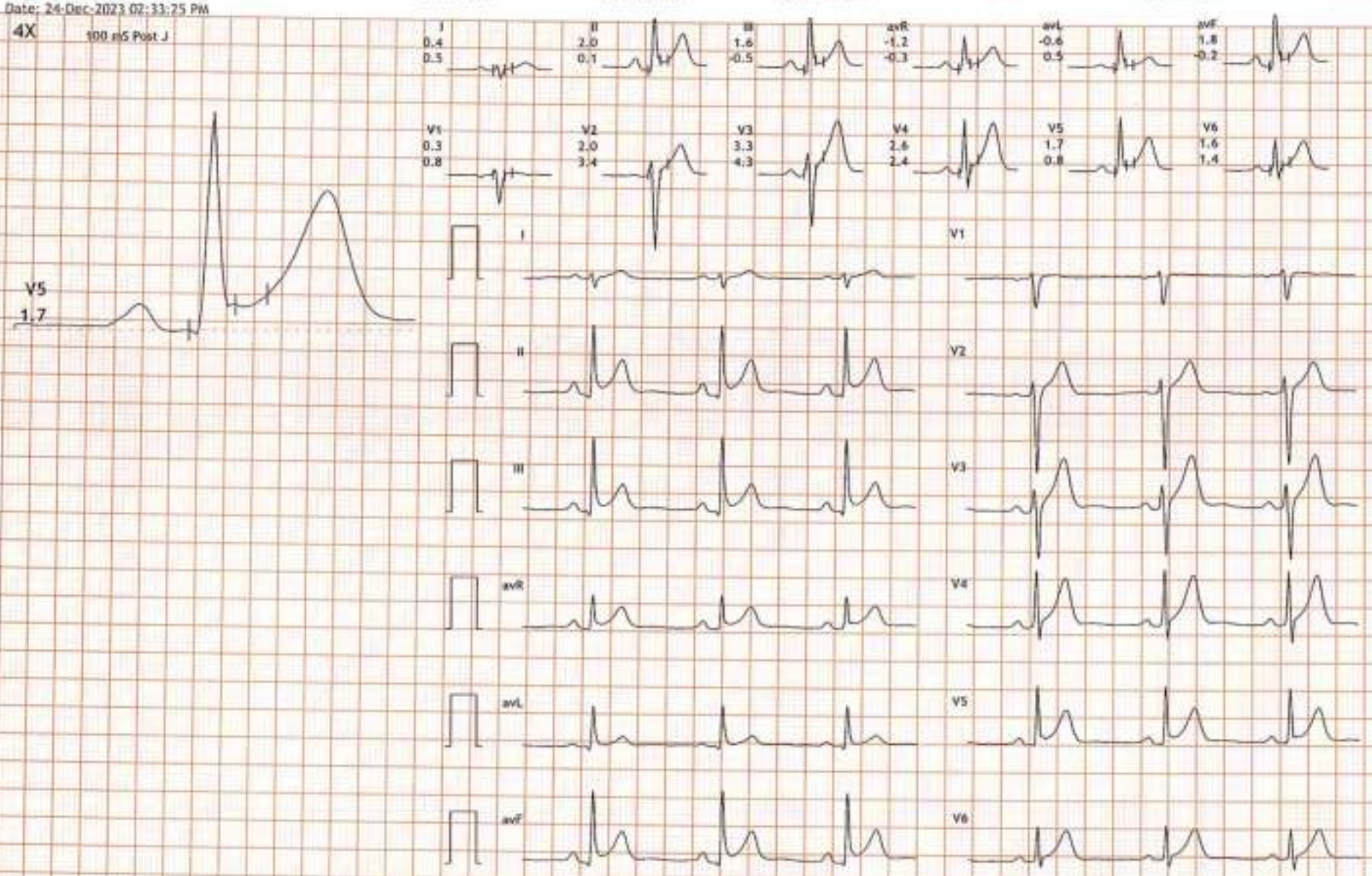
HR: 60 bpm
METS: 1.0
BP: 125/80

MPHR:33% of 180
Speed: 0.0 mph
Grade: 0.0%

Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 00:58
BLC :On
Notch :On

Standing
10.0 mm/mV
25 mm/Sec.



DR. NARESH MOHINKA

P3 HEALTH SOLUTIONS LLP
B-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur

12234237/Mr BIMAN SAHOO
40 Yrs/Male
0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

HR: 83 bpm
METS: 1.0
BP: 125/80

MPHR:46% of 180
Speed: 0.0 mph
Grade: 0.0%

Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 01:51
BLC :On
Notch :On

HV
10.0 mm/mV
25 mm/Sec.

12 Lead + Median

RMS

4X 100 mS Post J

I	II	III	avR	avL	avF
0.4	1.7	1.4	-1.1	-0.5	1.6
0.5	-0.8	-1.3	0.1	0.9	1.1

V1	V2	V3	V4	V5	V6
0.3	1.7	3.2	2.6	1.7	1.6
1.6	3.9	5.3	3.3	1.0	1.9

V5
1.7

I
II
III
avR
avL
avF

V1
V2
V3
V4
V5
V6

DR. NARESH MOHINKA

P3 HEALTH SOLUTIONS LLP

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

12 Lead + Median

12234237/Mr BIMAN SAHOO	HR: 78 bpm	MPHR:43% of 180	Raw ECG	Ex Time 02:08	ExStart
40 Yrs/Male	METS: 1.0	Speed: 0.0 mph	BRUCE	BLC :On	10.0 mm/mV
0 Kg/0 Cms	BP: 125/80	Grade: 0.0%	(0.05-100)Hz	Notch :On	25 mm/Sec.

Date: 24-Dec-2023 02:33:25 PM

DR. NARESH MOHINKA

P3 HEALTH SOLUTIONS LLP
B-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur
12234237/Mr BIMAN SAHOO
40 Yrs/Male
0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

12 Lead + Median

HR: 110 bpm
METS: 4.7
BP: 135/80

MPHR:61% of 180
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

RMS

4X 100 mS Post J

V5
1.9

I	II	III	avR	avL	avF
0.4	2.0	1.6	-1.2	-0.6	1.8
0.6	1.0	0.4	-0.8	0.1	0.7

V1	V2	V3	V4	V5	V6
0.4	1.7	3.9	3.2	1.9	2.0
0.3	3.0	5.2	3.8	1.8	2.2

I
II
III
avR
avL
avF
V1
V2
V3
V4
V5
V6

DR. NARESH MOHINKA

P3 HEALTH SOLUTIONS LLP

8-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur

12234237/Mr BIMAN SAHOO
40 Yrs/Male
0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

HR: 124 bpm
METS: 7.1
BP: 145/85

MPHR:68% of 180
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.

12 Lead + Median

RMS

4X 100 mS Post J

V5
1.4

I	II	III	avR	avL	avF
0.4 0.6	0.7 1.0	0.3 0.4	-0.6 -0.8	0.0 0.1	0.5 0.7

V1	V2	V3	V4	V5	V6
0.0 -0.1	1.9 3.2	3.4 5.6	2.5 4.3	1.4 2.0	1.6 2.6

I
II
III
avR
avL
avF
V1
V2
V3
V4
V5
V6

DR. NARESH MOHINKA

P3 HEALTH SOLUTIONS LLP

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

12234237/Mr BIMAN SAHOO
40 Yrs/Male
0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

HR: 157 bpm
METS: 9.7
BP: 155/85

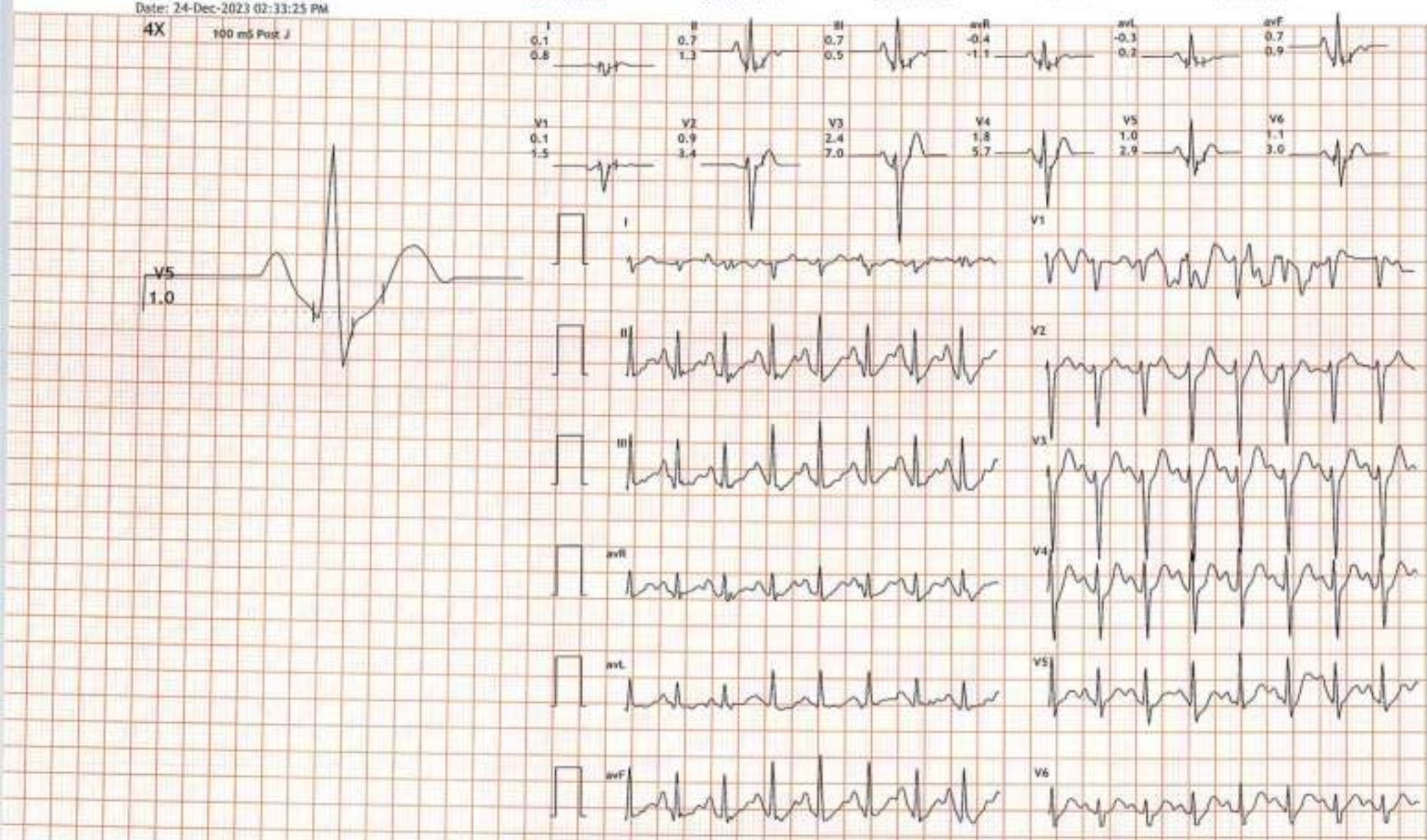
MPHR:87% of 180
Speed: 3.4 mph
Grade: 14.0%

Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 08:27
BLC :On
Notch :On

BRUCE:PeakEx(2:27)
10.0 mm/mV
25 mm/Sec.

12 Lead + Median



DR. NARESH MOHINKA

P3 HEALTH SOLUTIONS LLP

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

12234237/Mr BIMAN SAHOO
40 Yrs/Male
0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

HR: 107 bpm
METS: 1.3
BP: 155/85

MPHR:59% of 180
Speed: 0.0 mph
Grade: 0.0%

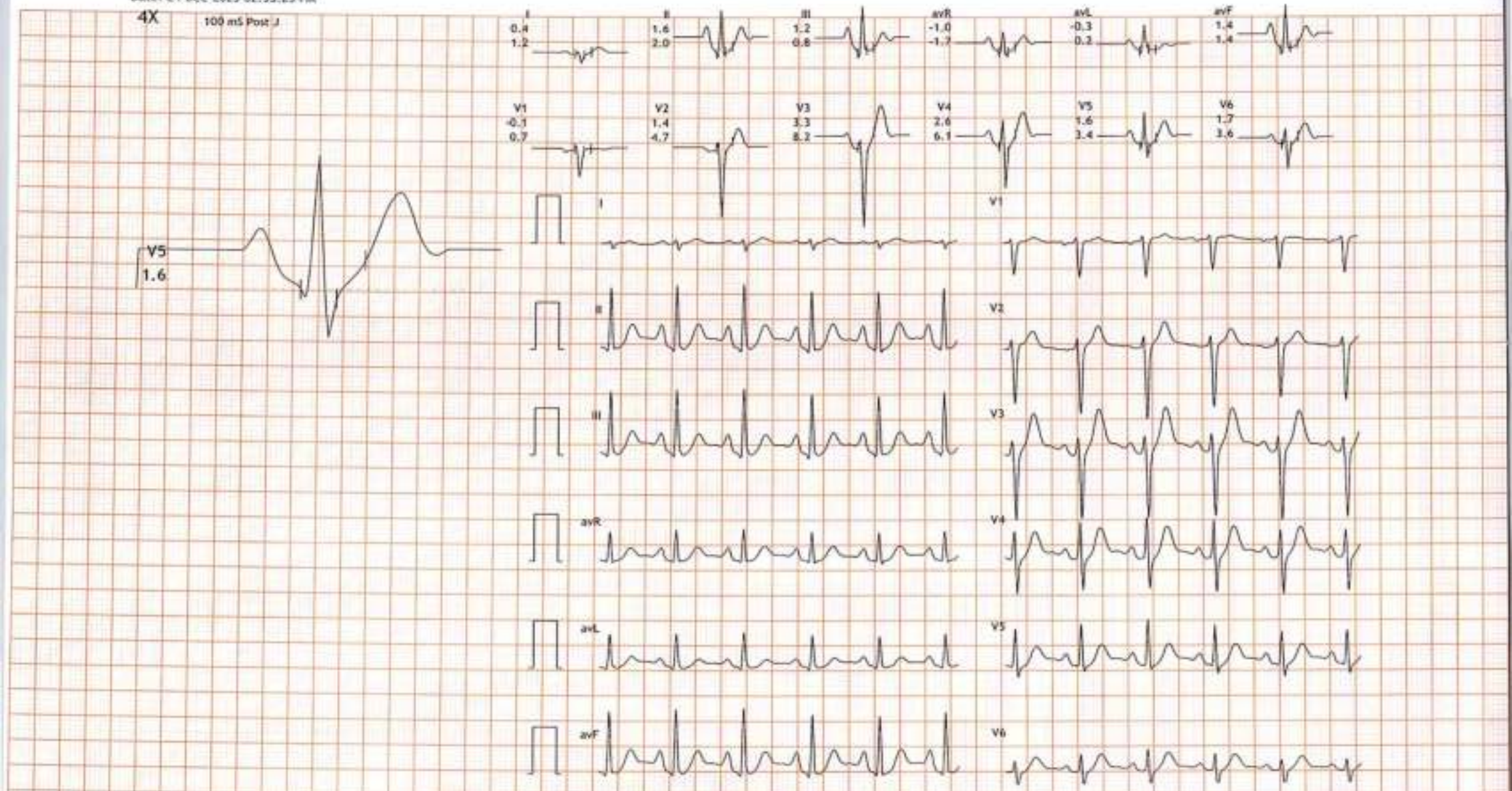
Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 08:29
BLC :On
Notch :On

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

12 Lead + Median

RMS



DR. NARESH MOHINKA

P3 HEALTH SOLUTIONS LLP

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

12 Lead + Median

12234237/Mr BIMAN SAHOO
40 Yrs/Male
0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

HR: 88 bpm
METS: 1.0
BP: 165/90

MPHR:48% of 180
Speed: 0.0 mph
Grade: 0.0%

Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 08:29
BLC :On
Notch :On

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

4X 100 mS Post J

V5
0.3

I	II	III	avR	avL	avF
0.2 0.5	0.2 0.4	0.0 -0.1	-0.2 -0.4	0.1 0.3	0.1 0.1

V1	V2	V3	V4	V5	V6
0.3 1.0	1.0 3.0	2.0 5.1	0.9 3.2	0.3 1.4	0.5 1.7

I
II
III
avR
avL
avF
V1
V2
V3
V4
V5
V6

DR. NARESH MOHINKA

P3 HEALTH SOLUTIONS LLP
B-14, Vidhyadhar Enclave-2, Vidhyadhar Nagar, Jaipur
12234237/Mr BIMAN SAHOO
40 Yrs/Male
0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

12 Lead + Median

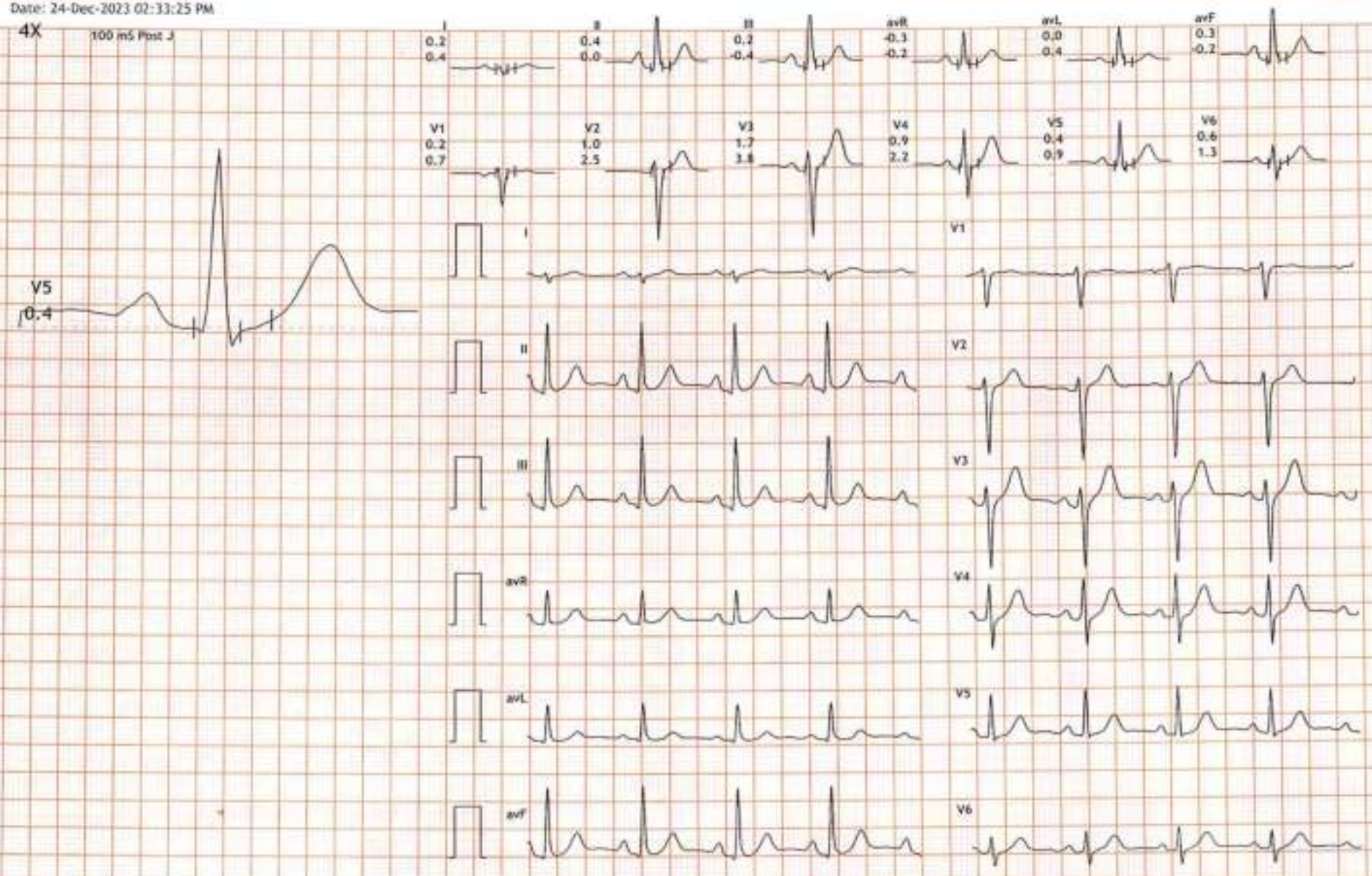
HR: 83 bpm
METS: 1.0
BP: 155/85

MPHR:46% of 180
Speed: 0.0 mph
Grade: 0.0%

Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 08:29
BLC :On
Notch :On

Recovery(3:00)
10.0 mm/mV
25 mm/Sec.



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12234237/Mr BIMAN SAHOO
40 Yrs/Male
0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

12 Lead + Median

HR: 81 bpm
METS: 1.0
BP: 145/85

MPHR:45% of 180
Speed: 0.0 mph
Grade: 0.0%

Raw ECG
BRUCE
(0.05-100)Hz

Ex Time 08:29
BLC :On
Notch :On

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

RMS

4X 100 ms Post J

V5
0.5

I	II	III	avR	avL	avF
0.3	0.7	0.4	-0.4	0.0	0.5
0.5	-0.3	-0.8	-0.1	0.6	-0.6

V1	V2	V3	V4	V5	V6
0.2	1.1	1.8	1.0	0.5	0.6
0.7	2.8	3.5	1.8	0.5	1.1

I
II
III
avR
avL
avF
V1
V2
V3
V4
V5
V6

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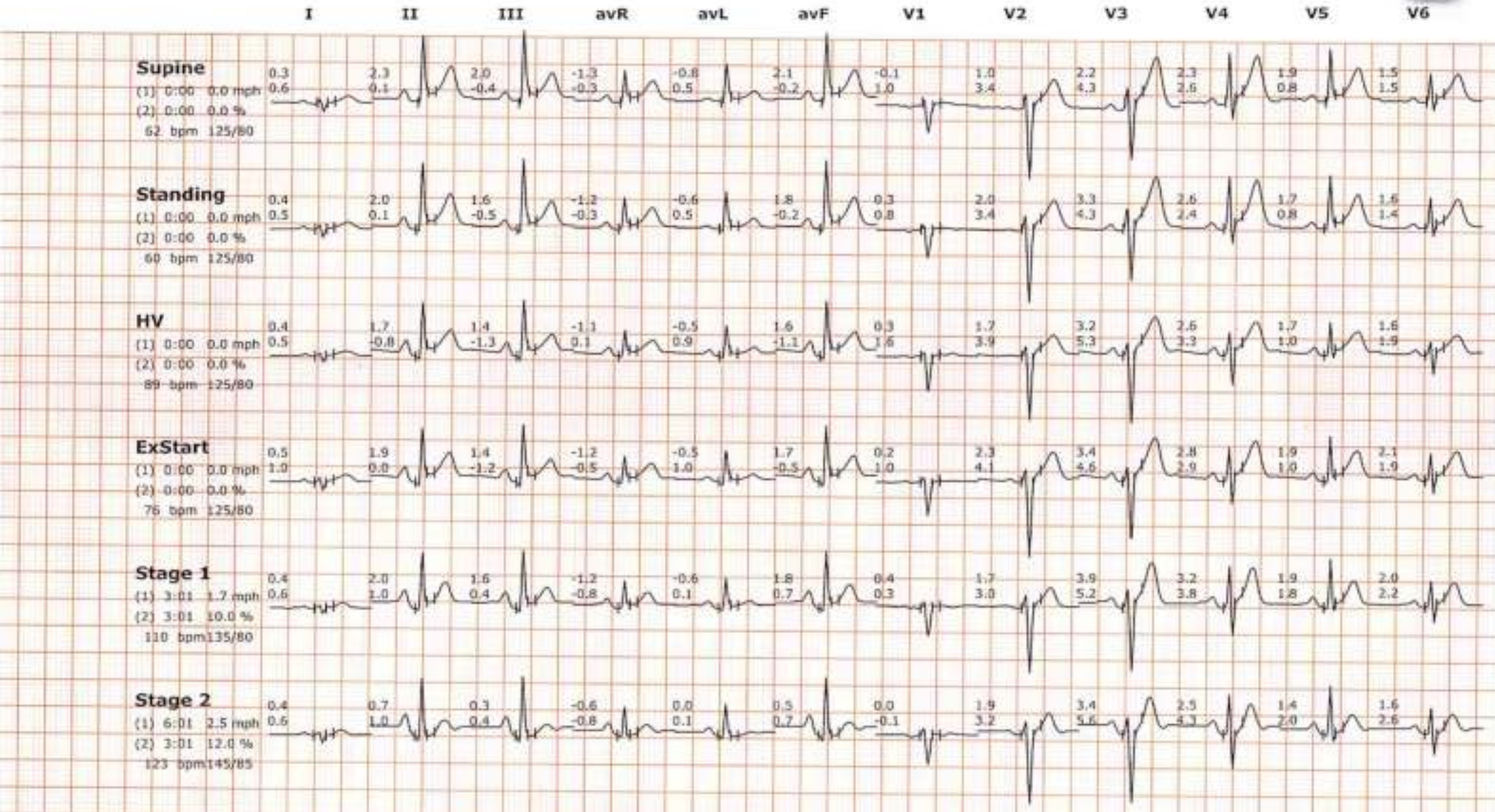
12234237/Mr BIMAN SAHOO 40 Yrs/Male 0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

Average

RMS



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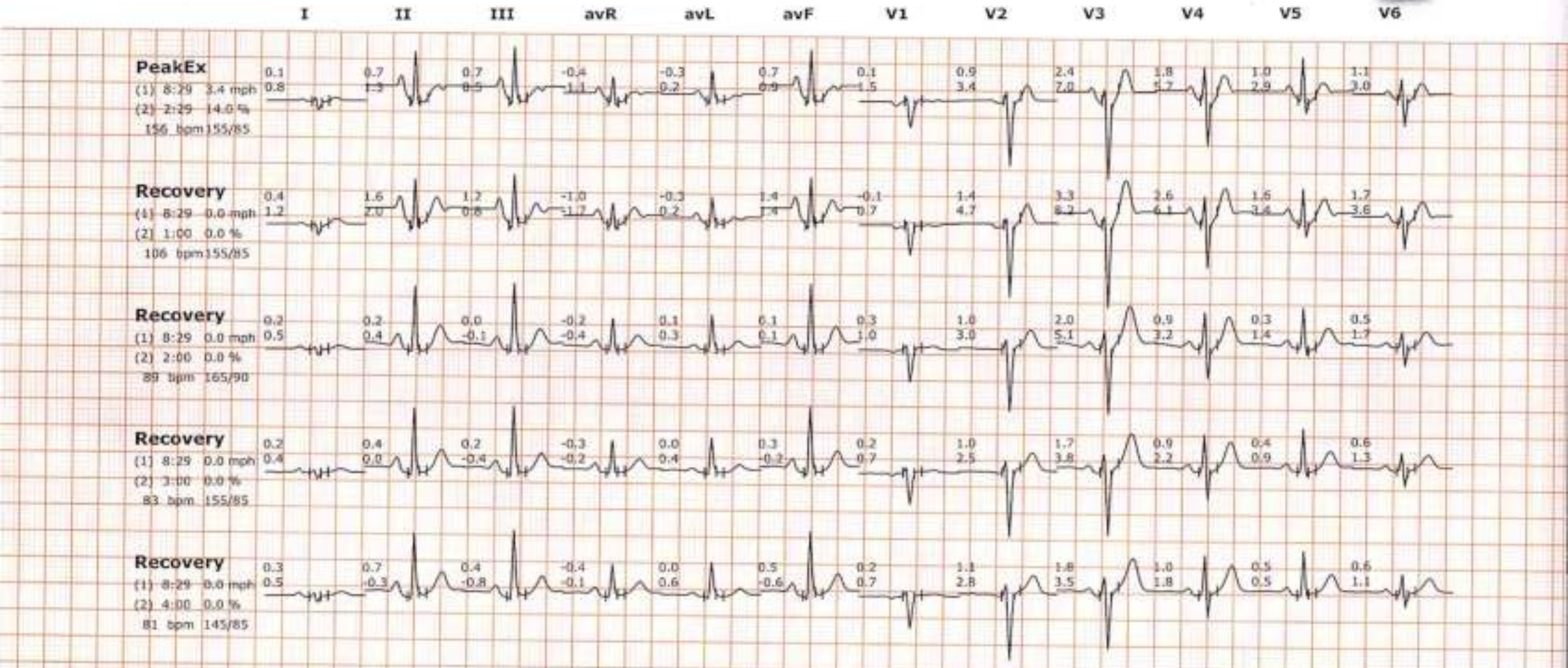
12234237/Mr BIMAN SAHOO 40 Yrs/Male 0 Kg/0 Cms
Date: 24-Dec-2023 02:33:25 PM

Average

RMS



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