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Name	: Mrs . SWATHI OLADRI	TID	: UMR1893441
Age/Gender	: 34 Years/Female	Registered On	: 26-Aug-2024 08:33 AM
Ref By	: Self	Reported On	: 26-Aug-2024 01:32 PM
Reg.No	: BIL4634115	Reference	: Arcofemi Health Care Ltd - Medi Whe

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DEPARTMENT OF ULTRASOUND  
**Ultrasound Whole Abdomen**

**LIVER** is normal shape, size (13.9 cms) and has uniform echopattern. No evidence of focal lesion. No intrahepatic biliary ductal dilatation. Hepatic and portal vein radicals are normal.

**GALL BLADDER** : Contracted. No evident calculi. CBD is of normal calibre.

**PANCREAS** : Head and body are normal.

**SPLEEN** shows normal shape, size (8.6 cms) and echopattern.

**KIDNEYS** move well with respiration and have normal shape, size and echopattern. Cortico- medullary differentiations are well madeout. No evidence of calculus or hydronephrosis. Right kidney measures : 8.8 x 4.0 cms, Left kidney measures : 8.6 x 4.7 cms.

**URINARY BLADDER** shows normal shape and wall thickness. It has clear contents. No evidence of diverticula.

**UTERUS** is anteverted has normal shape and size. It has uniform myometrial echopattern. Endometrial echo is of normal thickness : 10 mm. Uterus measures : 10.4 x 5.0 x 5.5 cms.

**OVARIES** are normal in size, shape and echotexture. Right ovary: 2.8 x 1.5 cms, Left ovary: 2.8 x 1.6 cms.

No evidence of free fluid in the abdomen and pelvis.



PLEASE SCAN QR CODE

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Name : Mrs . SWATHI OLADRI  
Age/Gender : 34 Years/Female  
Ref By : Self  
Reg.No : BIL4634115

TID : UMR1893441  
Registered On : 26-Aug-2024 08:33 AM  
Reported On : 26-Aug-2024 01:32 PM  
Reference : Arcofemi Health Care Ltd  
- Medi Whe

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**IMPRESSION:**

**\* NO SONOGRAPHIC ABNORMALITY DETECTED.**

- Suggested clinical correlation and follow up.

\*\*\* End Of Report \*\*\*

**Dr. Apoorva K**  
Consultant Radiologist



PLEASE SCAN QR CODE

Name : Mrs . SWATHI OLADRI  
Age/Gender : 34 Years/Female  
Ref By : Self  
Reg.No : BIL4634115

TID : UMR1893441  
Registered On : 26-Aug-2024 08:33 AM  
Reported On : 26-Aug-2024 04:48 PM  
Reference : Arcofemi Health Care Ltd  
- Medi Whe

DEPARTMENT OF X-RAY  
**X-Ray Chest PA View**

**CLINICAL DETAILS :** Health checkup.

**FINDINGS:**

Lung fields appear normal.

Cardiac size is within normal limits.

Aorta and pulmonary vasculature is normal.

Bilateral domes of diaphragm and costophrenic angles are normal.

Visualised bones and soft tissues appear normal.

**IMPRESSION:**

**\* NORMAL STUDY.**

- Suggested clinical correlation and follow up.

\*\*\* End Of Report \*\*\*

**Dr.Kiranchander**  
Reg.No - 58122  
Consultant Radiologist



Name	: MRS.SWATHI OLADRI	TID/SID	: UMR1893441/ 28141081
Age / Gender	: 34 Years / Female	Registered on	: 26-Aug-2024 / 08:33 AM
Ref.By	: SELF	Collected on	: 26-Aug-2024 / 08:33 AM
Req.No	: BIL4634115	Reported on	: 26-Aug-2024 / 10:30 AM
		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**DEPARTMENT OF CLINICAL PATHOLOGY**

**Complete Urine Examination (CUE), Urine**

Investigation	Result	Biological Reference Intervals
<b>Physical Examination</b>		
Colour Method:Physical	LightYellow	Straw to Yellow
Appearance Method:Physical	Clear	Clear
<b>Chemical Examination</b>		
Reaction and pH Method:Indicator	Acidic (5.0)	4.6-8.0
Specific gravity Method:Refractometry	1.002	1.000-1.035
Protein Method:Protein Error of pH indicators	Negative	Negative
Glucose Method:Glucose oxidase/Peroxidase	Negative	Negative
Blood Method:Peroxidase	Negative	Negative
Ketones Method:Sodium Nitroprusside	Negative	Negative
Bilirubin Method:Diazonium salt	Negative	Negative
Leucocytes Method:Esterase reaction	Negative	Negative
Nitrites Method:Modified Griess reaction	Negative	Negative
Urobilinogen Method:Diazonium salt	Negative	Up to 1.0 mg/dl (Negative)
<b>Microscopic Examination</b>		
Pus cells (leukocytes) Method:Flow Digital Imaging/Microscopy	2-3	2 - 3 /hpf
Epithelial cells Method:Flow Digital Imaging/Microscopy	2-3	2 - 5 /hpf
RBC (erythrocytes) Method:Flow Digital Imaging/Microscopy	Absent	Absent
Casts Method:Flow Digital Imaging/Microscopy	Absent	Occasional hyaline casts may be seen



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Registered on : 26-Aug-2024 / 08:33 AM  
Collected on : 26-Aug-2024 / 08:33 AM  
Reported on : 26-Aug-2024 / 10:30 AM  
Reference : Arcofemi Health Care Ltd -

**TEST REPORT**

Crystals	Absent	Phosphate, oxalate, or urate crystals may be seen
Method:Flow Digital Imaging/Microscopy		
Others	Nil	Nil
Method:Flow Digital Imaging/Microscopy		

**Method: Semi Quantitative test ,For CUE**

**Reference:** Godkar Clinical Diagnosis and Management by Laboratory Methods, First South Asia edition. Product kit literature.

**Interpretation:**

The complete urinalysis provides a number of measurements which look for abnormalities in the urine. Abnormal results from this test can be indicative of a number of conditions including kidney disease, urinary tract infection or elevated levels of substances which the body is trying to remove through the urine . A urinalysis test can help identify potential health problems even when a person is asymptomatic. All the abnormal results are to be correlated clinically.

\* Sample processed at National Reference Laboratory,  
Tenet Diagnostics,Hyderabad

--- End Of Report ---

**Dr Vikas Reddy**  
Consultant Pathologist





Name	: MRS.SWATHI OLADRI	TID/SID	: UMR1893441/ 28143068
Age / Gender	: 34 Years / Female	Registered on	: 26-Aug-2024 / 08:33 AM
Ref.By	: SELF	Collected on	: 26-Aug-2024 / 12:40 PM
Req.No	: BIL4634115	Reported on	: 27-Aug-2024 / 17:03 PM
		Reference	: Arcofemi Health Care Ltd -

TEST REPORT

DEPARTMENT OF CYTOPATHOLOGY

Pap Smear, Conventional

Cytology No	<b>C-9143/24</b>
Clinical Details	For screening.
Specimen Type	Conventional smear (Pap smear)
Specimen Adequacy	Satisfactory for evaluation without evidence of endocervical/transformation zone component
Microscopic Observations:	Smear contains superficial, intermediate and parabasal cells. Dense inflammation noted.
Organisms	Not present
Non-neoplastic findings	<b>Reactive cellular changes associated with inflammation</b>
Interpretation	<b>Negative for intraepithelial lesion or malignancy.</b>
.	<b>Inflammatory smear</b>
Note	Advice repeat after control of inflammation

**Method** : Pap staining & microscopy

Reported as per the 2014 Bethesda System

\* Sample processed at National Reference Laboratory,  
Tenet Diagnostics, Hyderabad

--- End Of Report ---

**Dr Shruti Reddy**  
Consultant Pathologist  
Reg No.TSMC/FMR/22656





Name	: MRS.SWATHI OLADRI	TID/SID	: UMR1893441/ 28141082
Age / Gender	: 34 Years / Female	Registered on	: 26-Aug-2024 / 08:33 AM
Ref.By	: SELF	Collected on	: 26-Aug-2024 / 08:33 AM
Req.No	: BIL4634115	Reported on	: 26-Aug-2024 / 12:47 PM
		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**DEPARTMENT OF HEMATOPATHOLOGY**

**Blood Grouping ABO And Rh Typing, EDTA Whole Blood**

Parameter	Results
Blood Grouping (ABO)	A
Rh Typing (D)	Positive
Method:Hemagglutination Tube Method by Forward & Reverse Grouping	

**Method:** Hemagglutination Tube Method by Forward & Reverse Grouping

**Reference:** Tulip kit literature

**Interpretation:** The ABO grouping and Rh typing test determines blood type grouping (A,B, AB, O ) and the Rh factor (positive or negative). A person's blood type is based on the presence or absence of certain antigens on the surface of their red blood cells and certain antibodies in the plasma. ABO antigens are poorly expressed at birth, increase gradually in strength and become fully expressed around 1 year of age. In case of Rh(D) - Du(weak positive) or Weak D positive, the individual must be considered as Rh positive as donor and Rh negative as recipient.

**Note:** Records of previous blood grouping/Rh typing not available. Please verify before transfusion.

\* Sample processed at National Reference Laboratory, Tenet Diagnostics,Hyderabad

--- End Of Report ---

**Dr Shruti Reddy**  
Consultant Pathologist  
Reg No.TSMC/FMR/22656





Name	: MRS.SWATHI OLADRI	TID/SID	: UMR1893441/ 28141082
Age / Gender	: 34 Years / Female	Registered on	: 26-Aug-2024 / 08:33 AM
Ref.By	: SELF	Collected on	: 26-Aug-2024 / 08:33 AM
Req.No	: BIL4634115	Reported on	: 26-Aug-2024 / 12:28 PM
		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**DEPARTMENT OF HEMATOPATHOLOGY**

**Erythrocyte Sedimentation Rate (ESR), Whole Blood**

Investigation	Observed Value	Biological Reference Intervals
ESR 1st Hour Method:Westergren/Vesmatic	<b>15</b>	<=12 mm/hour

**Complete Blood Count (CBC), EDTA Whole Blood**

Investigation	Observed Value	Biological Reference Intervals
Hemoglobin Method:Cyanide Free Lyse Hemoglobin	12.3	12.0-15.0 g/dL
PCV/HCT Method:Calculated	36.7	36.0-46.0 vol%
Total RBC Count Method:Electrical Impedance	<b>4.84</b>	3.80-4.80 mill /cu.mm
MCV Method:Calculated	<b>75.9</b>	83.0-101.0 fL
MCH Method:Calculated	<b>25.4</b>	27.0-32.0 pg
MCHC Method:Calculated	33.4	31.5-34.5 g/dL
RDW (CV) Method:Calculated	<b>17.0</b>	11.6-14.0 %
MPV Method:Calculated	8.1	7.0-10.0 fL
Total WBC Count Method:Electrical Impedance	5380	4000-10000 cells/cumm
Platelet Count Method:Electrical Impedance	2.86	1.50-4.10 lakhs/cumm
<b>Differential count</b>		
Neutrophils Method:Microscopy	50.3	40.0-80.0 %
Lymphocytes Method:Microscopy	38.3	20.0-40.0 %
Eosinophils	1.4	1.0-6.0 %
Monocytes	9.1	2.0-10.0 %
Basophils Method:Microscopy	<b>0.9</b>	< 1.0-2.0 %





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Reference : Arcofemi Health Care Ltd -

**TEST REPORT**

Absolute Neutrophil Count	2706	2000-7000 cells/cumm
Method:Calculated		
Absolute Lymphocyte Count (ALC)	2061	1000-3000 cells/cumm
Absolute Eosinophil Count (AEC)	75	20-500 cells/cumm
Absolute Monocyte Count	490	200-1000 cells/cumm
Method:Calculated		
Absolute Basophil Count	48	20-100 cells/cumm
Method:Calculated		
Neutrophil - Lymphocyte Ratio(NLR)	1.31	0.78-3.53
Method:Calculated		

**Method:** Automated Hematology Cell Counter, Microscopy

**Reference:** Dacie and Lewis Practical Hematology, 12th Edition.  
Wallach's interpretation of diagnostic tests, Soth Asian Edition.

**Interpretation:** A Complete Blood Picture (CBP) is a screening test which can aid in the diagnosis of a variety of conditions and diseases such as anemia, leukemia, bleeding disorders and infections. This test is also useful in monitoring a person's reaction to treatment when a condition which affects blood cells has been diagnosed. All the abnormal results are to be correlated clinically.

**Note:** These results are generated by a fully automated hematology analyzer and the differential count is computed from a total of several thousands of cells. Therefore the differential count appears in decimalised numbers and may not add upto exactly 100. It may fall between 99 and 101.

\* Sample processed at National Reference Laboratory,  
Tenet Diagnostics,Hyderabad

--- End Of Report ---

**Dr Shruti Reddy**  
Consultant Pathologist  
Reg No.TSMC/FMR/22656





Name	: MRS.SWATHI OLADRI	TID/SID	: UMR1893441/ 28141084F
Age / Gender	: 34 Years / Female	Registered on	: 26-Aug-2024 / 08:33 AM
Ref.By	: SELF	Collected on	: 26-Aug-2024 / 08:33 AM
Req.No	: BIL4634115	Reported on	: 26-Aug-2024 / 11:53 AM
		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

**Blood Urea Nitrogen (BUN), Serum**

Investigation	Observed Value	Biological Reference Interval
Blood Urea Nitrogen. Method:Calculated	13	6-20 mg/dL
Urea. Method:Urease/UV	26.8	12.8-42.8 mg/dL

**Interpretation:** Urea is a waste product formed in the liver when protein is metabolized. Urea is released by the liver into the blood and is carried to the kidneys, where it is filtered out of the blood and released into the urine. Since this is a continuous process, there is usually a small but stable amount of urea nitrogen in the blood. However, when the kidneys cannot filter wastes out of the blood due to disease or damage, then the level of urea in the blood will rise. The blood urea nitrogen (BUN) evaluates kidney function in a wide range of circumstances, to diagnose kidney disease, and to monitor people with acute or chronic kidney dysfunction or failure. It also may be used to evaluate a person's general health status as well.

**Reference:** Tietz Fundamentals of Clinical Chemistry and Molecular Diagnostics

**Creatinine, Serum**

Investigation	Observed Value	Biological Reference Interval
Creatinine. Method:Alkaline Picrate	0.7	0.50-0.90 mg/dL

**Interpretation:**

Creatinine is a nitrogenous waste product produced by muscles from creatine. Creatinine is majorly filtered from the blood by the kidneys and released into the urine, so serum creatinine levels are usually a good indicator of kidney function. Serum creatinine is more specific and more sensitive indicator of renal function as compared to BUN because it is produced from muscle at a constant rate and its level in blood is not affected by protein catabolism or other exogenous products. It is also not reabsorbed and very little is secreted by tubules making it a reliable marker. Serum creatinine levels are increased in pre renal, renal and post renal azotemia, active acromegaly and gigantism. Decreased serum creatinine levels are seen in pregnancy and increasing age.

**Glucose Fasting (FBS), Sodium Fluoride Plasma**

Investigation	Observed Value	Biological Reference Interval
Glucose Fasting Method:Hexokinase	94	Normal: <100 mg/dL Impaired FG: 100-125 mg/dL Diabetes mellitus: >=126 mg/dL

**Interpretation:** It measures the Glucose levels in the blood with a prior fasting of 9-12 hours. The test helps screen a symptomatic/ asymptomatic person who is at risk for Diabetes. It is also used for regular monitoring of glucose levels in people with Diabetes.

**Reference:** American Diabetes Association. Standards of Medical Care in Diabetes-2022



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Req.No	: BIL4634115	Reported on	: 26-Aug-2024 / 11:55 AM
		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**Glucose Post Prandial (PPBS), Sodium Fluoride Plasma**

Investigation	Observed Value	Biological Reference Interval
Glucose Post Prandial Method:Hexokinase	95	Normal : <140 mg/dL Impaired PG: 140-199 mg/dL Diabetes mellitus: >=200 mg/dL

**Interpretation:** This test measures the blood sugar levels 2 hours after a normal meal. Abnormally high blood sugars 2 hours after a meal reflect that the body is not producing sufficient insulin which is indicative of Diabetes.

**Reference:** American Diabetes Association. Standards of Medical Care in Diabetes-2022

**Glycosylated Hemoglobin (HbA1C), EDTA Whole Blood**

Investigation	Observed Value	Biological Reference Interval
Glycosylated Hemoglobin (HbA1c) Method:High-Performance Liquid Chromatography	5.3	Non-diabetic: <= 5.6 % Pre-diabetic: 5.7 - 6.4 % Diabetic: >= 6.5 %
Estimated Average Glucose (eAG) Method:Calculated	105	mg/dL

**Interpretation:**

It is an index of long-term blood glucose concentrations and a measure of the risk for developing microvascular complications in patients with diabetes. Absolute risks of retinopathy and nephropathy are directly proportional to the mean HbA1c concentration. In persons without diabetes, HbA1c is directly related to risk of cardiovascular disease.

1) Low glyated haemoglobin (below 4%) in a non-diabetic individual are often associated with systemic inflammatory diseases, chronic anaemia (especially severe iron deficiency & haemolytic), chronic renal failure and liver diseases. Clinical correlation suggested.

2) Interference of Hemoglobinopathies in HbA1c estimatiion:

- A. For HbF > 25%, an alternate platform (Fructosamine) is recommended for testing of HbA1c.
- B. Homozygous hemoglobinopathy is detected, fructosamine is recommended for monitoring diabetic status
- C. Heterozygous state detected (D10 is corrected for HbS and HbC trait).

3) In known diabetic patients, HbA1c can be considered as a tool for monitoring the glycemc control.

- Excellent Control - 6 to 7 %,
- Fair to Good Control - 7 to 8 %,
- Unsatisfactory Control - 8 to 10 %
- and Poor Control - More than 10 %.

**Reference:** American Diabetes Association. Standards of Medical Care in Diabetes-2022.

**Bun/Creatinine Ratio, Serum**

Investigation	Observed Value	Biological Reference Interval
BUN/Creatinine Ratio Method:Calculated	17.0	10-20



Name	: MRS.SWATHI OLADRI	TID/SID	: UMR1893441/ 28141083
Age / Gender	: 34 Years / Female	Registered on	: 26-Aug-2024 / 08:33 AM
Ref.By	: SELF	Collected on	: 26-Aug-2024 / 08:33 AM
Req.No	: BIL4634115	Reported on	: 26-Aug-2024 / 11:55 AM
		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**Interpretation:**

The BUN/Creatinine ratio blood test is used to diagnose acute or chronic renal disease. BUN (blood urea nitrogen) and creatinine are both filtered in the kidneys and excreted in urine. The two together are used to measure overall kidney function

1. Increased ratio (>20) with normal creatinine occurs in the following conditions:

- a) Increased BUN (prerenal azotemia), heart failure, salt depletion, dehydration
- b) Catabolic states with tissue breakdown
- c) GI hemorrhage
- d) Impaired renal function plus excess protein intake, production, or tissue breakdown

2. Increased ratio (>20) with elevated creatinine occurs in the following conditions:

- a) Obstruction of urinary tract
- b) Prerenal azotemia with renal disease

3. Decreased ratio (<10) with decreased BUN occurs in the following conditions:

- a) Acute tubular necrosis
- b) Decreased urea synthesis as in severe liver disease or starvation
- c) Repeated dialysis
- d) SIADH
- e) Pregnancy

4. Decreased ratio (<10) with increased creatinine occurs in the following conditions:

- a) Phenacemide therapy (accelerates conversion of creatine to creatinine)
- b) Rhabdomyolysis (releases muscle creatinine)
- c) Muscular patients who develop renal failure

\* Sample processed at National Reference Laboratory,  
Tenet Diagnostics,Hyderabad

--- End Of Report ---

**Dr Afreen Anwar**  
Consultant Biochemist





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Ref.By	: SELF	Collected on	: 26-Aug-2024 / 08:33 AM
Req.No	: BIL4634115	Reported on	: 26-Aug-2024 / 11:55 AM
		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

**Lipid Profile, Serum**

Investigation	Observed Value	Biological Reference Interval
Total Cholesterol Method:Cholesterol Oxidase	169	Desirable: <200 mg/dL Borderline: 200-239 mg/dL High: >=240 mg/dL
HDL Cholesterol Method:Direct Measurement	53	Low: <40 mg/dL High: >=60 mg/dL
VLDL Cholesterol Method:Calculated	10.80	6.0-38.0 mg/dL
LDL Cholesterol Method:Calculated	105.2	Optimum: <100 mg/dL Near/above optimum: 100-129 mg/dL Borderline: 130-159 mg/dL High: 160-189 mg/dL Very high: >=190 mg/dL
Triglycerides Method:Glycerol LPL/GK	54	Normal:<150 mg/dL Borderline: 150-199 mg/dL High: 200-499 mg/dL Very high: >=500 mg/dL
Chol/HDL Ratio Method:Calculated	<b>3.19</b>	Low Risk: 3.3-4.4 Average Risk: 4.5-7.1 Moderate Risk: 7.2-11.0
LDL Cholesterol/HDL Ratio Method:Calculated	1.98	Desirable: 0.5-3.0 Borderline Risk: 3.0-6.0 High Risk: >6.0

**Interpretation:** Lipids are fats and fat-like substances which are important constituents of cells and are rich sources of energy. A lipid profile typically includes total cholesterol, high density lipoproteins (HDL), low density lipoprotein (LDL), chylomicrons, triglycerides, very low density lipoproteins (VLDL), Cholesterol/HDL ratio .The lipid profile is used to assess the risk of developing a heart disease and to monitor its treatment. The results of the lipid profile are evaluated along with other known risk factors associated with heart disease to plan and monitor treatment. Treatment options require clinical correlation.

**Reference:** Third Report of the National Cholesterol Education program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III), JAMA 2001.

\* Sample processed at National Reference Laboratory,  
Tenet Diagnostics,Hyderabad

--- End Of Report ---

**Dr Afreen Anwar**  
Consultant Biochemist



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Age / Gender	: 34 Years / Female	Registered on	: 26-Aug-2024 / 08:33 AM
Ref.By	: SELF	Collected on	: 26-Aug-2024 / 08:33 AM
Req.No	: BIL4634115	Reported on	: 26-Aug-2024 / 11:55 AM
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**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

**Liver Function Test (LFT), Serum**

Investigation	Observed Value	Biological Reference Interval
Total Bilirubin. Method:Diazo method	0.42	<1.2 mg/dL
Direct Bilirubin. Method:Diazo method	0.24	<0.30 mg/dL
Indirect Bilirubin. Method:Calculated	0.18	<0.9 mg/dL
Alanine Aminotransferase ,(ALT/SGPT) Method:UV wihout P5P	14	<34 U/L
Aspartate Aminotransferase,(AST/SGOT) Method:UV wihout P5P	21	<31 U/L
ALP (Alkaline Phosphatase). Method:PNPP-AMP Buffer	63	35-104 U/L
Gamma GT. Method:Gamma-Glutamyl - 3 - Carbossi - 4 - Nitroanilide (GCNA)	14	6-42 U/L
Total Protein. Method:Biuret	7.1	6.6-8.7 g/dL
Albumin. Method:Bromocresol Green (BCG)	4.4	3.5-5.2 g/dL
Globulin. Method:Calculated	2.70	1.8-3.8 g/dL
A/GRatio. Method:Calculated	1.60	0.8-2.0

**Interpretation:** Liver functions tests help to identify liver disease, its severity, and its type. Generally these tests are performed in combination, are abnormal in liver disease, and the pattern of abnormality is indicative of the nature of liver disease. An isolated abnormality of a single liver function test usually means a non-hepatic cause. If several liver function tests are simultaneously abnormal, then hepatic etiology is likely.

\* Sample processed at National Reference Laboratory,  
Tenet Diagnostics,Hyderabad

--- End Of Report ---

**Dr Afreen Anwar**  
Consultant Biochemist



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**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

**Thyroid Profile (T3,T4,TSH), Serum**

Investigation	Observed Value	Biological Reference Interval
Triiodothyronine Total (T3) Method:ECLIA	1.28	0.80-2.00 ng/mL Pregnancy: 1st Trimester: 0.81 - 1.90 ng/mL 2nd & 3rd Trimester: 1.00 - 2.60 ng/mL
Thyroxine Total (T4) Method:ECLIA	9.4	5.1-14.1 µg/dL
Thyroid Stimulating Hormone (TSH) Method:ECLIA	1.46	0.27-4.20 µIU/mL Pregnancy: 1st Trimester: 0.1 - 2.5 µIU/mL 2nd Trimester: 0.2 - 3.0 µIU/mL 3rd Trimester: 0.3 - 3.0 µIU/mL

**Interpretation:**

A thyroid profile is used to evaluate thyroid function and/or help diagnose hypothyroidism and hyperthyroidism due to various thyroid disorders. T4 and T3 are hormones produced by the thyroid gland. They help control the rate at which the body uses energy, and are regulated by a feedback system. TSH from the pituitary gland stimulates the production and release of T4 (primarily) and T3 by the thyroid. Most of the T4 and T3 circulate in the blood bound to protein. A small percentage is free (not bound) and is the biologically active form of the hormones.

**Reference:** Tietz textbook of Clinical Chemistry and Molecular Diagnostics, Nader Rifa, Andrea Ritas Horvath, Carl T. Wittwer.

\* Sample processed at National Reference Laboratory,  
Tenet Diagnostics,Hyderabad

--- End Of Report ---

**Dr Afreen Anwar**  
Consultant Biochemist





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**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

**Uric Acid, Serum**

Investigation	Observed Value	Biological Reference Interval
Uric Acid. Method:Uricase	4.5	2.4-5.7 mg/dL

**Interpretation**

It is the major product of purine catabolism. Hyperuricemia can result due to increased formation or decreased excretion of uric acid which can be due to several causes like metabolic disorders, psoriasis, tissue hypoxia, pre-eclampsia, alcohol, lead poisoning, acute or chronic kidney disease, etc. Hypouricemia may be seen in severe hepato cellular disease and defective renal tubular reabsorption of uric acid.

\* Sample processed at National Reference Laboratory,  
Tenet Diagnostics,Hyderabad

--- End Of Report ---

**Dr Afreen Anwar**  
Consultant Biochemist





**TENET MEDCORP PVT LTD**  
**GACHIBOWLI, HYDERABAD.**

4634115/SWATHI OLADRI 34 Yrs/Female 56 Kg/157 Cms

Date: 26-Aug-2024 12:35:51 PM

Ref. By : ARCOFEMI HEALTH CARE

Medication : Nil

Objective :

**Summary**

Protocol : BRUCE  
 History : Nil

Stage	StageTime (Min:Sec)	PhaseTime (Min:Sec)	Speed (kmph)	Grade (%)	METS	H.R. (bpm)	B.P. (mmHg)	R.P.P. (x100)	PVC	Comments
Supine					1.0	84	---/---	0	-	
Standing					1.0	91	---/---	0	-	
ExStart					1.0	86	---/---	0	-	
Stage 1	3:01	3:02	2.7	10.0	4.6	101	---/---	0	-	
Stage 2	3:01	6:02	4.0	12.0	7.0	117	120/80	140	-	
Stage 3	3:01	9:02	5.5	14.0	10.2	150	140/80	210	-	
PeakEx	0:19	9:20	6.8	16.0	10.6	164	140/80	229	-	
Recovery	1:00		0.0	0.0	4.3	141	150/80	211	-	
Recovery	3:00		0.0	0.0	1.0	108	130/80	140	-	

**Findings :**

Exercise Time : 9:19 minutes  
 Max HR attained : 160 bpm 86% of Max Predictable HR 186  
 Max BP : 150/80(mmHg)  
 Workload attained : 10.6 (Good Effort Tolerance )  
 No significant ST segment changes noted during exercise or recovery.  
 No Angina / Arrhythmia / SOB

Final Impression:\*\*\* TEST IS NEGATIVE FOR EXERCISE INDUCIBLE SICHEMIA \*\*\*

REPORTED BY DR RAJESH K  
 Dr. Rajesh K. S. (London)

Advicer/Comments:



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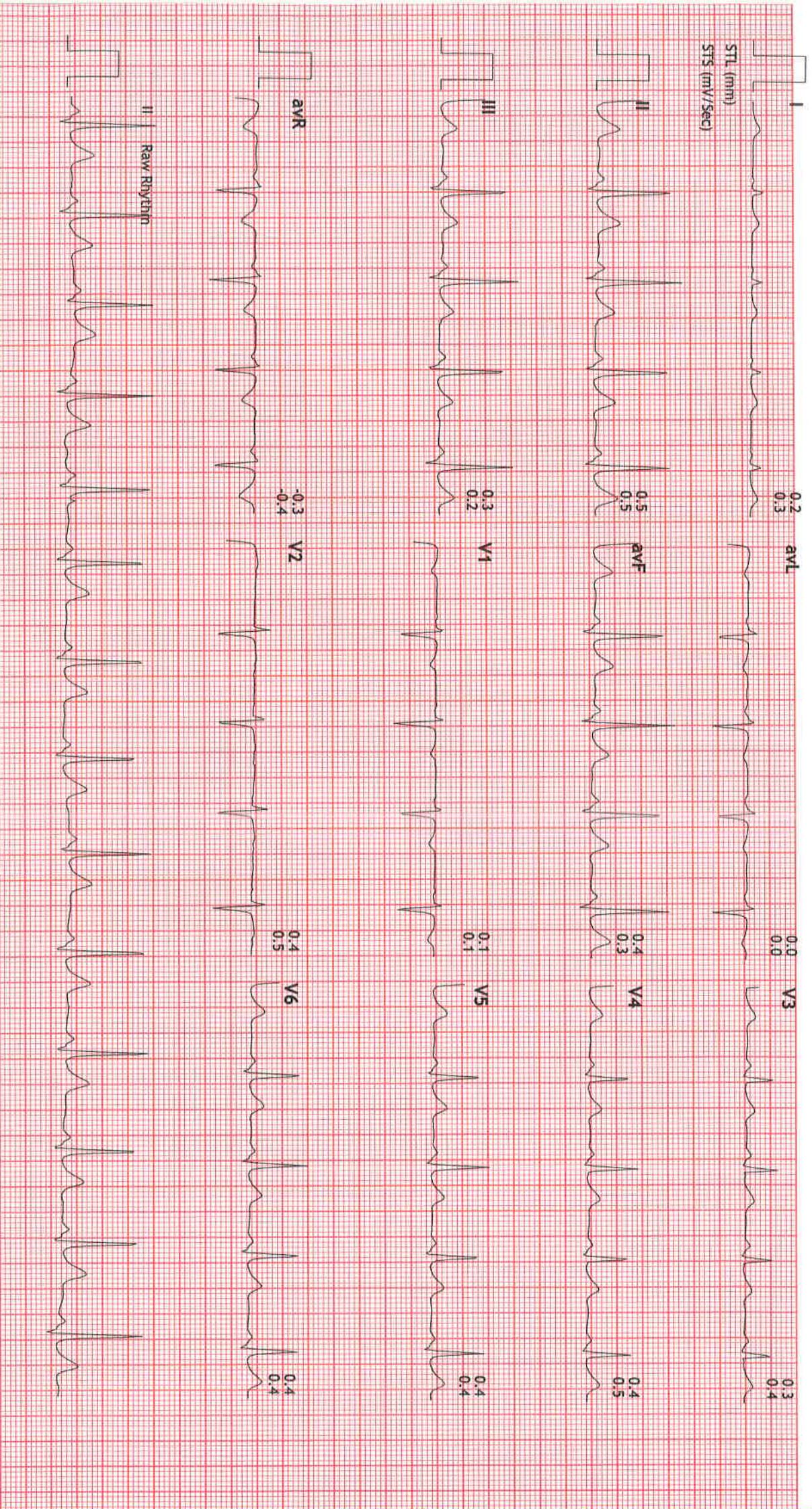
4634115/SWATHI OLADRI  
34 Yrs/Female  
56 Kg/157 Cms  
Date: 26-Aug-2024 12:35:51 PM

HR: 84 bpm  
METS: 1.0  
BP: ---/---  
Stage Report Time: 26-Aug-2024 12:38:15 PM

Linked Medians Report  
BRUCE  
(0.05-100)Hz

Ex Time 01:04  
BLC :On  
Notch :On

SUPINE  
10.0 mm/mV  
25 mm/Sec.



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4634115/SWATHI OLADRI  
34 Yrs/Female  
56 Kg/157 Cms  
Date: 26-Aug-2024 12:35:51 PM

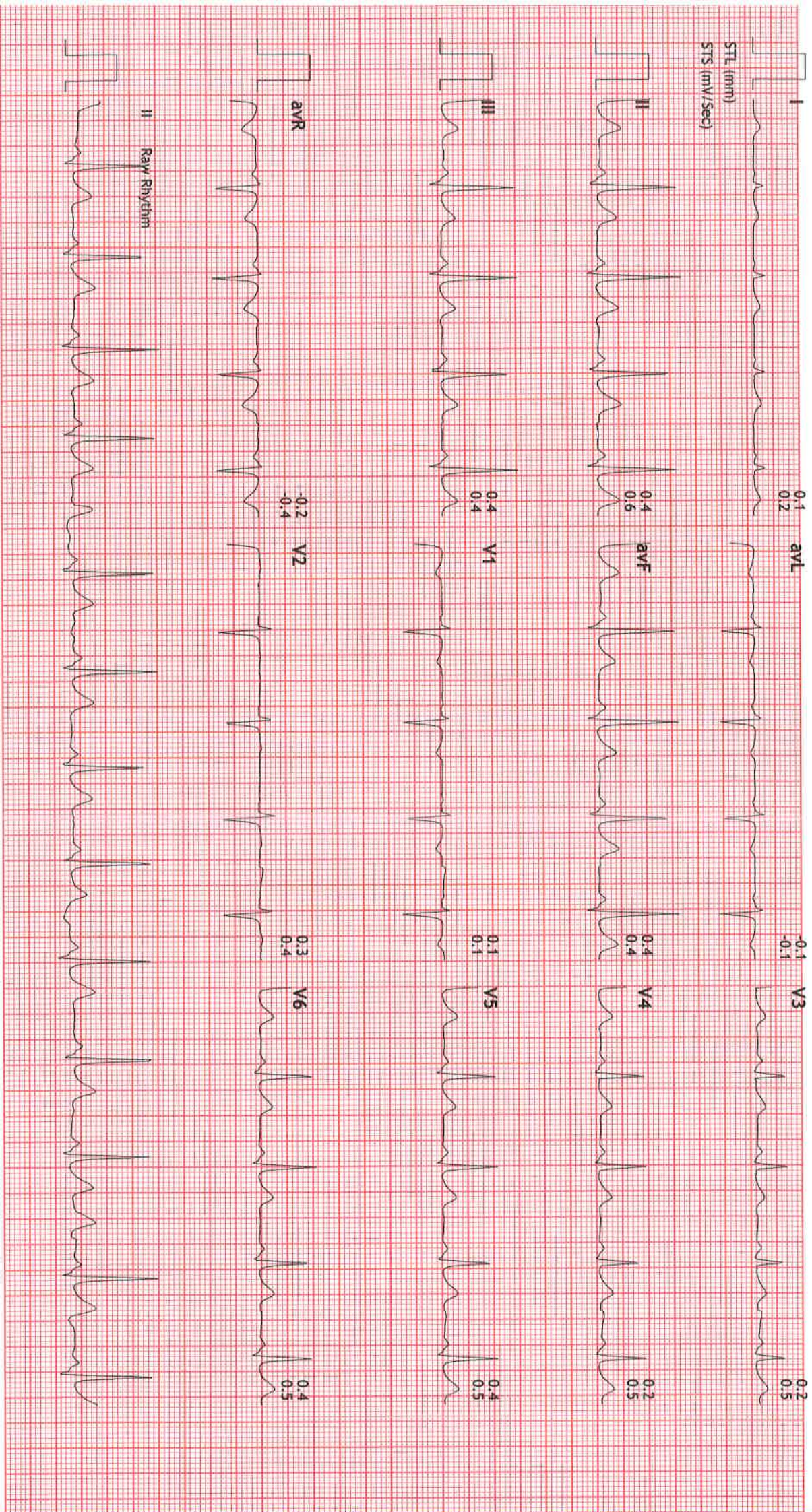
Linked Medians Report

HR: 91 bpm  
METS: 1.0  
BP: ---/---  
Stage Report Time: 26-Aug-2024 12:38:24 PM

MPHR: 48% of 186  
Speed: 0.0 kmph  
Grade: 0.0%

BRUCE  
(0.05-100)Hz  
Ex Time 01:13  
BLC :On  
Notch :On

STANDING  
10.0 mm/mV  
25 mm/Sec.



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4634115/SWATHI OLADRI  
34 Yrs/Female  
56 Kg/157 Cms  
Date: 26-Aug-2024 12:35:51 PM

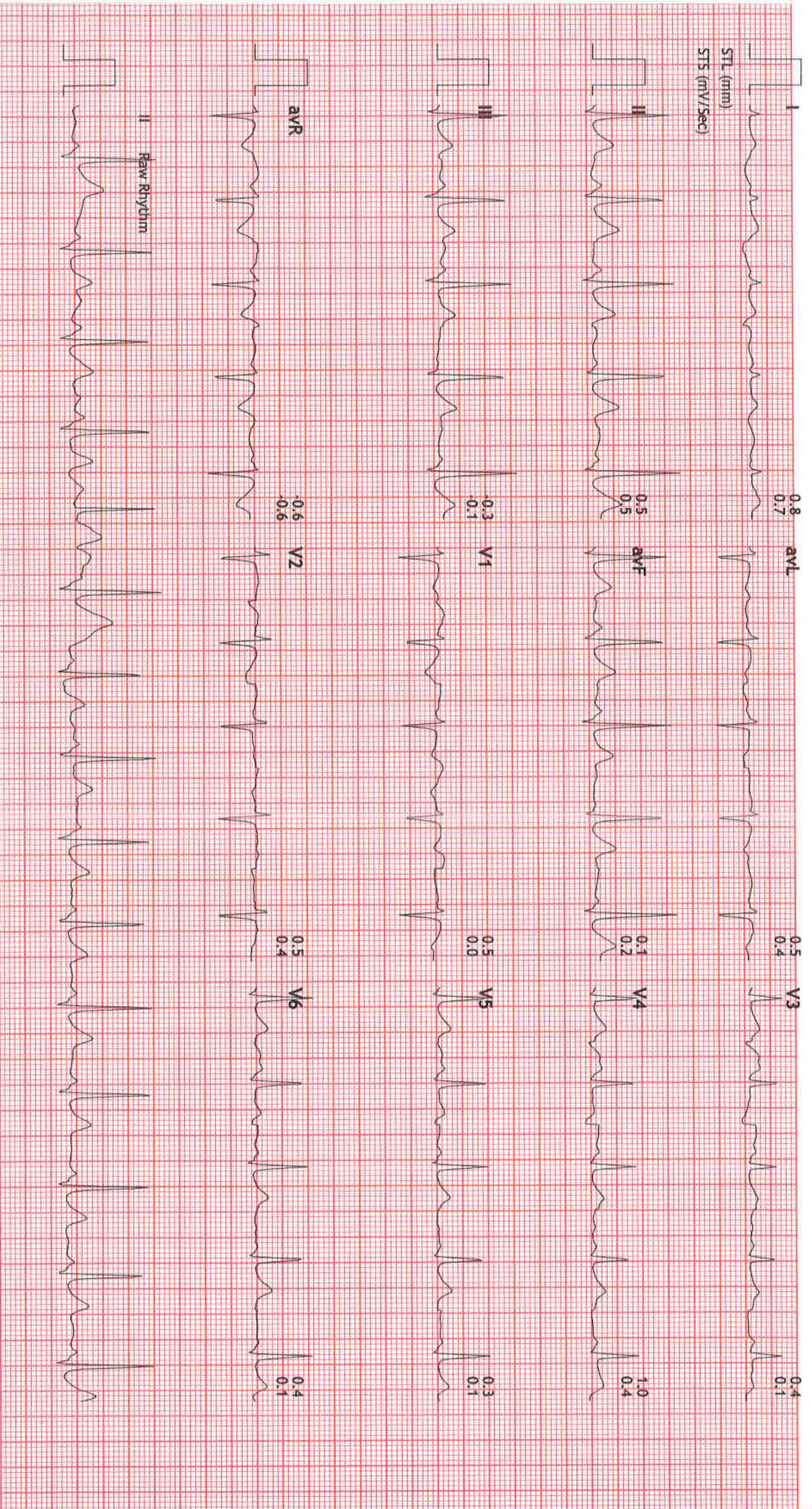
Linked Medians Report

HR: 86 bpm  
METs: 1.0  
BP: ----/----  
MPPHR: 46% of 186  
Speed: 0.0 kmph  
Grade: 0.0%Stage Report Time: 26-Aug-2024 12:39:17 PM

BRUCE  
(0.05-100)Hz

Ex Time 00:00  
BLC : On  
Notch : On

ExStart  
10.0 mm/mV  
25 mm/Sec.



463415/SWATHI OLADRI  
34 Yrs/Female  
56 Kg/157 Cms  
Date: 26-Aug-2024 12:35:51 PM

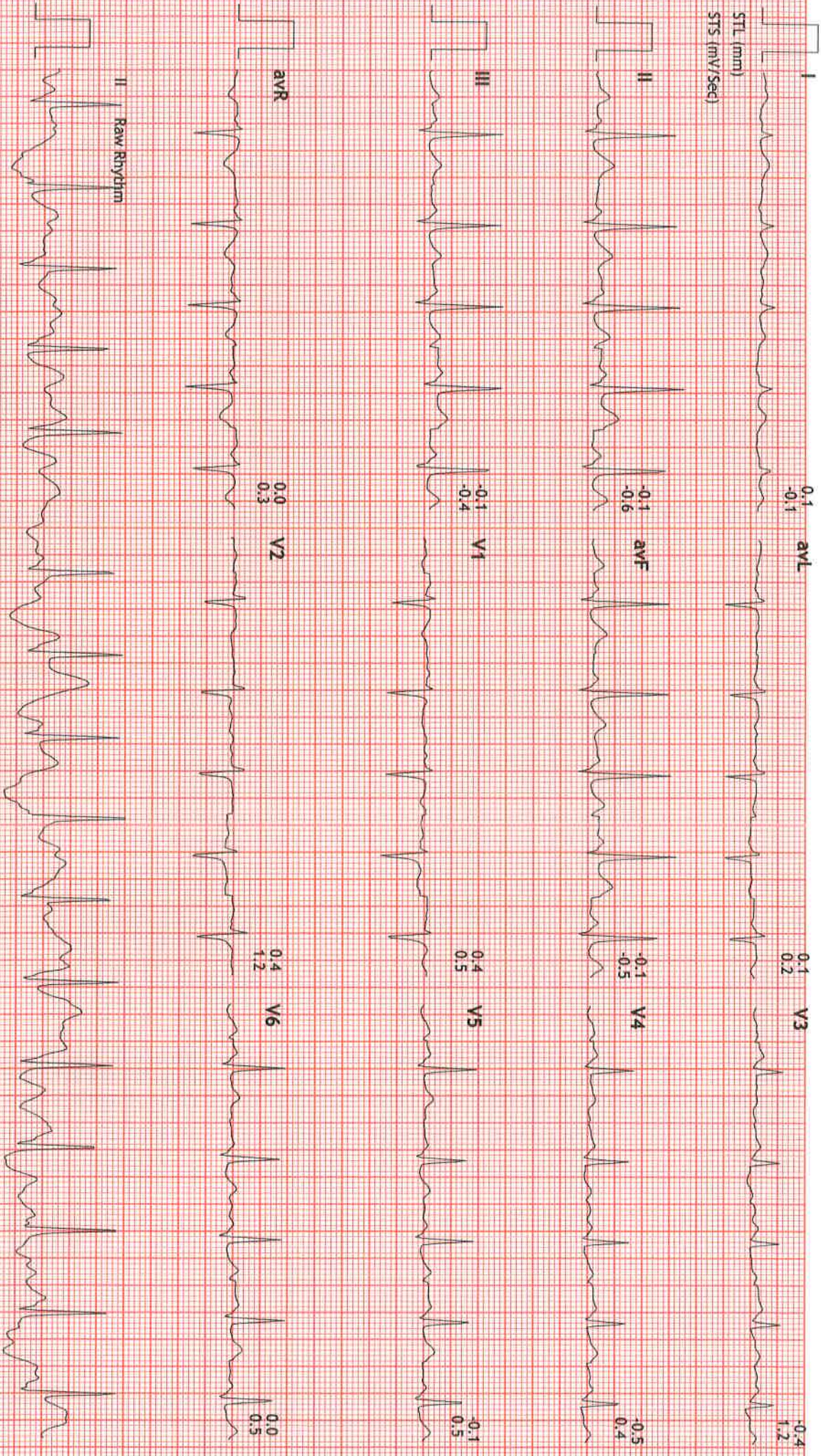
HR: 101 bpm  
METS: 4.6  
BP: ---/---  
Stage Report Time: 26-Aug-2024 12:42:16 PM

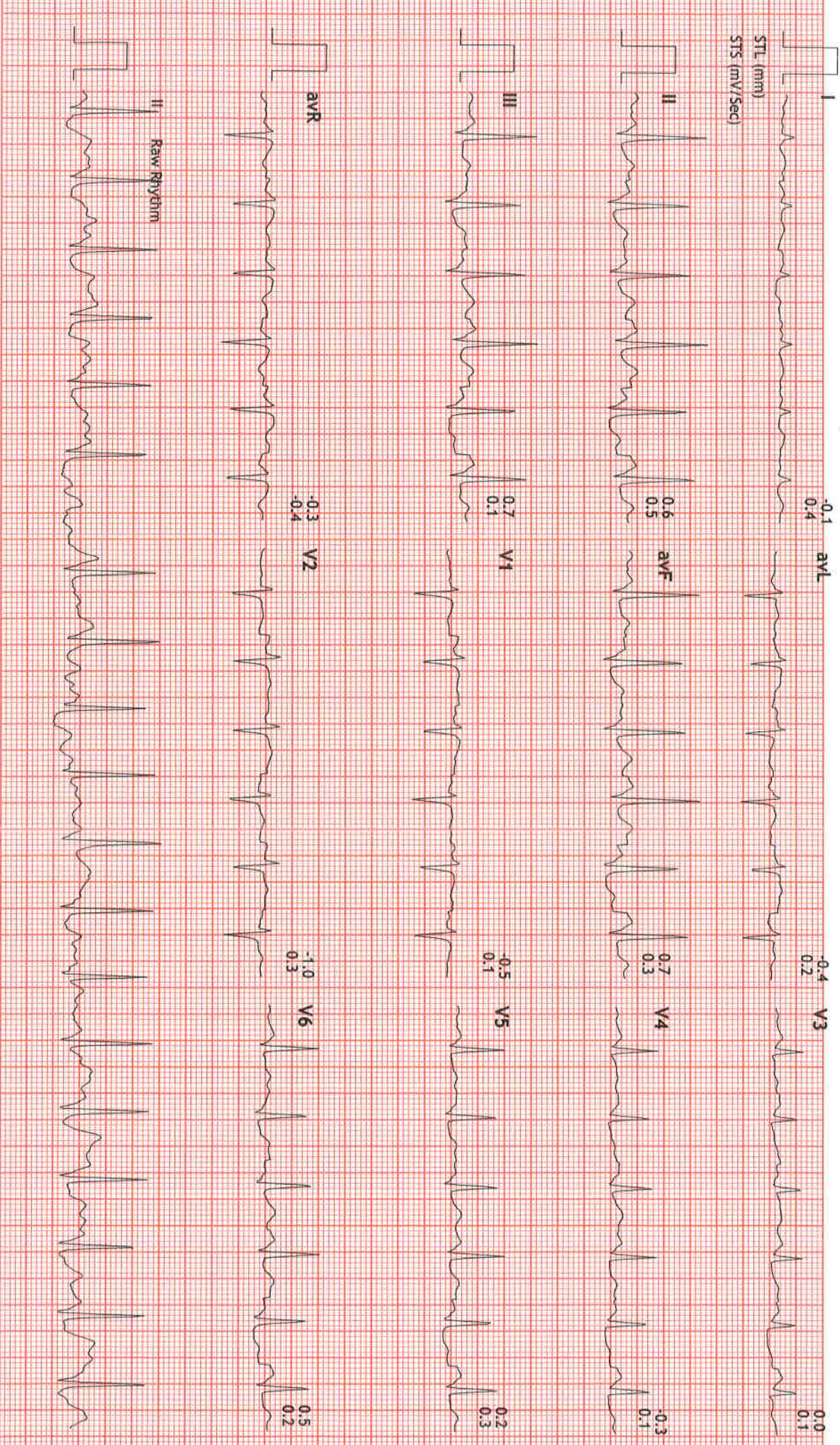
Linked Medians Report

BRUCE  
(0.05-100)Hz

Ex Time 03:00  
BLC :On  
Notch :On

Stage 1 ( 03:00 )  
10.0 mm/mV  
25 mm/Sec.





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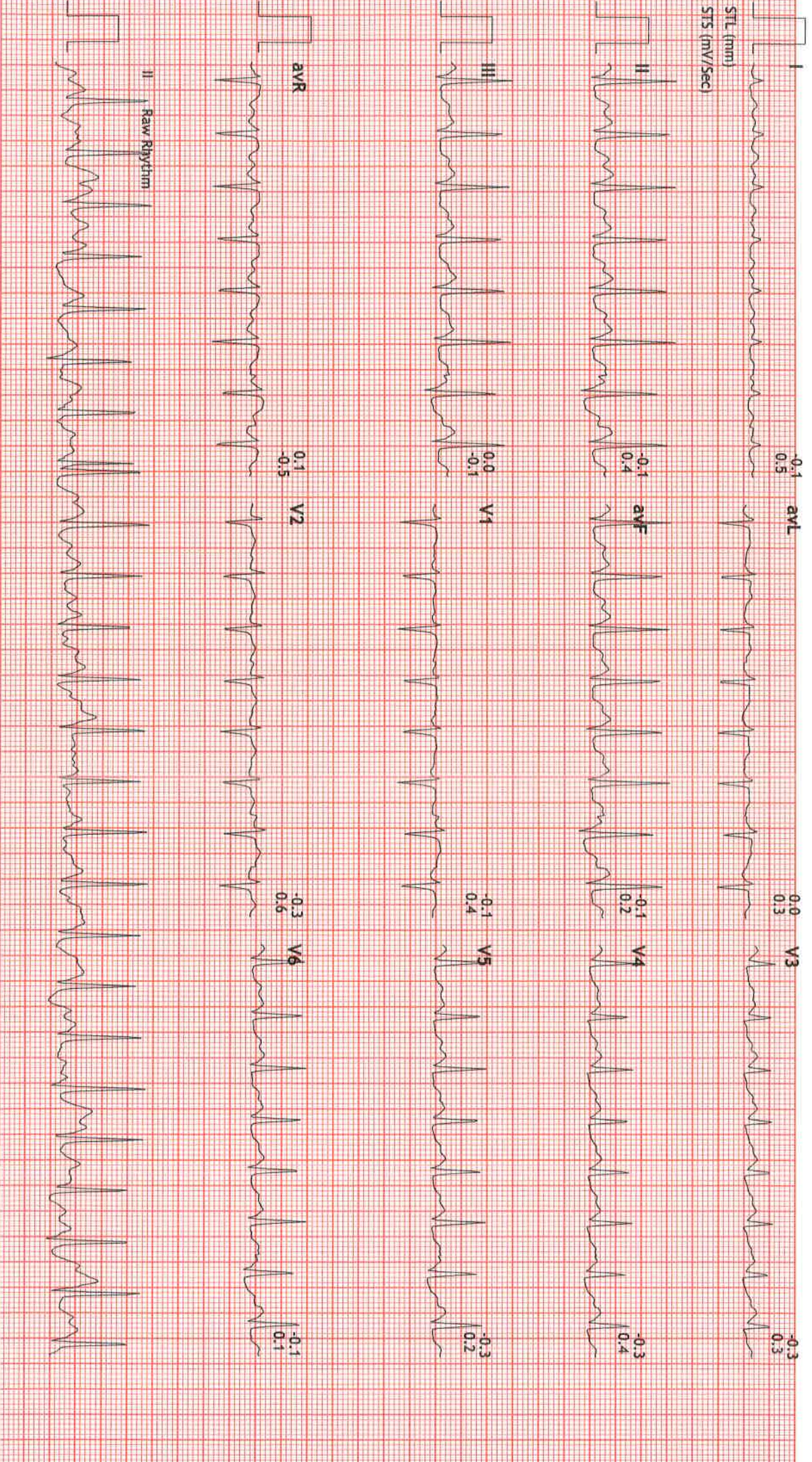
4634115/SWATHI OLADRI  
34 Yrs/Female  
56 Kg/157 Cms  
Date: 26-Aug-2024 12:35:51 PM

HR: 150 bpm  
METS: 10.2  
BP: 140/80  
Stage Report Time: 26-Aug-2024 12:48:17 PM

Linked Medians Report  
BRUCE  
(0.05-100)Hz

Ex Time 09:00  
BLC : On  
Notch : On

Stage 3 ( 03:00 )  
10.0 mm/mV  
25 mm/Sec.

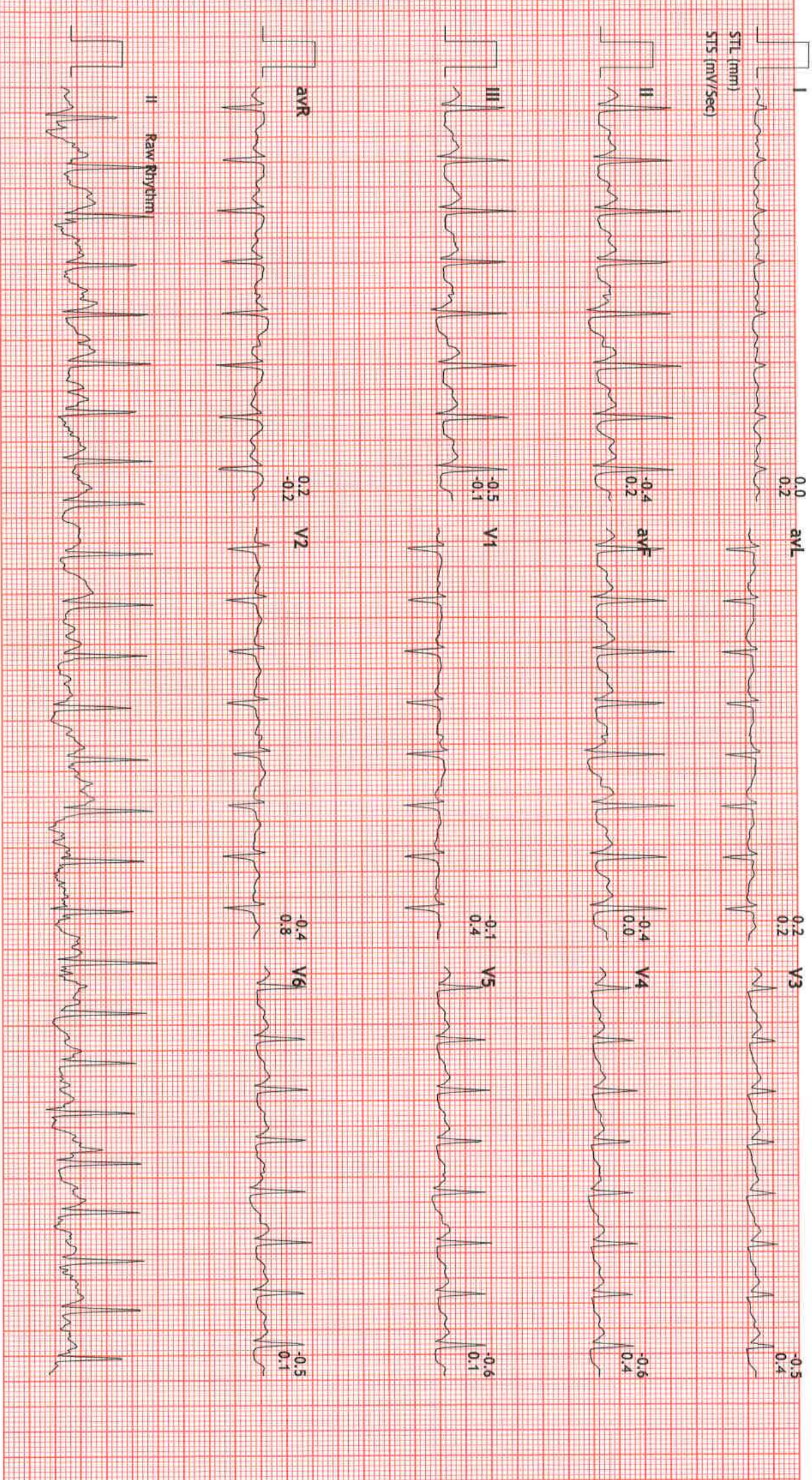


HR: 164 bpm  
METs: 10.6  
BP: 140/80  
Stage Report Time: 26-Aug-2024 12:48:36 PM

MPHR: 88% of 186  
Speed: 6.8 kmph  
Grade: 16.0%

BRUCE  
(0.05-100)Hz  
Ex Time 09:18  
BLC : On  
Notch : On

PeakEx  
10.0 mm/mV  
25 mm/Sec.





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4634115/SWATHI OLADRI  
34 Yrs/Female  
56 Kg/157 Cms  
Date: 26-Aug-2024 12:35:51 PM

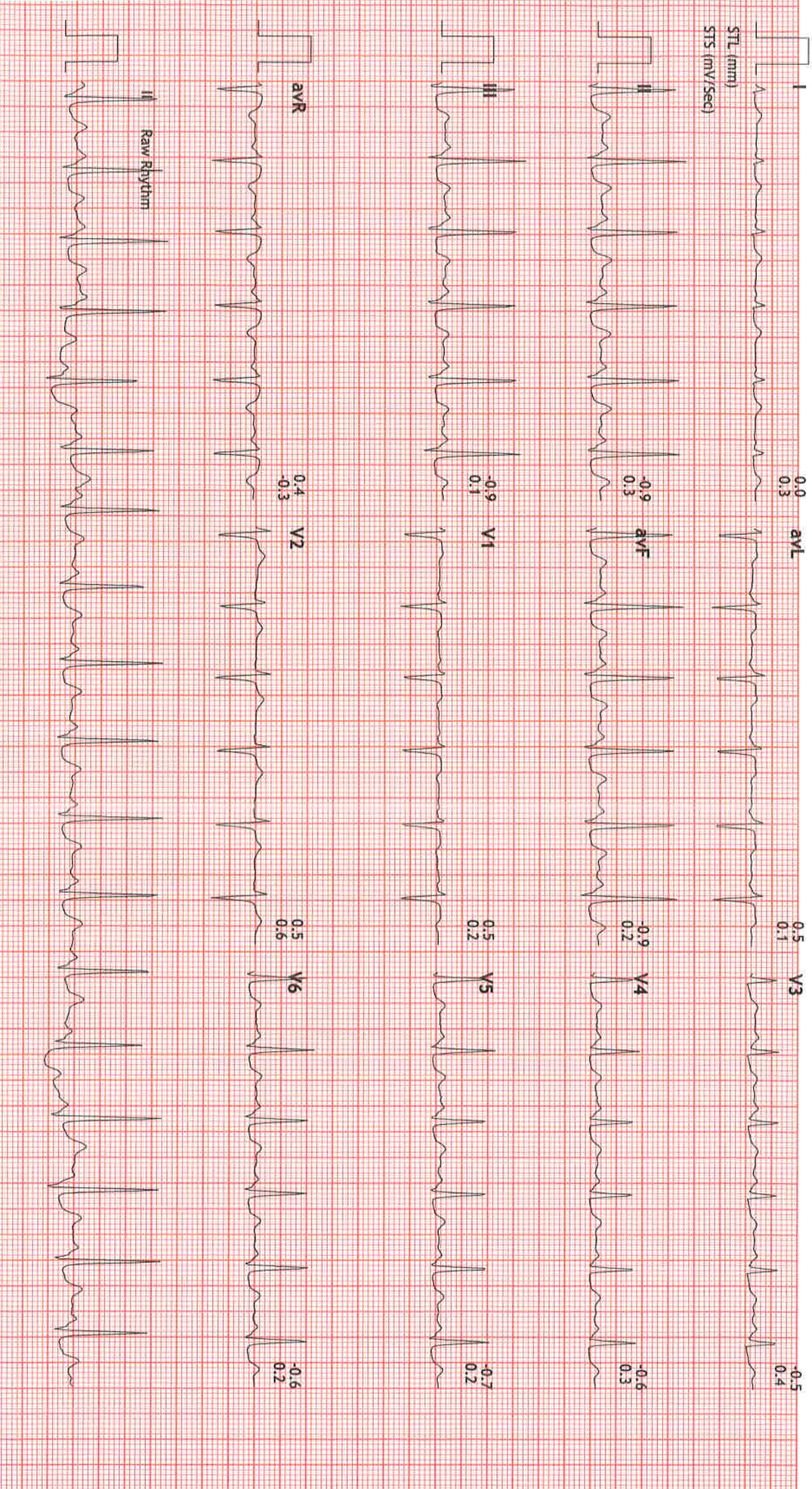
HR: 108 bpm  
METs: 1.0  
BP: 130/80  
Stage Report Time: 26-Aug-2024 12:51:35 PM

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

Linked Medians Report

BRUCE  
(0.05-100)Hz  
Ex Time 09:18  
BLC : On  
Notch : On

Recovery : ( 02:59 )  
10.0 mm/mV  
25 mm/Sec.



34 Years

Female

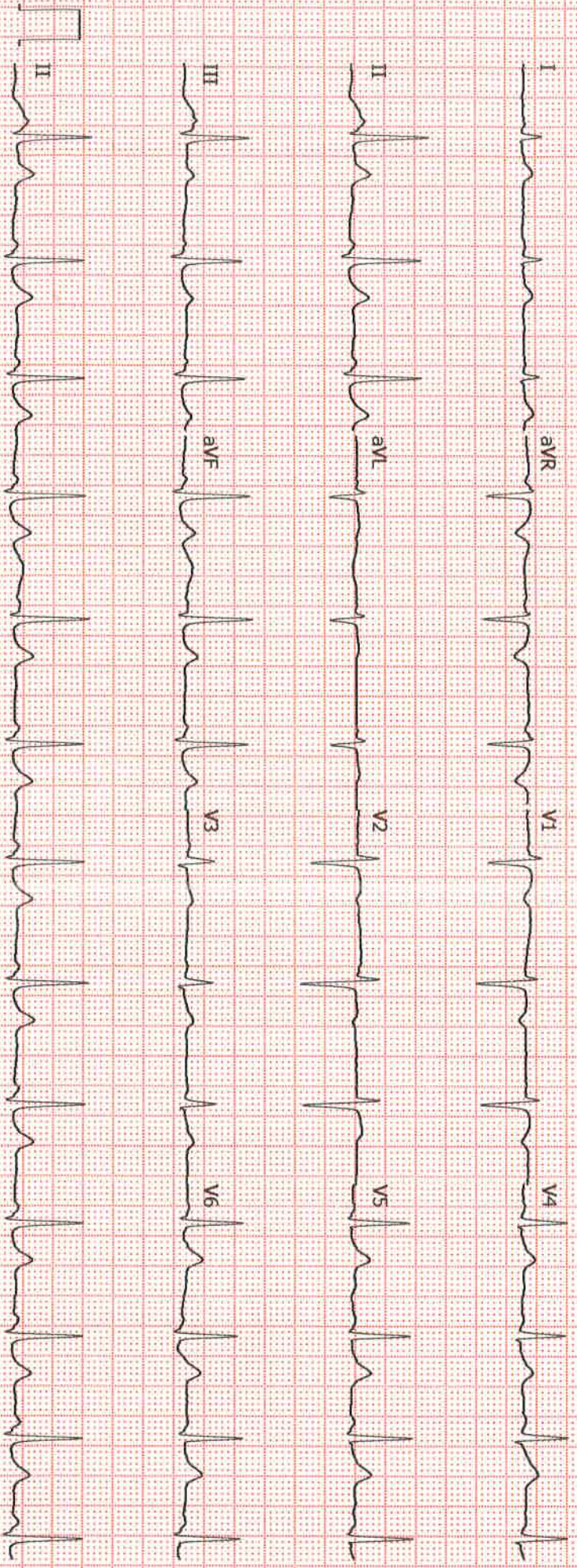
QRS : 84 ms  
 QT / QTcBaz : 376 / 425 ms  
 PR : 100 ms  
 P : 78 ms  
 RR / pp : 780 / 779 ms  
 P / QRS / T : 53 / 81 / 53 degrees

Sinus rhythm with sinus arrhythmia with short PR  
 Otherwise normal ECG

Technician: js  
 Ordering Ph:  
 Referring Ph:  
 Attending Ph:

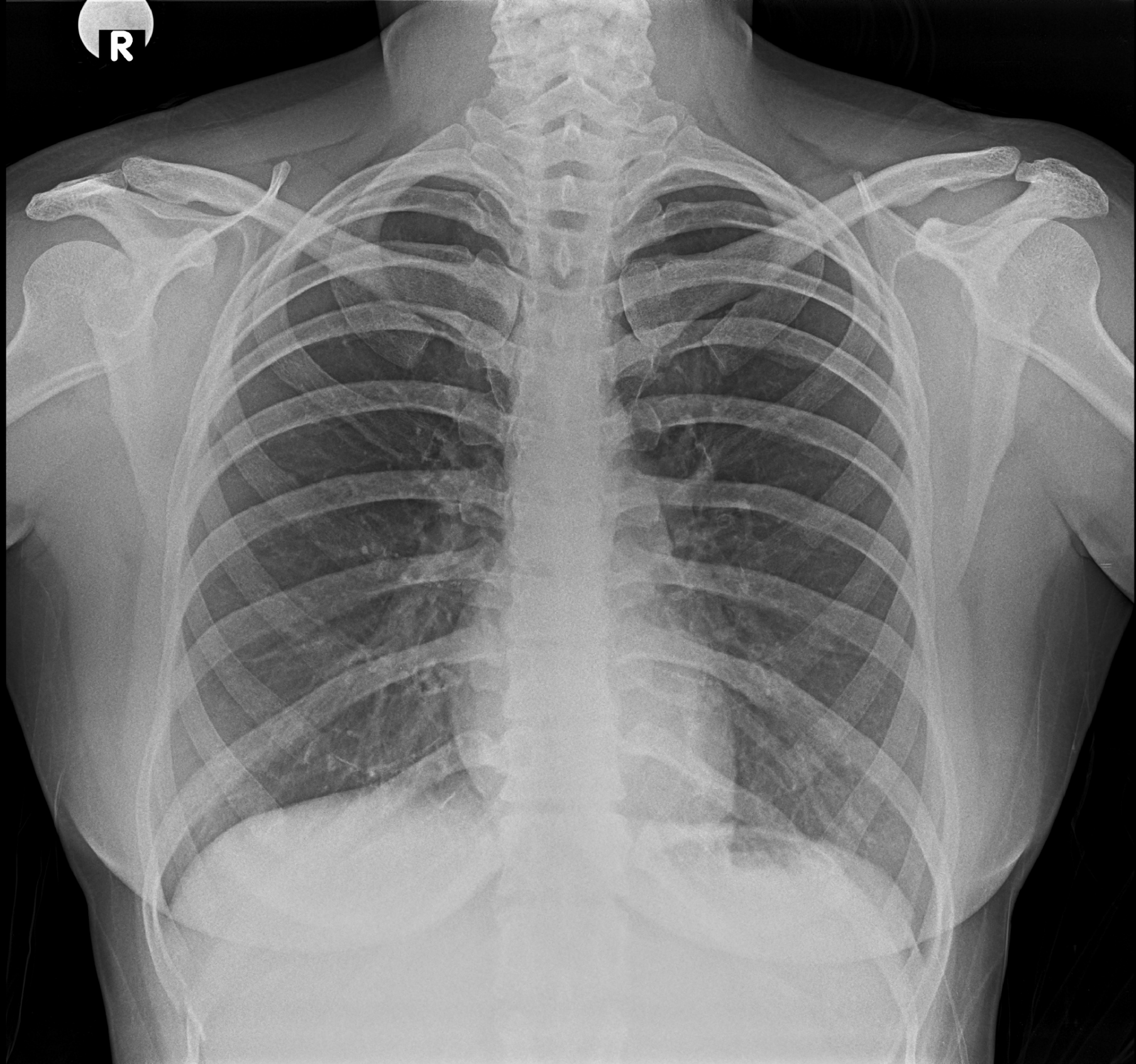
**D/D RAVI KUMAR**  
 MD (S) ENT  
 Registrar  
 Vertas Rehal Hospital  
 Khanapur, Kothapalli Hyderabad-500064 Telangana

T. Anurag Kumar  
 MD (S) ENT  
 Registrar  
 Vertas Rehal Hospital  
 Khanapur, Kothapalli Hyderabad-500064 Telangana



Unconfirmed

R



SWATHI OLADRI BIL4634115 22401258 CHEST PA 8/26/2024  
TENET DIAGNOSTICS GACHIBOWLI HYD