



Lab No.	012411200230	Age/Gender	54 YRS/MALE	Coll. ON	20/Nov/2024 09:16AM
NAME	Mr. JAGDISH			Reg. ON	20/Nov/2024
Ref. Dr.	MEDIWEEL	BarcodeNo	01200230	Approved ON	20/Nov/2024 02:49PM
Rpt. Centre	undefined			Printed ON	20/Nov/2024 06:52PM

Test Name	Value	Unit	Biological Reference Interval
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Complete Haemogram, EDTA whole blood

Haemoglobin (Hb) <i>Method : Colorimetry</i>	14.70	gm/dl	13.0 - 17.0
RBC count <i>Method : Electrical impedance</i>	4.45	Millions/cmm	4.5 - 5.5
PCV / Haematocrit <i>Method : Calculated</i>	41.80	%	40.0 - 50.0
MCV <i>Method : Calculated</i>	93.80	fl	83.0 - 101.0
MCH <i>Method : Calculated</i>	33.00	picogram	27.0 - 32.0
MCHC <i>Method : Calculated</i>	35.10	%	31.5 - 34.5
RDW - CV <i>Method : Calculated</i>	13.80	%	11.6 - 14.0
Mentzer Index <i>Method : Calculated</i>	21.08		>= 13.0

The Mentzer index (MCV/ RBC count) is a useful tool for initial screening of patients with a microcytic hypochromic blood picture to rule out a thalassemia trait. If the index is less than 13, thalassemia is said to be more likely. If the result is greater than 13, then iron-deficiency anemia is said to be more likely. All patients with a low normal to low hemoglobin and a Mentzer index below 13 should be screened for thalassemia trait by HPLC.

TLC (Total Leucocyte Count) <i>Method : Flowcytometry</i>	7,050	/cmm	4000 - 10000
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DLC (Flowcytometry)

Neutrophils	45.80	%	35.0 - 75.0
Lymphocytes	43.40	%	25.0 - 45.0
Eosinophils	4.30	%	1.0 - 5.0
Monocytes	6.00	%	1.0 - 6.0
Basophils	0.50	%	0 - 1

Absolute Leucocyte Count (Calculated)

Absolute Neutrophil Count	3,228.90	/cmm	2000 - 7000
Absolute Lymphocyte Count	3,059.70	/cmm	1000 - 3000
Absolute Eosinophil count	303.15	/cmm	20 - 500
Absolute Monocyte count	423.00	/cmm	200 - 1000
Absolute Basophil count	35.25	/cmm	0 - 100

Platelet count <i>Method : Electrical impedance</i>	1.67	Lakh/cmm	1.5 - 4.1
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ESR (Erythrocyte Sedimentation Rate) <i>Method : Westergren method</i>	20	mm/1st hr	0 - 22
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Peripheral Smear

RBCs are normocytic and normochromic.
Leucocytic series is numerically and morphologically within normal limits.
Platelets are adequate in number and are normal in morphology.
No atypical cells or haemoparasites are seen.
Impression: Normal peripheral smear.

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Blood Group (ABO + RH)

Blood Group , EDTA blood O
 Method : Slide agglutination (Forward & Reverse grouping)

Rh type , EDTA blood Positive
 Method : Slide agglutination



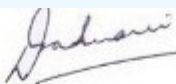
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Test Name	Value	Unit	Biological Reference Interval
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Glucose Fasting, plasma Method : GOD POD	90.10	mg/dL	60 - 100
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Interpretation (In accordance with the American diabetes association guidelines):

- A fasting plasma glucose level below 100 mg/dl is considered normal.
- A fasting plasma glucose level between 100-126 mg/dl is considered as glucose intolerant or pre diabetic. A fasting and post-prandial blood sugar test (after consumption of 75 gm of glucose) is recommended for all such patients.
- A fasting plasma glucose level of above 126 mg/dl is highly suggestive of a diabetic state. A repeat fasting test is strongly recommended for all such patients. A fasting plasma glucose level in excess of 126 mg/dl on both the occasions is confirmatory of a diabetic state.

Glucose PP, plasma Method : GOD POD	125.10	mg/dL	90 - 140
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Interpretation (In accordance with the American diabetes association guidelines):

- A post-prandial plasma glucose level below 140 mg/dl is considered normal.
- A post-prandial plasma glucose level between 140-199 mg/dl is considered as glucose intolerant or pre diabetic. A fasting and post-prandial blood sugar test (after consumption of 75 gm of glucose) is recommended for all such patients.
- A post-prandial plasma glucose level of above 200 mg/dl is highly suggestive of a diabetic state. A repeat post-prandial test is strongly recommended for all such patients. A post-prandial plasma glucose level in excess of 200 mg/dl on both the occasions is confirmatory of a diabetic state.

Blood Urea Nitrogen (BUN), serum Method : Calculated	7.92	mg/dl	7.8 - 20.2
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Serum Creatinine Method : Jaffe kinetic	0.82	mg/dl	0.7 - 1.2
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Serum Uric Acid Method : Uricase-Peroxidase	6.16	mg/dl	3.6 - 8.2
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HbA1c (Glycosylated haemoglobin) , EDTA whole blood <i>Method : HPLC</i>	5.80	%	< 5.7
Estimated average plasma Glucose <i>Method : Calculated</i>	119.76	mg/dL	65 - 136

The test is approved by NGSP for patient sample testing.

Interpretation:

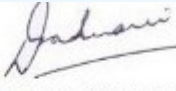
Metabolically normal patients	%	< 5.7
Pre-diabetic	%	5.7 - 6.4
Diabetic	%	> 6.4

Glycosylated hemoglobin or HbA1C is a reliable indicator of mean plasma glucose levels for a period of 8-12 weeks preceding the date on which the test is performed and is a more reliable indicator of overall blood sugar control in known diabetic patients than blood sugar levels. A value of less than 5.7 % is usually seen in metabolically normal patients, however diabetics with very good control can also yield similar values. The HbA1c test, thus can not be used to differentiate between diabetic patients with very good control over the plasma glucose levels from metabolically normal, non-diabetic subjects as both groups may reveal very similar values in the assay.



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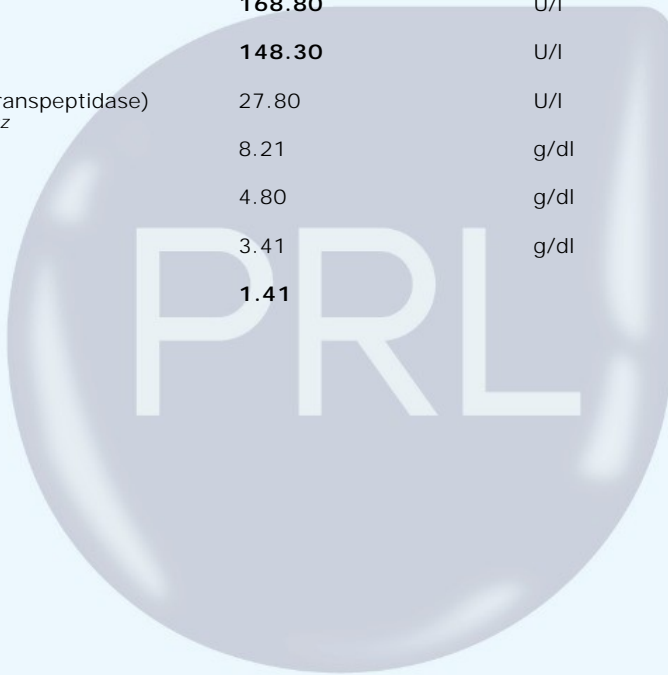


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LFT (Liver Function Test)

Serum Bilirubin Total <i>Method : Diazotized Sulfanilic Acid (DSA)</i>	0.71	mg/dl	0.1 - 1.2
Serum Bilirubin Direct <i>Method : Diazotized Sulfanilic Acid (DSA)</i>	0.21	mg/dl	0.0 - 0.3
Serum Bilirubin Indirect <i>Method : Calculated</i>	0.50	mg/dl	0.1 - 1.1
Serum SGOT/AST <i>Method : IFCC without P5P</i>	104.10	U/l	<= 35.0
Serum SGPT/ALT <i>Method : IFCC without P5P</i>	168.80	U/l	<= 45.0
Serum Alkaline Phosphatase <i>Method : PNP, AMP Buffer</i>	148.30	U/l	30.0 - 120.0
Serum GGT (Gamma Glutamyl Transpeptidase) <i>Method : UV-assay according to Szasz</i>	27.80	U/l	11.0 - 61.0
Serum total Protein <i>Method : Biuret</i>	8.21	g/dl	6.6 - 8.3
Serum Albumin <i>Method : Bromo Cresol Green</i>	4.80	g/dl	3.5 - 5.2
Serum Globulin <i>Method : Calculated</i>	3.41	g/dl	2.0 - 3.5
Albumin / Globulin ratio <i>Method : Calculated</i>	1.41		1.5 - 2.5



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Lipid Profile basic (direct HDL,calculated LDL)

Total Cholesterol, , serum Method : CHOD-POD	216.50	mg/dl	< 200.0
Triglycerides , serum Method : GPO-POD	127.50	mg/dl	< 150
HDL Cholesterol , serum Method : Direct measure PEG (CHE-CHO)	44.20	mg/dl	> 40
VLDL Cholesterol , serum Method : Calculated	25.50	mg/dl	< 30
L.D.L Cholesterol , serum Method : Calculated	146.80	mg/dl	< 100
Cholesterol, Non HDL , serum Method : Calculated	172.30	mg/dl	< 130
Total Cholesterol / HDL Cholesterol Ratio , serum Method : Calculated	4.90		< 5.0
LDL / HDL Cholesterol ratio , serum Method : Calculated	3.32		< 3.5

Interpretation:

National Lipid Association Recommendation (NLA-2014)	
Total Cholesterol Desirable: <200 mg/dL Borderline high: 200-239 mg/dL High: > or =240 mg/dL	Triglycerides Normal: <150 mg/dL Borderline high: 150-199 mg/dL High: 200-499 mg/dL Very high: > or =500 mg/dL
Non HDL Cholesterol Desirable: <130 mg/dL Borderline high: 130-159 mg/dL High: 160-189 mg/dL Very high: > or =190 mg/dL	LDL Cholesterol Optimal: <100 mg/dL Near Optimal: 100-129 mg/dL Borderline high: 130-159 mg/dL High: 160-189 mg/dL Very high: > or =190 mg/dL
HDL Cholesterol Low (Men) <40 mg/dL Low (Women) <50 mg/dL	

Phosphorus (inorganic), serum Method : Phosphomolybdate Method	2.14	mg/dl	2.5 - 4.5
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Interpretation:

Eighty-eight percent of the phosphorus contained in the body is localized in bone in the form of hydroxyapatite. The remainder is involved in intermediary carbohydrate metabolism and in physiologically important substances such as phospholipids, nucleic acids, and adenosine triphosphate (ATP). Phosphorus occurs in blood in the form of inorganic phosphate and organically bound phosphoric acid. The small amount of extracellular organic phosphorus is found exclusively in the form of phospholipids. Serum phosphate concentrations are dependent on meals and variation in the secretion of hormones such as parathyroid hormone (PTH) and may vary widely. Hypophosphatemia may have 4 general causes: shift of phosphate from extracellular to intracellular, renal phosphate wasting, loss from the gastrointestinal tract, and loss

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from intracellular stores.
 Hyperphosphatemia is usually secondary to an inability of the kidneys to excrete phosphate. Other factors may relate to increased intake or a shift of phosphate from the tissues into the extracellular fluid.
 Phosphate levels may be used in the diagnosis and management of a variety of disorders including bone, parathyroid and renal disease.
 Hypophosphatemia is relatively common in hospitalized patients. Levels less than 1.5 mg/dL may result in muscle weakness, hemolysis of red cells, coma, and bone deformity and impaired bone growth.
 The most acute problem associated with rapid elevations of serum phosphate levels is hypocalcemia with tetany, seizures, and hypotension. Soft tissue calcification is also an important long-term effect of high phosphorus levels.
 Phosphorus levels less than 1.0 mg/dL are potentially life-threatening and are considered a critical value.
 Note: Phosphorus has a very strong biphasic circadian rhythm. Values are lowest in the morning, peak first in the late afternoon and peak again in the late evening. The second peak is quite elevated and results may be outside the reference range.



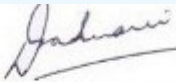
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Test Name	Value	Unit	Biological Reference Interval
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Vitamin B 12, serum 614.21 pg/ml 183.0 - 822.0
 Method : CLIA Microparticles

Please note change in biological reference interval.

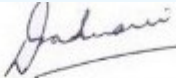
Interpretation:

Vitamin B12 (cobalamin) is necessary for hematopoiesis and normal neuronal function. In humans, it is obtained only from animal proteins and requires intrinsic factor (IF) for absorption. The body uses its vitamin B12 stores very economically, reabsorbing vitamin B12 from the ileum and returning it to the liver; very little is excreted. Vitamin B12 deficiency may be due to lack of IF secretion by gastric mucosa (eg, gastrectomy, gastric atrophy) or intestinal malabsorption (eg, ileal resection, small intestinal diseases). Vitamin B12 deficiency frequently causes macrocytic anemia, glossitis, peripheral neuropathy, weakness, hyperreflexia, ataxia, loss of proprioception, poor coordination, and affective behavioral changes. These manifestations may occur in any combination; many patients have the neurologic defects without macrocytic anemia. Serum methylmalonic acid and homocysteine levels are also elevated in vitamin B12 deficiency states. Follow-up testing for antibodies to intrinsic factor (IF) is recommended to identify this potential cause of vitamin B12 malabsorption. A normal serum concentration of vitamin B12 does not rule out tissue deficiency of vitamin B12. The most sensitive test for vitamin B12 deficiency at the cellular level is the assay for MMA. If clinical symptoms suggest deficiency, measurement of MMA and homocysteine should be considered, even if serum vitamin B12 concentrations are normal. The commonest cause of increased level of vitamin B12 is therapeutic intake of vitamin B12 in the form of multivitamin tablets or as intramuscular injections. Many other conditions are known to cause an increase or decrease in the serum vitamin B12 concentration including:

Increased Serum B12	Decreased Serum B12
Ingestion of vitamin C	Pregnancy
Ingestion of estrogens	Aspirin
Ingestion of vitamin A	Anticonvulsants
Hepatocellular injury	Colchicine
Myeloproliferative disorder	Ethanol ingestion
Uremia	Contraceptive hormones
	Smoking
	Hemodialysis
	Multiple myeloma

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Vitamin D (25 Hydroxy), serum 21.72 ng/ml 30.0 - 100.0
 Method : CLIA Microparticles

Interpretation:

Deficiency	ng/ml	< 20
Insufficiency	ng/ml	21 - 29
Sufficiency	ng/ml	30 - 100
Intoxication	ng/ml	> 150

Vitamin D compounds are derived from dietary ergocalciferol (from plants, VitD2) or cholecalciferol (from animals, VitD3), or by conversion of 7-dihydrocholesterol to VitD3 in the skin upon ultraviolet exposure. VitD2 and VitD3 are subsequently 25-hydroxylated in the liver to 25-OH-VitD. 25-OH-VitD represents the main body reservoir and transport form of vitamin D, being stored in adipose tissue and tightly bound by a transport protein while in circulation. A fraction of circulating 25-OH-VitD is converted to its active metabolites 1,25-dihydroxy vitamin D2 and D3 (1,25-OH-VitD), mainly by the kidneys. This process is regulated by parathyroid hormone (PTH). VitD plays a primary role in the maintenance of calcium homeostasis. It promotes intestinal calcium absorption and, in concert with PTH, skeletal calcium deposition, or less commonly, calcium mobilization. Renal calcium and phosphate reabsorption are also promoted. In addition to its effects on calcium and bone metabolism, 1,25-OH-VitD regulates the expression of a multitude of genes in many other tissues including immune cells, muscle, vasculature, and reproductive organs. The exact 25-OH-VitD level reflecting optimal body stores remains unknown. Mild-to-modest deficiency can be associated with osteoporosis or secondary hyperparathyroidism. Severe deficiency may lead to failure to mineralize newly formed osteoid in bone, resulting in rickets in children and osteomalacia in adults. The consequences of vitamin D deficiency on organs other than bone are not fully known, but may include increased susceptibility to infections, muscular discomfort, and an increased risk of colon, breast, and prostate cancer.

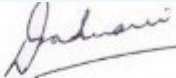
Reasons for suboptimal 25-OH-VitD levels include lack of sunshine exposure, a particular problem in India; inadequate intake; malabsorption (eg, due to Celiac disease); depressed hepatic vitamin D 25-hydroxylase activity, secondary to advanced liver disease; and enzyme-inducing drugs, in particular many antiepileptic drugs, including phenytoin, phenobarbital, and carbamazepine, that increase 25-OH-VitD metabolism.

Hypervitaminosis D is rare, and is only seen after prolonged exposure to extremely high doses of vitamin D. When it occurs, it can result in severe hypercalcemia and hyperphosphatemia.

Caution: Replacement therapy in deficient individuals must be monitored by periodic assessment of Vitamin D levels in order to prevent hypervitaminosis D.

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PSA Total, serum <i>Method : ECLIA</i>	0.38	ng/mL	0 - 3.1
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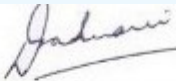
Interpretation:
Prostate-specific antigen (PSA) is a glycoprotein that is produced by the prostate gland, the lining of the urethra, and the bulbourethral gland. Normally, very little PSA is secreted in the blood. Increases in glandular size and tissue damage caused by benign prostatic hypertrophy, prostatitis, or prostate cancer may increase circulating PSA levels.
In patients with previously diagnosed prostate cancer, PSA testing is advocated as an early indicator of tumor recurrence and as an indicator of response to therapy. The test is also useful for initial screening for prostate cancer:
Total PSA levels < 2 ng/ml almost rule out the possibility of prostatic malignancy.
Total PSA levels between 2 and 10 ng/ml lie in the grey zone. Such values may be obtained in prostatitis, benign hyperplasia and malignancy. Further testing including a free PSA/PSA ratio and prostate biopsy is recommended for these patients for confirmation of the diagnosis.
Total PSA values >10 ng/ml are highly suspicious for prostate cancer but further testing, such as prostate biopsy, is needed to diagnose the exact pathology.



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Thyroid Profile Total (T3, T4, TSH)

T3, (Triiodothyronine) , serum Method : ECLIA	1.07	ng/mL	0.80 - 2.0
T4, (Thyroxine) , serum Method : ECLIA	6.55	ug/dL	5.1 - 14.1
TSH (Thyroid Stimulating Hormone) , serum Method : ECLIA	1.29	uIU/ml	0.27 - 4.2

Interpretation:

- Primary hyperthyroidism is accompanied by elevated serum T3 and T4 values alongwith depressed TSH levels
- Primary hypothyroidism is accompanied by depressed serum T3 and T4 values and elevated serum TSH levels.
- High T3 levels coupled with normal T4 and suppressed TSH may be seen in T3 toxicosis.

Note: Total T3 and total T4 are highly bound to plasma proteins and are amenable to fluctuations with plasma protein content as well as due to binding defects in the thyroid hormone binding proteins.

The following ranges are recommended for pregnant females:

Gestation period	TSH (uIU/ml)
First trimester	0.1 - 2.5
Second trimester	0.2 - 3.0
Third trimester	0.3 - 3.0

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Dr. Smita Sadwani
MD(Biochemistry)
Technical Director

Dr. Mayank Gupta
MD, DNB Pathology
Consultant Pathologist

Jadhav
Dr. Deepak Sadwani
MD(Pathology)
Lab Director

Dr. Moushmi Mukherjee
MBBS,MD (Pathology)
Consultant Pathologist

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Mobile:9313817732



Lab No.	012411200230	Age/Gender	54 YRS/MALE	Coll. ON	20/Nov/2024 09:16AM
NAME	Mr. JAGDISH			Reg. ON	20/Nov/2024
Ref. Dr.	MEDIWEEL	BarcodeNo	01200230	Approved ON	20/Nov/2024 01:05PM
Rpt. Centre	undefined			Printed ON	20/Nov/2024 06:52PM

Test Name	Value	Unit	Biological Reference Interval
-----------	-------	------	-------------------------------

Urine Routine & Microscopic Examination

Physical examination

Volume	40	mL	
Colour	Pale Yellow		Pale yellow
Transparency	Clear		Clear
Specific gravity	1.015		1.003 - 1.035

Method : pKa change

Chemical examination

Protein	Nil		Nil
Method : error-of-indicator			
Glucose	Nil		Nil
Method : GOD-POD			
pH	5.0		
Method : Double indicator			
Bilirubin	Negative		Negative
Method : Azo-coupling reaction			
Urobilinogen	Normal		Normal
Method : Azo- coupling reaction			
Ketone	Negative		Negative
Method : Legals test			
Erythrocytes	Absent		Absent
Method : Peroxidase			
Nitrite	Negative		Negative
Method : Griess reaction			
Leukocytes	Absent	Leu/uL	Negative
Method : Esterase activity of granulocytes			

Microscopic examination

WBC	0 - 1	/ HPF	0 - 2
RBC	Nil	/ HPF	0 - 2
Casts	Nil	/ HPF	Nil
Crystals	Nil	/ HPF	Nil
Epithelial cells	0 - 1	/ HPF	0 - 15
Bacteria	Absent		Absent
Others	Nil		

Method : Light microscopy

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Consultant Pathologist

Dr. Moushmi Mukherjee
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Consultant Pathologist

Mobile:9313817732

Lab No.	012411200230	Age/Gender	54 YRS/MALE	Coll. ON	20/Nov/2024 09:16AM
NAME	Mr. JAGDISH			Reg. ON	20/Nov/2024
Ref. Dr.	MEDIWEEL	BarcodeNo	01200230	Approved ON	20/Nov/2024 01:05PM
Rpt. Centre	undefined			Printed ON	20/Nov/2024 06:52PM

Test Name	Value	Unit	Biological Reference Interval
Urine Sugar fasting <i>Method : Hexokinase</i>	Nil		Nil
Urine Sugar PP <i>Method : Hexokinase</i>	NIL		NIL



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Lab Director

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Consultant Pathologist

Dr. Moushmi Mukherjee
MD Pathology
Consultant Pathologist

Lab No.	012411200230	Age/Gender	54 YRS/MALE	Coll. ON	20/Nov/2024 09:16AM
NAME	Mr. JAGDISH			Reg. ON	20/Nov/2024
Ref. Dr.	MEDIWEEL	BarcodeNo	01200230	Approved ON	20/Nov/2024 12:14PM
Rpt. Centre	Courier			Printed ON	20/Nov/2024 06:52PM

ECG Electro-cardiography

Normal ECG.



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Sadwani

Dr. Smita Sadwani
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Director
DMC Regd. No. 48732

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MD(Pathology)
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Dr. Ashish Gautam
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MBBS,MD (Pathology)
Consultant Pathologist

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Lab No.	012411200230	Age/Gender	54 YRS/MALE	Coll. ON	20/Nov/2024 09:16AM
NAME	Mr. JAGDISH			Reg. ON	20/Nov/2024
Ref. Dr.	MEDIWEEL	BarcodeNo	01200230	Approved ON	20/Nov/2024 01:38PM
Rpt. Centre	Courier			Printed ON	20/Nov/2024 06:52PM


TMT (Treadmill Test)
Negative For RMI.



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Dr. Anil Sahoo
MD. PGDCO
Reg. No.33201

Lab No.	012411200230	Age/Gender	54 YRS/MALE	Coll. ON	20/Nov/2024 09:16AM
NAME	Mr. JAGDISH			Reg. ON	20/Nov/2024
Ref. Dr.	MEDIWEEL	BarcodeNo	01200230	Approved ON	20/Nov/2024 01:35PM
Rpt. Centre	Courier			Printed ON	20/Nov/2024 06:52PM

Eye Vision		
	Right Eye	Left Eye
NEAR VISION	N/6 (With Glass)	N/6 (With Glass)
DISTANCE VISION	6/6 (With Glass)	6/6 (With Glass)
COLOR VISION	Normal	Normal

MER

General Condition	Fair, no pallor, no icterus, no anemia observed
Height (cm)	178
Weight (kg)	80
Pulse (bpm)	73
BP (mm/hg)	147/88

Please note: Kindly review with clinician in view of abnormal reports (if any).

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 Processing Centre : Prognosis Laboratories,515-516, Sector-19, Dwarka, Behind Gupta Properties.

 *Sadwani*
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Dr. Moushmi Mukherjee
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 Consultant Pathologist

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Mobile:9313817732

Lab No.	012411200230	Age/Gender	54 YRS/MALE	Coll. ON	20/Nov/2024 09:16AM
NAME	Mr. JAGDISH			Reg. ON	20/Nov/2024
Ref. Dr.	MEDIWEEL	BarcodeNo	01200230	Approved ON	20/Nov/2024 11:06AM
Rpt. Centre	Courier			Printed ON	20/Nov/2024 06:52PM

X-Ray Chest PA view

Trachea and mediastinum are central.

Bilateral lung fields are clear.

Bilateral hilar shadows are normal.

Bilateral costophrenic angles are clear.

Cardiac shadow is normal.

Soft tissue shadows and bony rib cage is normal.

Impression: No significant abnormality seen .

Please correlate clinically

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DR AMIT JAISWAL
MBBS,DMRD.DNB (RADIO DIAGNOSIS)
DMC No. 55709

Lab No.	012411200230	Age/Gender	54 YRS/MALE	Coll. ON	20/Nov/2024 09:16AM
NAME	Mr. JAGDISH			Reg. ON	20/Nov/2024
Ref. Dr.	MEDIWEEL	BarcodeNo	01200230	Approved ON	20/Nov/2024 10:37AM
Rpt. Centre	Courier			Printed ON	20/Nov/2024 06:52PM

SONOGRAPHY OF ABDOMEN AND PELVIS

The liver is normal in size (13.6 cm) *and shows mild diffuse increased parenchymal echogenicity.* There is no evidence of any focal hepatic lesion. The hepatic and portal veins are normal. There is no intrahepatic biliary dilatation.

The gall bladder is adequately distended. There is no evidence of any calculi. There is no evidence of any wall thickening seen. The CBD is not dilated.

The pancreas is well visualized and shows a normal parenchymal echotexture. There is no evidence of any focal mass, calcification or ductal dilatation seen. There is no peripancreatic fluid collection seen.

The spleen is normal in size (9.3 cm) and shows a normal parenchymal echotexture. There is no focal lesion seen.

The right kidney measures 11.2 x 4.4 cm and the left kidney measures 11.3 x 5.1 cm. Both kidneys are normal in size and shape. The kidneys show normal echotexture with a well-maintained cortical thickness. There is no evidence of hydronephrosis, cortical scarring or calculus disease in right kidney.

Left kidney shows few simple cortical cysts largest of them measuring 18 x 14 mm at interpolar region.

There is no ascites or bowel wall thickening.

The urinary bladder shows normal contours.

The prostate is normal in size.

IMPRESSION

- **Grade I fatty liver.**

Kindly correlate clinically.

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*** Partial Report ***



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DR AMIT JAISWAL
MBBS,DMRD.DNB (RADIO DIAGNOSIS)
DMC No. 55709

Mobile:9313817732

PROGNOSIS LABORATORIES

A SUBSIDIARY OF MEDGENOME

515-516 DWARKA SEC19 NEW DELHI 110075

Mr. JAGDISH

ID. : 361

AGE/SEX : 54 Yr /M

HT/WT : /

DATE : 20-11-2024 10:43:43 AM

REF.BY : Dr.MEDIWEEL

MACHINE INTERPRETATION : Normal ECG.

RATE : 61 bpm

BP : N/A

P Axis : -5 deg.

QRS Axis : -8 deg.

T Axis : -1 deg.

P Duration : 125 ms

PR Duration : 168 ms

QRS Duration : 85 ms

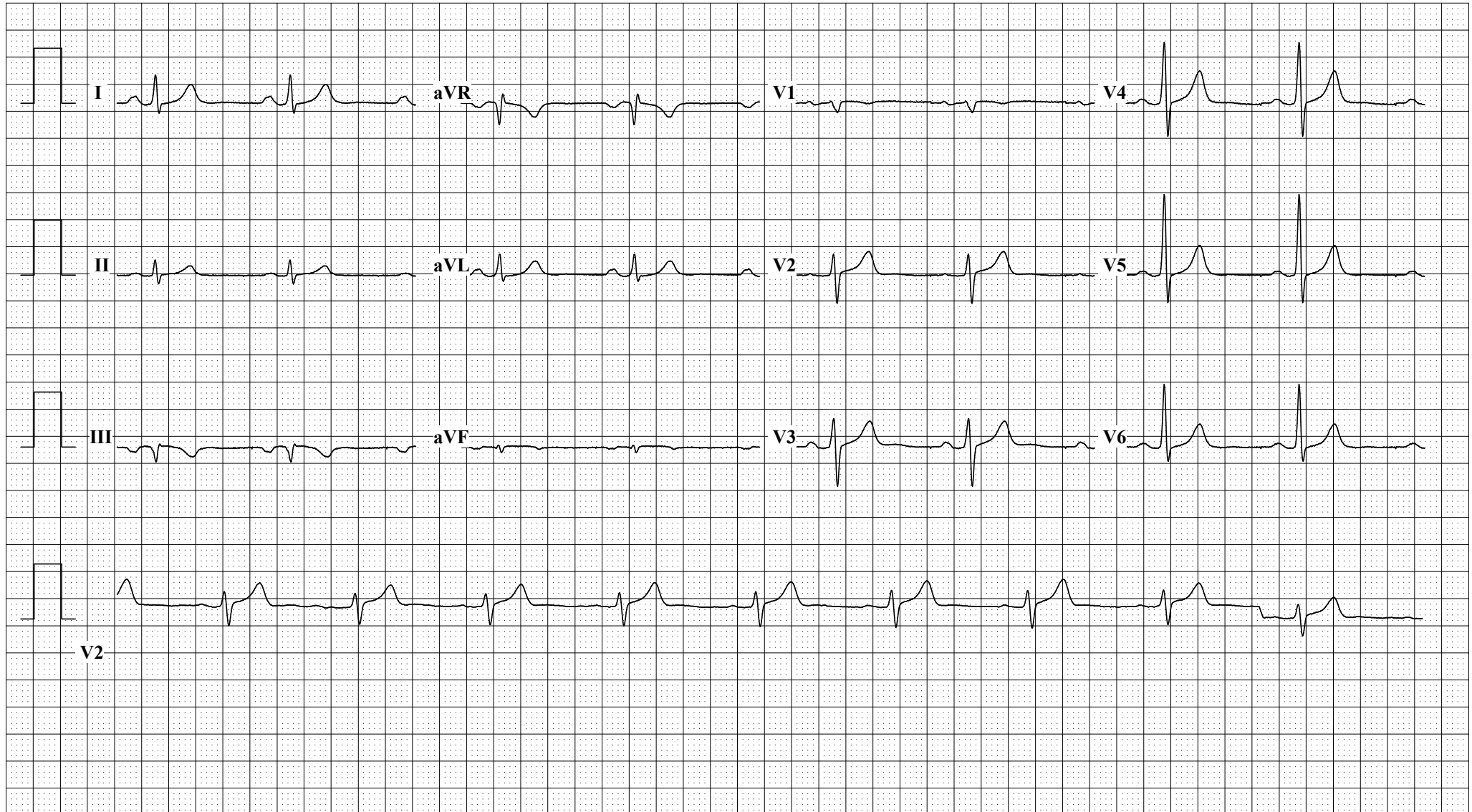
QT Interval : 350 ms

QTc Interval : 354 ms

Linked Median

Speed : 25 mm/s

Sensitivity : 10 mm/mV



PROGNOSIS LABORATORIES

JAGDISH
I.D. 583
Age 54/M
Date 20-11-2024

RATE 68bpm
B.P. 140/88

PRETEST
SUPINE

ST @ 10mm/mV
80ms PostJ

LINKED MEDIAN



PROGNOSIS LABORATORIES

JAGDISH
I.D. 583
Age 54/M
Date 20-11-2024

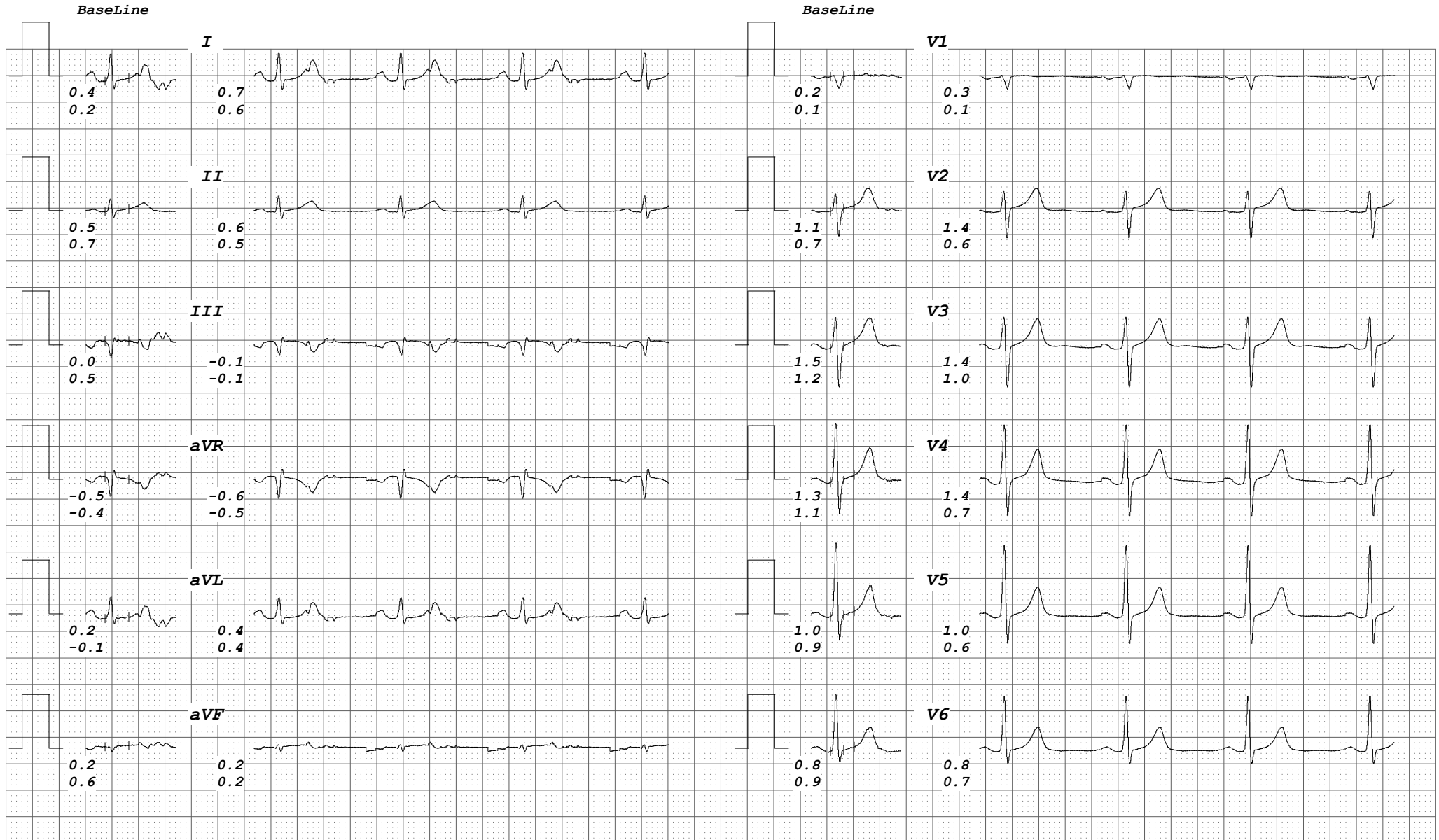
RATE 66bpm
B.P. 140/88

PRETEST
HYPERVENT

ST @ 10mm/mV
80ms PostJ

PHASE TIME 0:16

LINKED MEDIAN



PROGNOSIS LABORATORIES

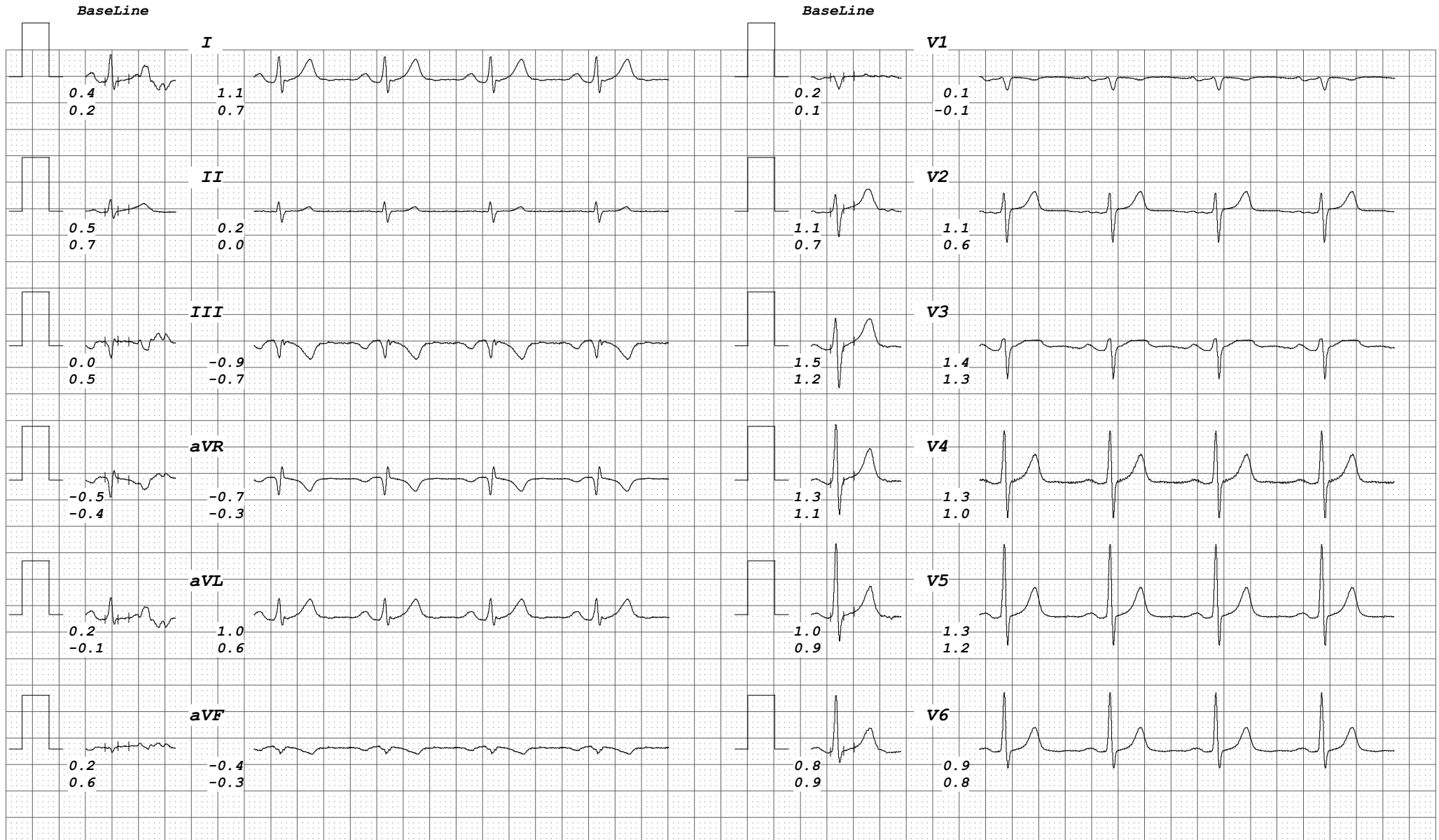
JAGDISH
I.D. 583
Age 54/M
Date 20-11-2024

RATE 76bpm
B.P. 140/88

PRETEST
STANDING

ST @ 10mm/mV
80ms PostJ

LINKED MEDIAN



PROGNOSIS LABORATORIES

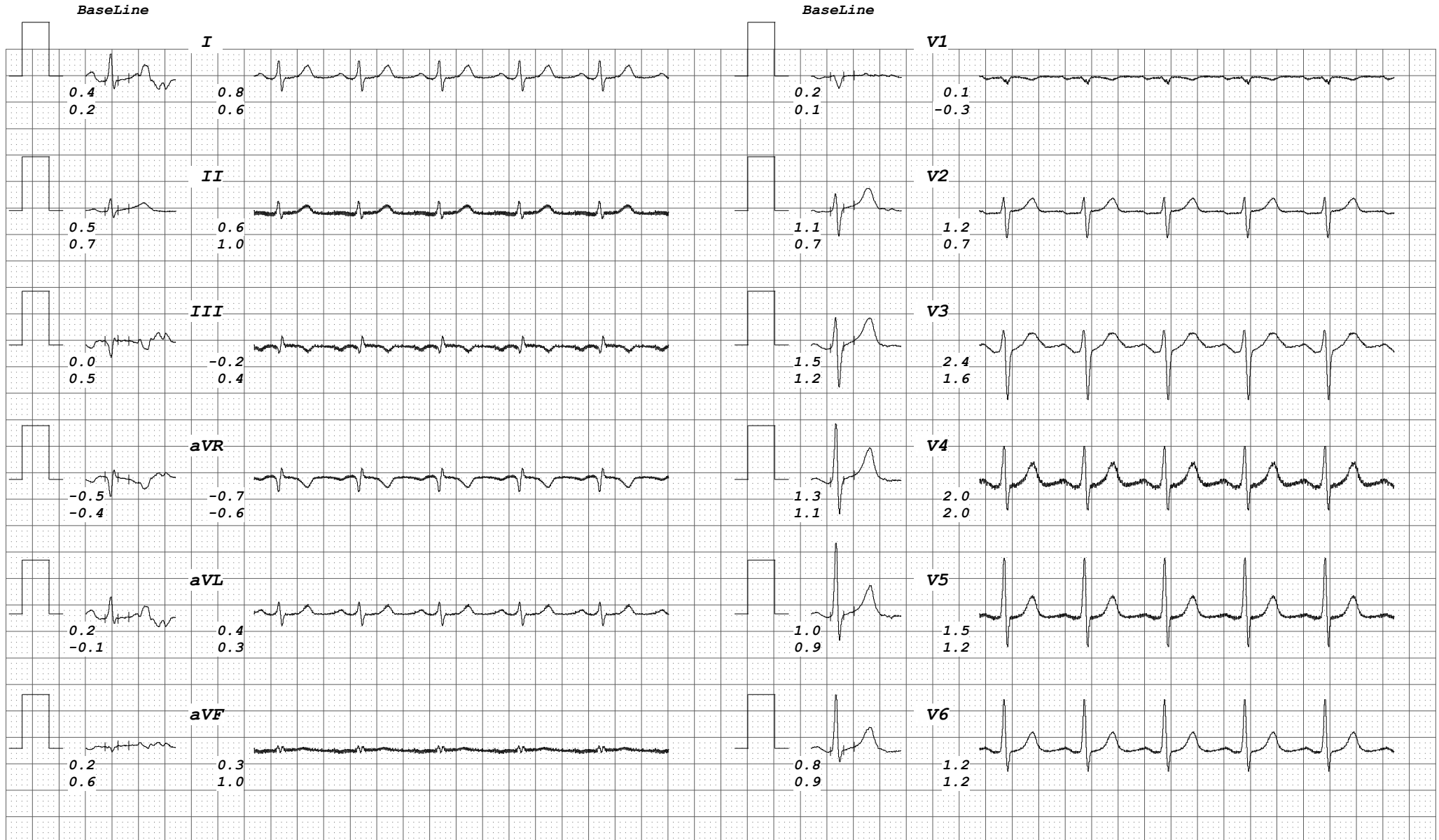
JAGDISH
I.D. 583
Age 54/M
Date 20-11-2024

RATE 100bpm
B.P. 142/90

Bruce
Stage 1
TOTAL TIME 2:55
PHASE TIME 2:55

ST @ 10mm/mV
80ms PostJ
Speed 2.7 km/hr
SLOPE 10 %

LINKED MEDIAN



PROGNOSIS LABORATORIES

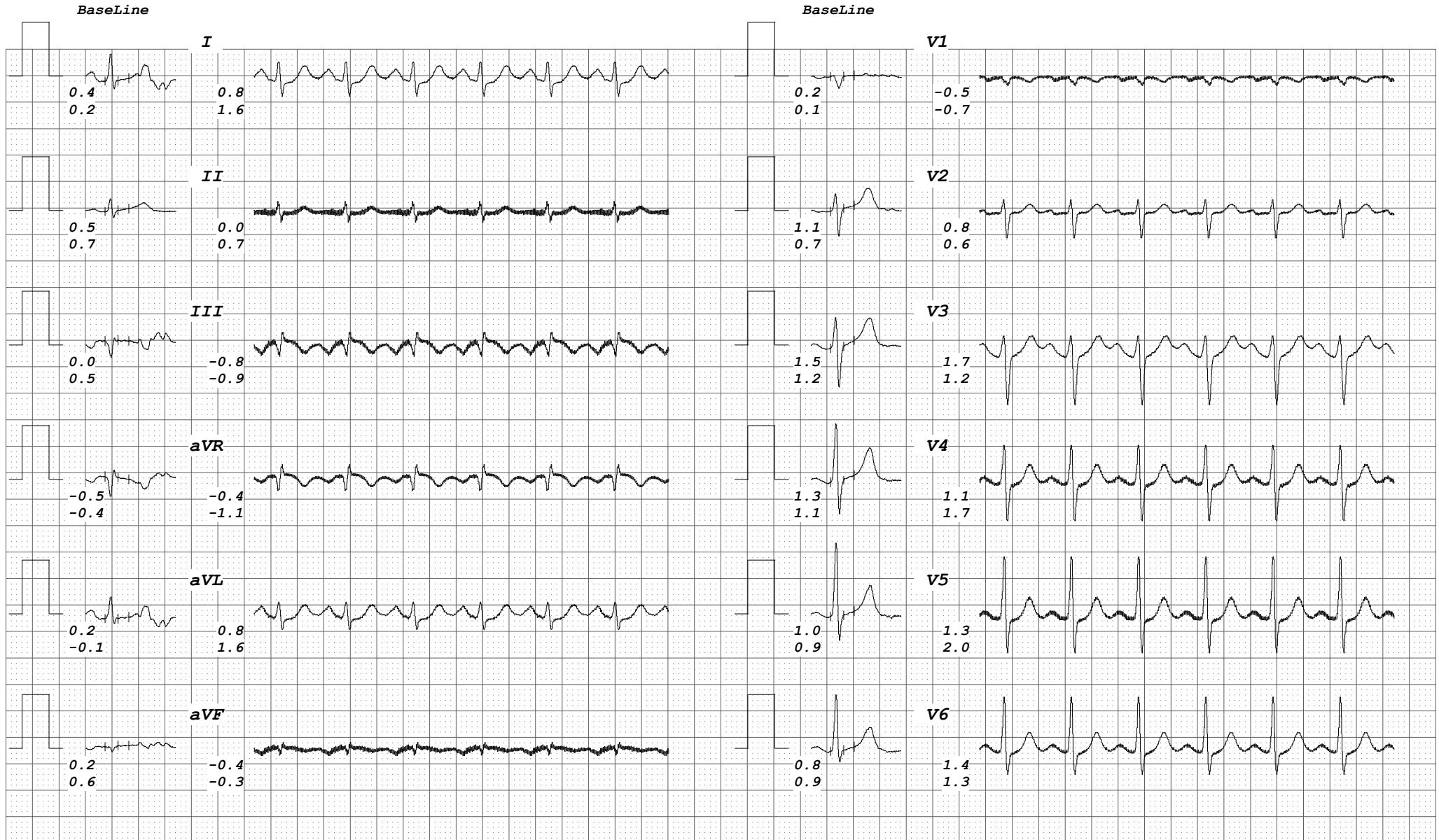
JAGDISH
I.D. 583
Age 54/M
Date 20-11-2024

RATE 120bpm
B.P. 144/92

Bruce
Stage 2
TOTAL TIME 5:55
PHASE TIME 2:55

ST @ 10mm/mV
80ms PostJ
Speed 4 km/hr
SLOPE 12 %

LINKED MEDIAN



PROGNOSIS LABORATORIES

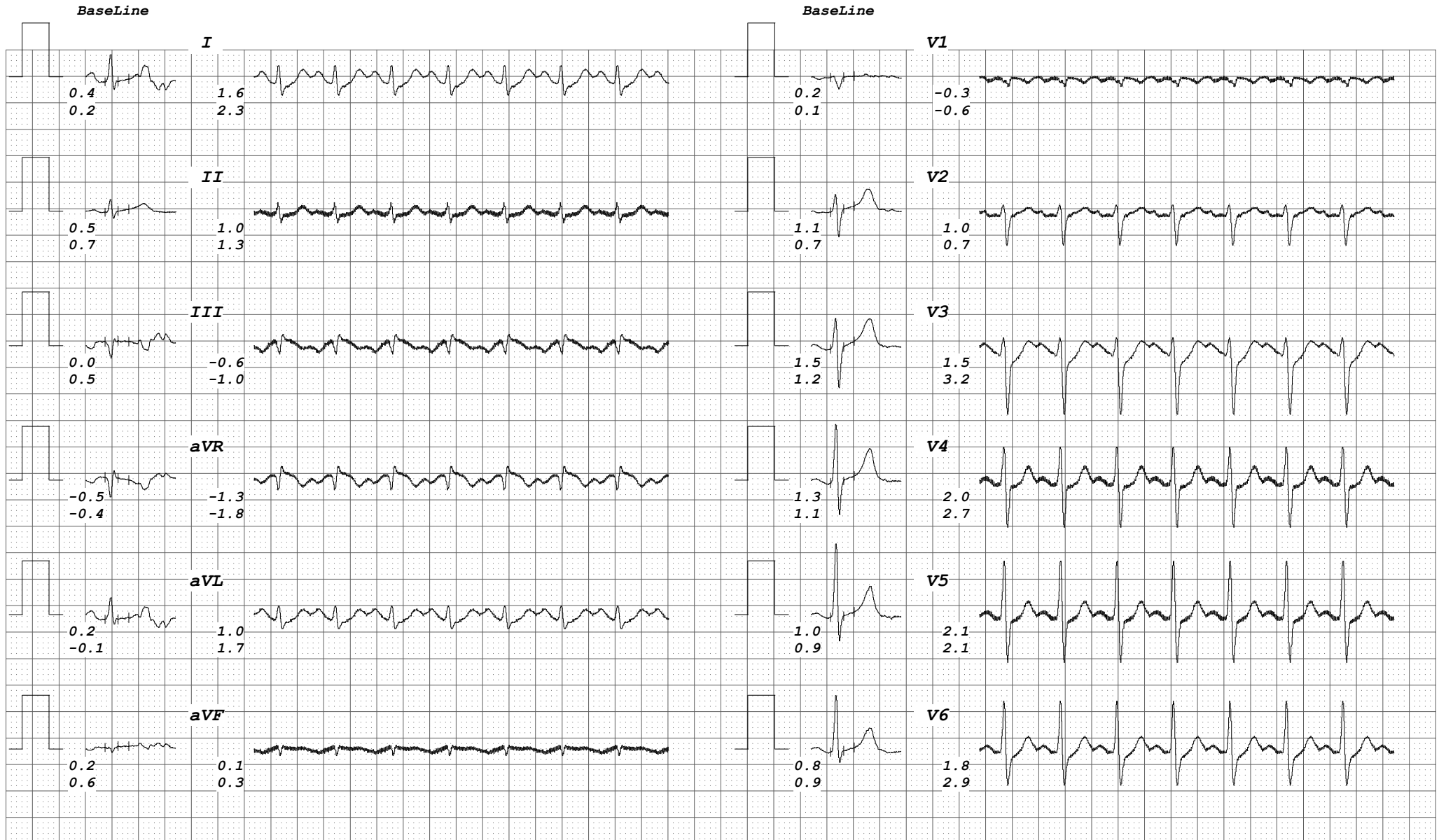
JAGDISH
I.D. 583
Age 54/M
Date 20-11-2024

RATE 142bpm
B.P. 146/94

Bruce
PK-EXERCISE
TOTAL TIME 7:47
PHASE TIME 1:47

ST @ 10mm/mV
80ms PostJ
Speed 5.4 km/hr
SLOPE 14 %

LINKED MEDIAN



PROGNOSIS LABORATORIES

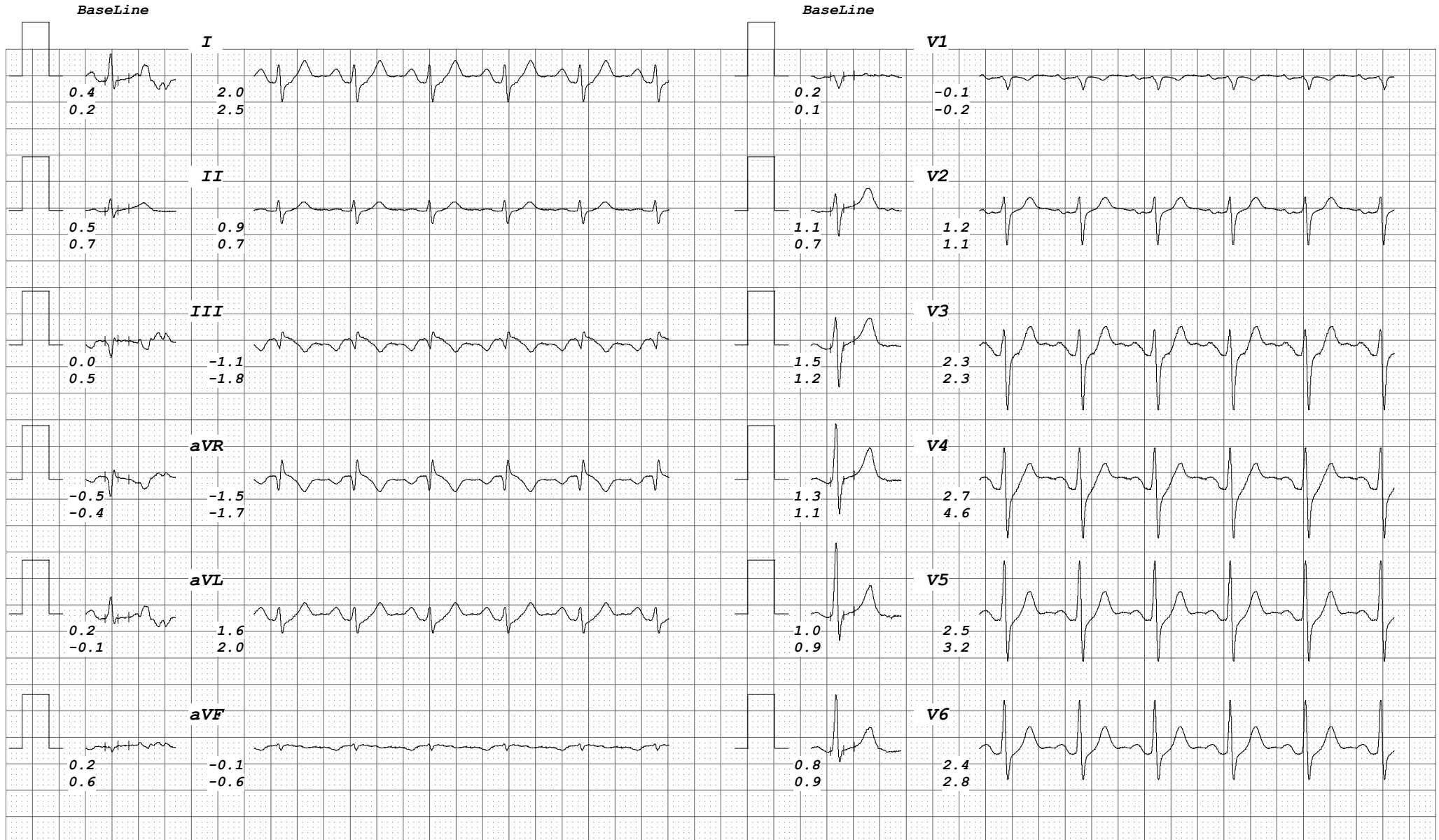
JAGDISH
I.D. 583
Age 54/M
Date 20-11-2024

RATE 107bpm
B.P. 144/92

Bruce
RECOVERY
TOTAL TIME 8:57
PHASE TIME 0:55

ST @ 10mm/mV
80ms PostJ

LINKED MEDIAN



PROGNOSIS LABORATORIES

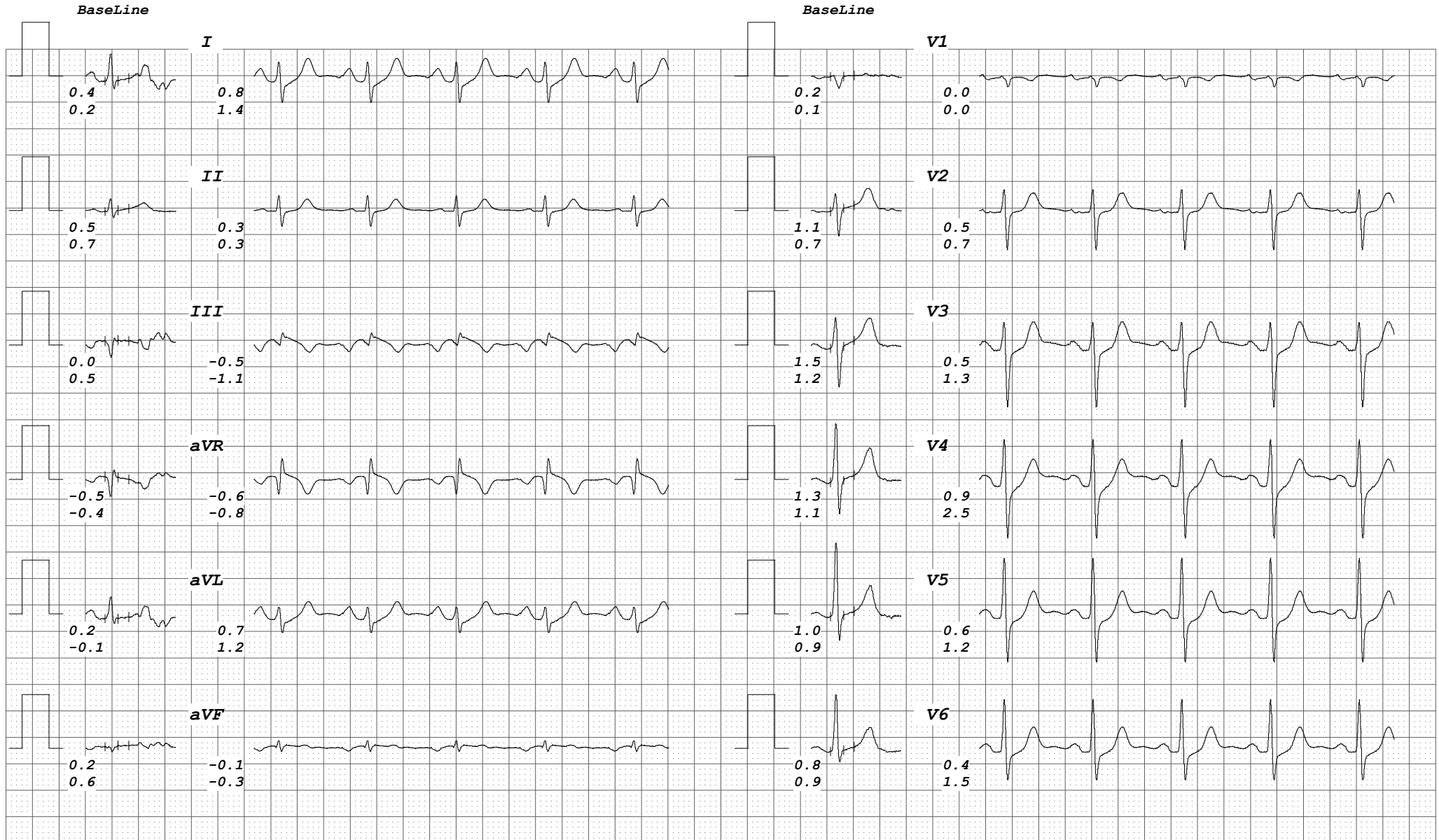
JAGDISH
I.D. 583
Age 54/M
Date 20-11-2024

RATE 90bpm
B.P. 140/88

Bruce
RECOVERY
TOTAL TIME 9:57
PHASE TIME 1:55

ST @ 10mm/mV
80ms PostJ

LINKED MEDIAN



PROGNOSIS LABORATORIES

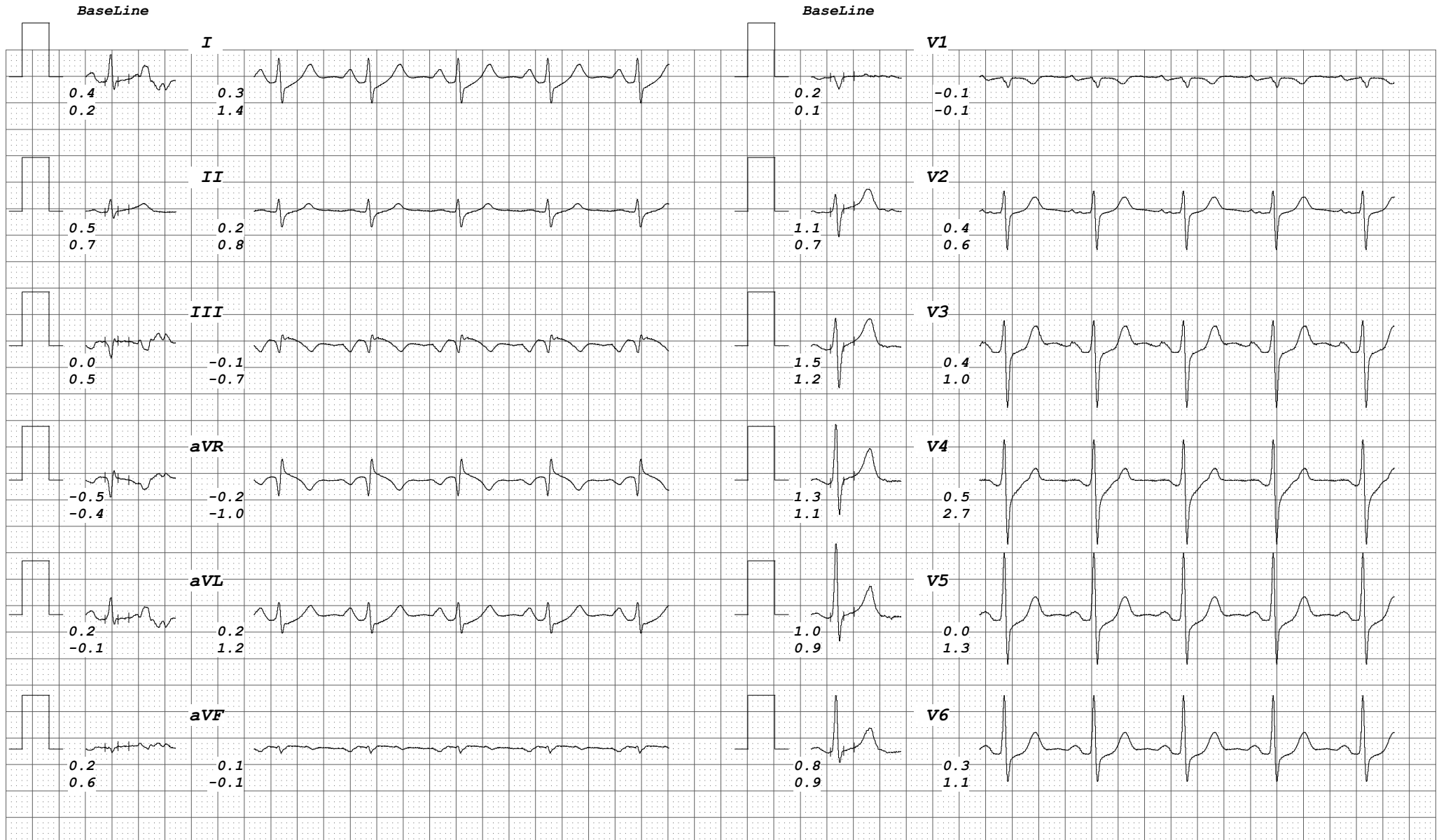
JAGDISH
I.D. 583
Age 54/M
Date 20-11-2024

RATE 90bpm
B.P. 140/88

Bruce
RECOVERY
TOTAL TIME 10:57
PHASE TIME 2:55

ST @ 10mm/mV
80ms PostJ

LINKED MEDIAN





भारतीय विशिष्ट पहचान प्राधिकरण
भारत सरकार
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Government of India



E-Aadhaar Letter

नामांकन क्रमांक/Enrolment No.: 1452/18091/00122

Jagdish Prasad Yadav (जगदीश प्रसाद यादव)
RZF-907/1 S/F, M.G MARG, Raj Nagar-2, Bagdola,
South West Delhi,
Delhi - 110077

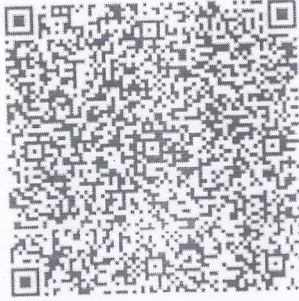
सूचना

- आधार पहचान का प्रमाण है, नागरिकता का नहीं।
- पहचान का प्रमाण ऑनलाइन ऑथेंटिकेशन द्वारा प्राप्त करें।
- यह एक इलेक्ट्रॉनिक प्रक्रिया द्वारा बना हुआ पत्र है।

Date: 29/06/2015

आपका आधार क्रमांक/ Your Aadhaar No.:

4284 2981 5051



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- To establish identity, authenticate online.
- This is electronically generated letter.

आधार-आम आदमी का अधिकार



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1800 300 1947



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Signature Not Verified
Digitally signed by Sandeep Bhardwaj
Date: 2015.06.29 11:58:23 IST

- आधार देश भर में मान्य है।
- आधार के लिए आपको एक ही बार नामांकन दर्ज करवाने की आवश्यकता है।
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UNIQUE IDENTIFICATION AUTHORITY OF INDIA



जगदीश प्रसाद यादव
Jagdish Prasad Yadav
जन्म तिथि/ DOB: 02/01/1970
पुरुष / MALE



पता:

आरजेडमार्ग-907/1 एस/एफ,
एम.जी मार्ग, राज नगर-2,
बगडोला, दक्षिण पश्चिमी
दिल्ली,
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Address:

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Delhi - 110077

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आधार-आम आदमी का अधिकार

Aadhaar-Aam Admi ka Adhikar