







Name : MRS.MADHAVI N .
Age / Gender : 40 Years / Female

Ref.By : SELF

Reg.No : BIL4816279

TID/SID : UMR2060615/ 28393500 Registered on : 11-Oct-2024 / 08:56 AM Collected on : 11-Oct-2024 / 10:06 AM Reported on : 11-Oct-2024 / 15:07 PM

TEST REPORT Reference : Arcofemi Health Care Ltd -

DEPARTMENT OF CLINICAL PATHOLOGY

Complete Urine Examination (CUE)

Investigation	Result	Biological Reference Intervals
Physical Examination		
Colour	Yellow	Straw to Yellow
Method:Physical		
Appearance	Clear	Clear
Method:Physical		
Chemical Examination		
Reaction and pH Method:Indicator	Acidic (5.5)	4.6-8.0
Specific gravity	1.027	1.000-1.035
Method:Refractometry		
Protein	Negative	Negative
Method:Protein Error of pH indicators		
Glucose	Negative	Negative
Method:Glucose oxidase/Peroxidase		
Blood	Negative	Negative
Method:Peroxidase	Namathia	Negative
Ketones	Negative	Negative
Method:Sodium Nitroprusside	Negative	Negative
Bilirubin Method:Diazonium salt	Negative	rvegative
	Negative	Negative
Leucocytes Method:Esterase reaction	regative	Nogalivo
Nitrites	Negative	Negative
Method:Modified Griess reaction	g	
Urobilinogen	Negative	Up to 1.0 mg/dl
Method:Diazonium salt	-	(Negative)
Microscopic Examination		
Pus cells (leukocytes)	2-3	2 - 3 /hpf
Method:Flow Digital Imaging/Microscopy		
Epithelial cells	2-3	2 - 5 /hpf
Method:Flow Digital Imaging/Microscopy		
RBC (erythrocytes)	Absent	Absent
Method:Flow Digital Imaging/Microscopy		
Casts	Absent	Occasional hyaline casts may be seen
Method:Flow Digital Imaging/Microscopy		







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TEST REPORT Reference : Arcofemi Health Care Ltd -

Crystals

Absent

Phosphate, oxalate, or urate crystals may

be seen

Others

Method:Flow Digital Imaging/Microscopy

Nil

Nil

Method:Flow Digital Imaging/Microscopy

Method: Semi Quantitative test ,For CUE

Reference: Godka**r** 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 infecation 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 Shruti Reddy Consultant Pathologist Reg No.TSMC/FMR/22656









N A R

Name : MRS.MADHAVI N .

Age / Gender : 40 Years / Female

Ref.By : SELF

Reg.No : BIL4816279

TID/SID : UMR2060615/ 28393056 Registered on : 11-Oct-2024 / 08:56 AM

Collected on : 11-Oct-2024 / 09:09 AM

Reported on : 11-Oct-2024 / 13:38 PM

TEST REPORT Reference : Arcofemi Health Care Ltd -

DEPARTMENT OF HEMATOPATHOLOGY

Blood Grouping ABO And Rh Typing, EDTA Whole Blood

Parameter Results

Blood Grouping (ABO)

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 expresses 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







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Reg.No : BIL4816279

TID/SID : UMR2060615/ 28393056 Registered on : 11-Oct-2024 / 08:56 AM Collected on : 11-Oct-2024 / 09:09 AM

Reported on : 11-Oct-2024 / 12:53 PM

TEST REPORT Reference : Arcofemi Health Care Ltd -

DEPARTMENT OF HEMATOPATHOLOGY

Erythrocyte Sedimentation Rate (ESR), Whole Blood

Investigation	Observed Value	Biological Reference Intervals
ESR 1st Hour	30	<=12 mm/hour

Method:Westergren/Vesmatic

Complete Blood Count (CBC), EDTA Whole Blood

Investigation	Observed Value	Biological Reference Intervals
Hemoglobin	13.7	12.0-15.0 g/dL
Method:Cyanide Free Lyse Hemoglobin		
PCV/HCT	40.1	36.0-46.0 vol%
Method:Calculated		
Total RBC Count	4.90	3.80-4.80 mill /cu.mm
Method:Electrical Impedance		
MCV	81.9	83.0-101.0 fL
Method:Calculated		
MCH	27.9	27.0-32.0 pg
Method:Calculated		
MCHC	34.1	31.5-34.5 g/dL
Method:Calculated		
RDW (CV)	15.6	11.6-14.0 %
Method:Calculated		
MPV	7.5	7.0-10.0 fL
Method:Calculated		
Total WBC Count	5660	4000-10000 cells/cumm
Method:Electrical Impedance		
Platelet Count	3.02	1.50-4.10 lakhs/cumm
Method:Electrical Impedance		
Differential count		
Neutrophils	50.8	40.0-80.0 %
Method:Microscopy		
Lymphocytes	40.9	20.0-40.0 %
Method:Microscopy		
Eosinophils	2.2	1.0-6.0 %
Monocytes	5.8	2.0-10.0 %
Basophils	0.3	< 1.0-2.0 %
Method:Microscopy		







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TEST REPORT Reference : Arcofemi Health Care Ltd -

	0075	0000 7000 # /
Absolute Neutrophil Count	2875	2000-7000 cells/cumm
Method:Calculated		
Absolute Lymphocyte Count (ALC)	2315	1000-3000 cells/cumm
Absolute Eosinophil Count (AEC)	125	20-500 cells/cumm
Absolute Monocyte Count	328	200-1000 cells/cumm
Method:Calculated		
Absolute Basophil Count	17	20-100 cells/cumm
Method:Calculated		
Neutrophil - Lymphocyte Ratio(NLR)	1.24	0.78-3.53
Method:Calculated		
monioa.ouioaiatoa		

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
Age / Gender

Method:Urease/UV

: MRS.MADHAVI N .

: 40 Years / Female

Ref.By : SELF

Req.No : BIL4816279

TID/SID : UMR2060615/ 28393058F Registered on : 11-Oct-2024 / 08:56 AM Collected on : 11-Oct-2024 / 09:09 AM

Reported on : 11-Oct-2024 / 12:58 PM

TEST REPORT Reference : Arcofemi Health Care Ltd -

Blood Urea Nitrogen (BUN), Serum Investigation Observed Value Biological Reference Interval Blood Urea Nitrogen. 9 6-20 mg/dL Method:Calculated Urea. 18.9 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.	0.70	0.50-0.90 mg/dL
Method:Alkaline Picrate		

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

	<u> </u>	
Investigation	Observed Value	Biological Reference Interval
Glucose Fasting Method:Hexokinase	83	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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Ref.By : SELF Collected on : 11-Oct-2024 / 09:10 AM

Req.No : BIL4816279 Reported on : 11-Oct-2024 / 13:37 PM

TEST REPORT Reference : Arcofemi Health Care Ltd -

Glucose Post Prandial (PPBS), Sodium Fluoride Plasma

Investigation	Observed Value	Biological Reference Interval
Glucose Post Prandial Method:Hexokinase	71	Normal : <140 mg/dL Impaired PG: 140-199 mg/dL Diabetes mellitus: >/=200 mg/dL
Note	The discordant post prandial blood glucose values levels are observed in some of the conditions related to defective absorption, insufficient dietary intake, endocrine disorders, hypoglycemic drug overdose and reactive hypoglycemia etc.	

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	4.9	Non-diabetic: <= 5.6 % Pre-diabetic: 5.7 - 6.4 % Diabetic: >= 6.5 %
Estimated Average Glucose (eAG)	94	mg/dL
Method:Calculated		

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 glycated 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 estimation:
- 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 glycemic 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

Ban/orcatiline ratio, cerain			
Investigation	Observed Value		
BUN/Creatinine Ratio Method:Calculated	11	10-20	





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TEST REPORT Reference : Arcofemi Health Care Ltd -

TID/SID

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 ---







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Reported on . 11-Oct-2024 / 13:37

TEST REPORT Reference : Arcofemi Health Care Ltd -

TID/SID

DEPARTMENT OF CLINICAL CHEMISTRY I

Lipid Profile, Serum

Lipia i romo, coram		
Investigation	Observed Value	Biological Reference Interval
Total Cholesterol Method:Cholesterol Oxidase	183	Desirable: <200 mg/dL Borderline: 200-239 mg/dL High: >/=240 mg/dL
HDL Cholesterol Method:Direct Measurement	28	Low: <40 mg/dL High: >/=60 mg/dL
VLDL Cholesterol Method:Calculated	15	6.0-38.0 mg/dL
LDL Cholesterol Method:Calculated	140	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	75	Normal:<150 mg/dL Borderline: 150-199 mg/dL High: 200-499 mg/dL Very high: >/=500 mg/dL
Chol/HDL Ratio Method:Calculated	6.54	Low Risk: 3.3-4.4 Average Risk: 4.5-7.1 Moderate Risk: 7.2-11.0
LDL Cholesterol/HDL Ratio Method:Calculated	5	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 ---







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TEST REPORT Reference : Arcofemi Health Care Ltd -

DEPARTMENT OF CLINICAL CHEMISTRY I

Liver Function Test (LFT), Serum

Investigation	Observed Value	Biological Reference Interval
Total Bilirubin. Method:Diazo method	0.58	<1.2 mg/dL
Direct Bilirubin. Method:Diazo method	0.29	<0.30 mg/dL
Indirect Bilirubin. Method:Calculated	0.29	<0.9 mg/dL
Alanine Aminotransferase ,(ALT/SGPT) Method:UV wtihout P5P	24	<34 U/L
Aspartate Aminotransferase,(AST/SGOT) Method:UV wtihout P5P	20	<31 U/L
ALP (Alkaline Phosphatase). Method:PNPP-AMP Buffer	91	35-104 U/L
Gamma GT. Method:Gamma-Glutamyl - 3 - Carbossi - 4 - Nitroanilide	115	6-42 U/L
(GCNA)	7.6	6.6-8.7 g/dL
Total Protein. Method:Biuret		-
Albumin. Method:Bromocresol Green (BCG)	4.3	3.5-5.2 g/dL
Globulin.	3.30	1.8-3.8 g/dL
Method:Calculated A/GRatio. Method:Calculated	1.30	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.

--- End Of Report ---

^{*} Sample processed at National Reference Laboratory, Tenet Diagnostics, Hyderabad







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TEST REPORT Reference : Arcofemi Health Care Ltd -

DEPARTMENT OF CLINICAL CHEMISTRY I

Thyroid Profile (T3,T4,TSH), Serum

Triyroid Frome (15,14,1511), Serum		
Investigation	Observed Value	Biological Reference Interval
Triiodothyronine Total (T3) Method:ECLIA	1.14	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	10.1	5.1-14.1 μg/dL
Thyroid Stimulating Hormone (TSH) Method:ECLIA	1.06	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 Clinial Chemistry and Molecular Diagnostics, Nader Rifia, Andrea Ritas Horvath, Carl T. Wittwer.

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

--- End Of Report ---







TO VERIFY THE REPORT ONLINE

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: 40 Years / Female

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:UMR2060615/ 28393057 Registered on: 11-Oct-2024 / 08:56 AM

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Reference : Arcofemi Health Care Ltd -

DEPARTMENT OF CLINICAL CHEMISTRY I					
Uric Acid, Serum					
Observed Value	Biological Reference Interval				
3.1	2.4-5.7 mg/dL				

TID/SID

Interpretation

Method:Uricase

Investigation

Uric Acid.

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, preeclampsia, 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.

TEST REPORT

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

--- End Of Report ---







PLEASE SCAN OR CODE

Name : Mrs. MADHAVI N. TID : UMR2060615

Age/Gender: 40 Years/FemaleRegistered On: 11-Oct-2024 08:56 AMRef By: SelfReported On: 11-Oct-2024 02:35 PMReg.No: BIL4816279Reference: Arcofemi Health Care Ltd

- Medi Whe

Mammography Bilateral

Mediolateral oblique and craniocaudal views was performed.

BILATERAL MAMMOGRAPHY:

Bilateral breasts show symmetrical fibroglandular parenchyma.

Few medium dense lesions are noted at the upper inner quadrant of the left breast, few of the lesions are showing punctate calcifications

No evidence of focal soft tissue lesion in right breast.

No evidence of cluster microcalcification.

Subcutaneous fat deposition is within normal limits.

IMPRESSION:

* Few medium dense lesions at the upper inner quadrant of the left breast, few of the lesions are showing punctate calcifications - Likely Fibroadenoma

BI-RADS CATEGORY - 2

BI-RADS CLASSIFICATION CATEGORY RESULT

AIEGURT	RESULT
0	Assessment incomplete. Need additional imaging evaluation
1	Negative. Routine mammogram in 1 year recommended.
2	Benign finding. Routine mammogram in 1 year recommended.
3	Probably benign finding. Short interval follow-up suggested.
4	Suspicious - 4A : Low suspicion for malignancy (2 - 9%)
	4B: Moderate suspicion for malignancy (10 - 49%)
	4C : High suspicion for malignancy (50 - 94%)
	Biopsy should be considered.
5	Highly suggestive of malignancy. Appropriate action should be taken.
6.	Known biopsy proven malignancy.

Suggested clinical correlation and follow up

*** End Of Report ***

B. KOWA BELIDE
Consultant Radiologist



: BIL4816279

Reg.No



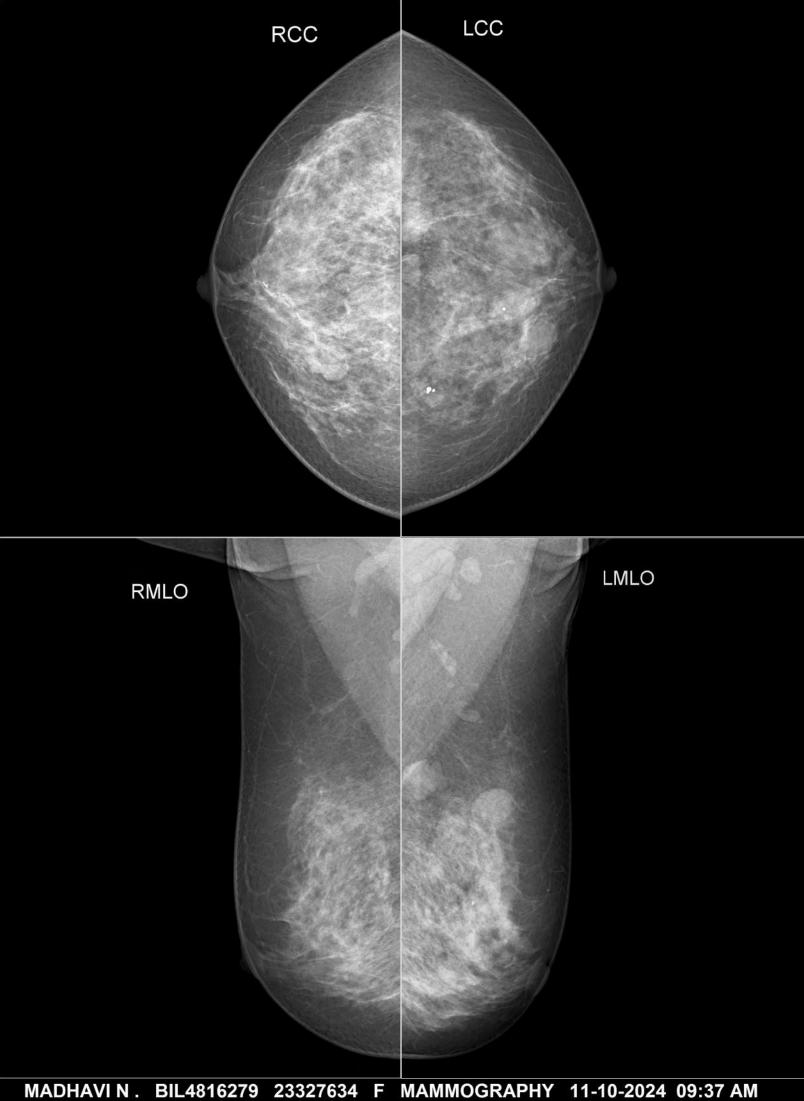
PLEASE SCAN QR CODE

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Reference : Arcofemi Health Care Ltd

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MADHAVI N. BIL4816279 23327634 F MAMMOGRAPHY 11-10-2024 09:37 AM TENET DIAGNOSTICS, VIKARAMPURI, SECUNDERABAD

RMS

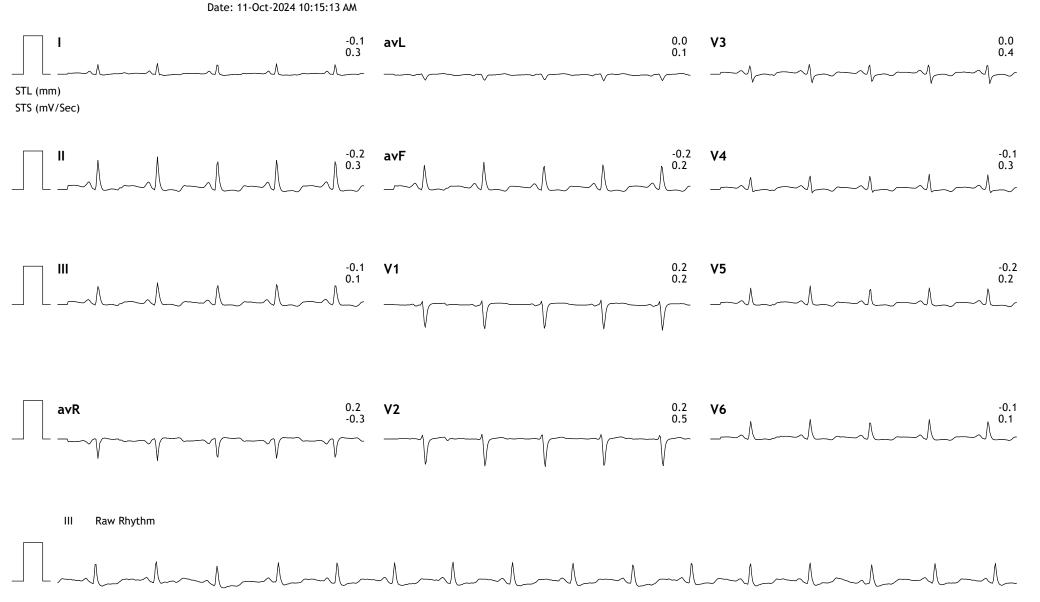
Linked Medians Report

TENET DIAGNOSTIC VIKRAMPURI, HYDERABAD

4816279/MRS N MADHAV**HR: 97 bpm** 40 Yrs/Female METS: 1.0

0 Kg/0 Cms BP: 110/70

MPHR:53% of 180 BRUCE Speed: 0.0 mph (1.0-100)Hz Grade: 0.0% Ex Time 00:51 BLC :On Notch :On SUPINE 10.0 mm/mV 25 mm/Sec.

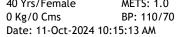


RMS

Linked Medians Report

TENET DIAGNOSTIC VIKRAMPURI, HYDERABAD

4816279/MRS N MADHA**VHR: 101 bpm** 40 Yrs/Female METS: 1.0 MPHR:56% of 180 BRUCE Speed: 0.0 mph (1.0-100)Hz Grade: 0.0% Ex Time 00:52 BLC :On Notch :On STANDING 10.0 mm/mV 25 mm/Sec.















Linked Medians Report

TENET DIAGNOSTIC VIKRAMPURI, HYDERABAD BRUCE

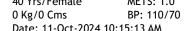
(1.0-100)Hz

4816279/MRS N MADHAVHR: 95 bpm 40 Yrs/Female METS: 1.0

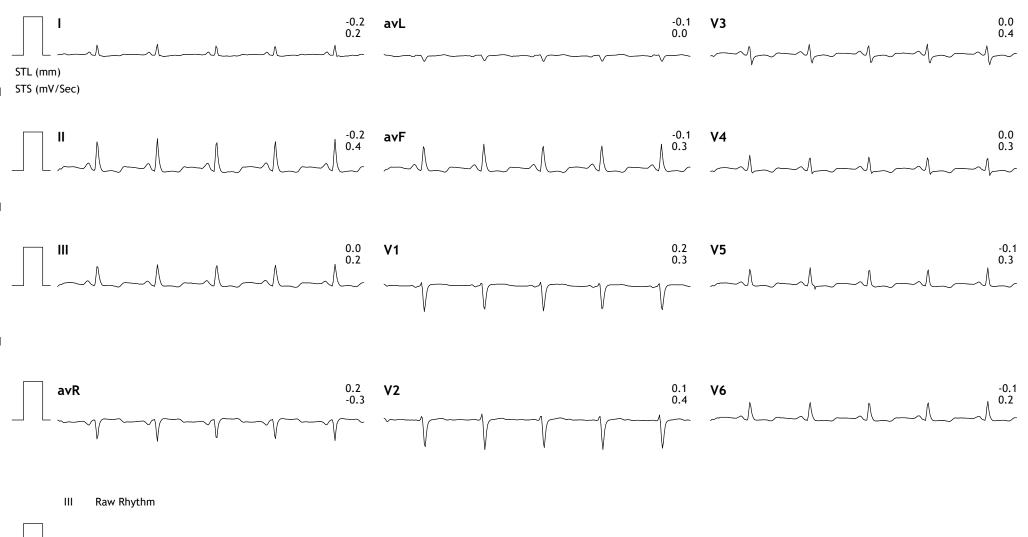
MPHR:52% of 180 Speed: 0.0 mph Grade: 0.0%

Ex Time 00:00 BLC:On Notch:On

ExStrt 10.0 mm/mV 25 mm/Sec.



Date: 11-Oct-2024 10:15:13 AM



RMS

Linked Medians Report

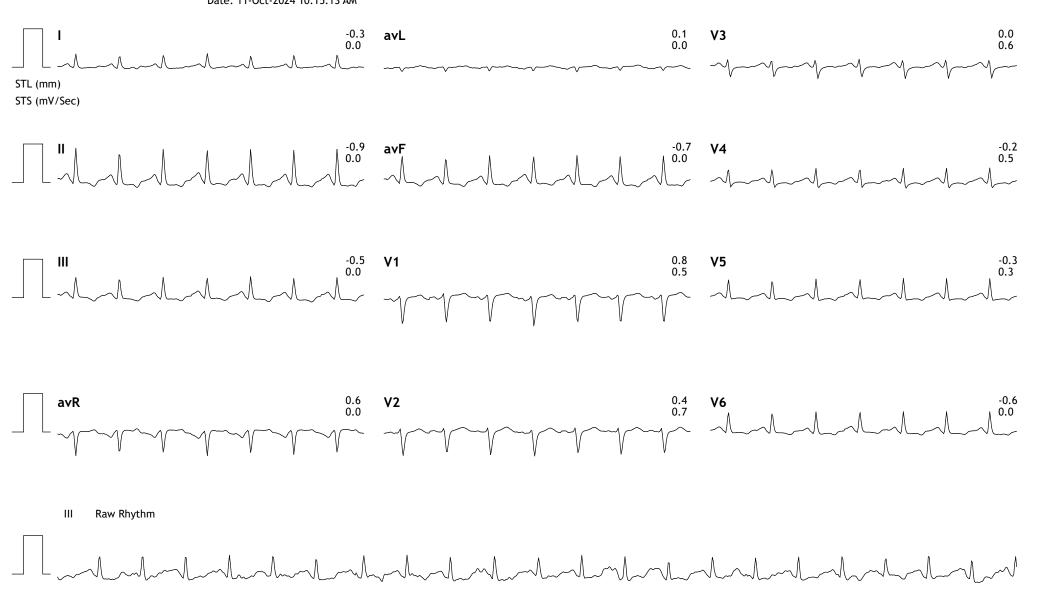
TENET DIAGNOSTIC VIKRAMPURI, HYDERABAD

4816279/MRS N MADHAVHR: 133 bpm 40 Yrs/Female METS: 4.7

0 Kg/0 Cms BP: 120/70 Date: 11-Oct-2024 10:15:13 AM

MPHR:73% of 180 BRUCE Speed: 1.7 mph (1.0-10

Speed: 1.7 mph (1.0-100)Hz Grade: 10.0% Ex Time 03:00 BLC :On Notch :On Stage 1 (03:00) 10.0 mm/mV 25 mm/Sec.



RMS

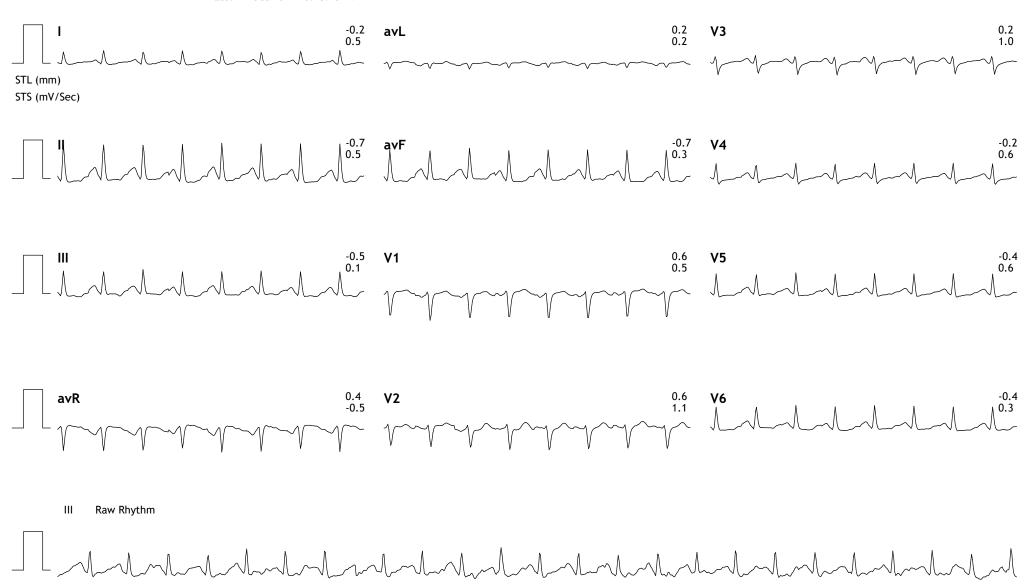
Linked Medians Report

TENET DIAGNOSTIC VIKRAMPURI, HYDERABAD

4816279/MRS N MADHAVHR: 147 bpm 40 Yrs/Female METS: 7.1

0 Kg/0 Cms BP: 130/70 Date: 11-Oct-2024 10:15:13 AM

MPHR:81% of 180 BRUCE Speed: 2.5 mph (1.0-100)Hz Grade: 12.0% Ex Time 06:00 BLC :On Notch :On Stage 2 (03:00) 10.0 mm/mV 25 mm/Sec.



Linked Medians Report

TENET DIAGNOSTIC VIKRAMPURI, HYDERABAD

4816279/MRS N MADHAVHR: 156 bpm 40 Yrs/Female METS: 7.4 0 Kg/0 Cms BP: 130/70

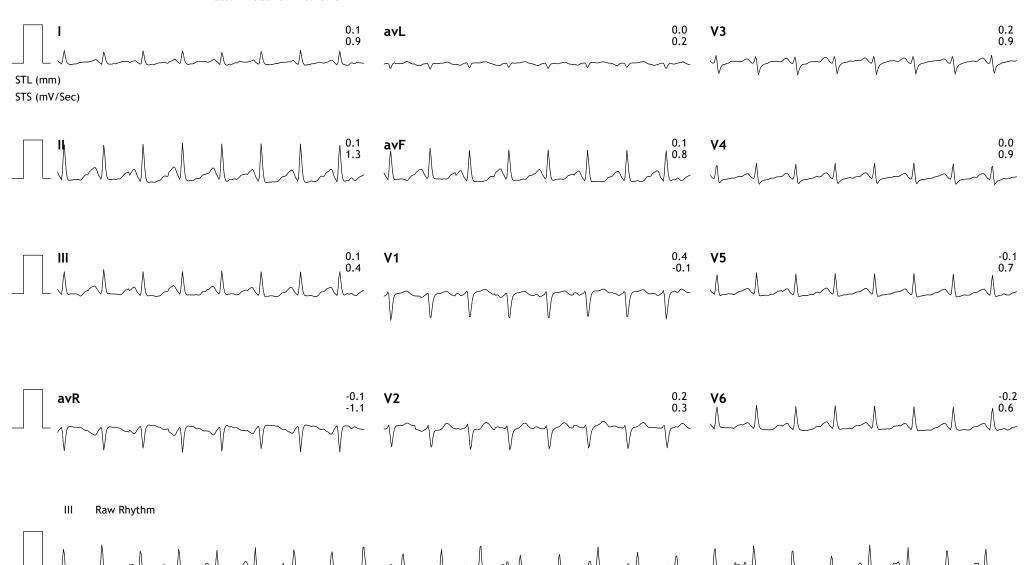
Date: 11-Oct-2024 10:15:13 AM

BRUCE MPHR:86% of 180 Speed: 3.4 mph (1.0-100)Hz

Grade: 14.0%

Ex Time 06:15 BLC:On Notch:On

PeakEx 10.0 mm/mV 25 mm/Sec.



Linked Medians Report

TENET DIAGNOSTIC VIKRAMPURI, HYDERABAD

4816279/MRS N MADHAVHR: 131 bpm 40 Yrs/Female METS: 1.1

0 Kg/0 Cms Date: 11-Oct-2024 10:15:13 AM

BRUCE MPHR:72% of 180 Speed: 0.0 mph (1.0-100)Hz Ex Time 06:15 BLC:On Notch:On

Recovery: (00:59) 10.0 mm/mV

25 mm/Sec.

BP: 130/70 Grade: 0.0%

avL

0.0 ٧3

0.0 0.1 STL (mm) STS (mV/Sec)

0.4 0.5





Ш Raw Rhythm

RMS

Linked Medians Report

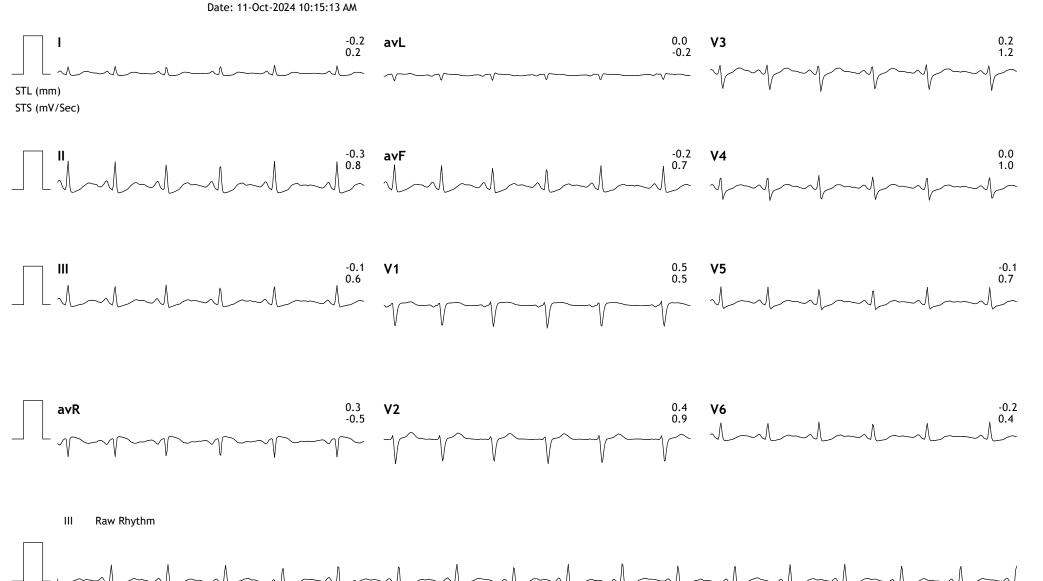
0 Kg/0 Cms

TENET DIAGNOSTIC VIKRAMPURI, HYDERABAD

4816279/MRS N MADHAVHR: 103 bpm 40 Yrs/Female METS: 1.0

METS: 1.0 Speed: 0.0 mph BP: 120/70 Grade: 0.0% BRUCE (1.0-100)Hz Ex Time 06:15 BLC :On Notch :On Recovery: (01:59)

10.0 mm/mV 25 mm/Sec.



MPHR:57% of 180

Summary



TENET DIAGNOSTIC VIKRAMPURI, HYDERABAD

4816279/MRS N MADHAVI 40 Yrs/Female 0 Kg/0 Cms

Date: 11-Oct-2024 10:15:13 AM

Protocol : BRUCE

Print Date: 11-Oct-2024

History:

Ref.By : Medication : Objective :

Stage	StageTime (Min:Sec)	PhaseTime (Min:Sec)	Speed (mph)	Grade (%)	METs	H.R.	B.P. (mmHg)	R.P.P. ×100	PVC	Comments
Supine					1.0	97	110/70	106	-	
Standing					1.0	97	110/70	106	-	
ExStart					1.0	96	110/70	105	-	
Stage 1	3:00	3:01	1.7	10.0	4.7	133	120/70	159	-	
Stage 2	3:00	6:01	2.5	12.0	7.1	147	130/70	191	-	
PeakEx	0:15	6:16	3.4	14.0	7.4	156	130/70	202	-	
Recovery	1:00		0.0	0.0	1.2	131	130/70	170	-	
Recovery	2:00		0.0	0.0	1.0	103	120/70	123	-	
Recovery	2:05		0.0	0.0	1.0	100	110/70	110	-	

Findings:

Exercise Time : 06:16

Max HR attained $\,$: 156 bpm $\,$ 87% of Max Predictable HR 180 $\,$

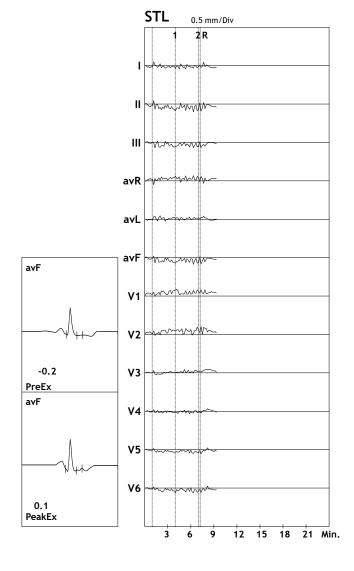
Max BP: 130/70(mmHg)
WorkLoad attained: 7.4 (

No significant ST segment changes noted during exercise or recovery.

No Angina/Arrhythmia/S3/murmur

Final Impression

Maxmum Depression: --:--



Advice/Comments:





PLEASE SCAN OR CODE

Name: Mrs. MADHAVI N. TID: UMR2060615

Age/Gender: 40 Years/FemaleRegistered On: 11-Oct-2024 08:56 AMRef By: SelfReported On: 11-Oct-2024 12:54 PMReg.No: BIL4816279Reference: Arcofemi Health Care Ltd

- Medi Whe

Ultrasound Whole Abdomen

LIVER is normal shape, size (12.7 cms) and has uniform echopattern. No evidence of focal lesion. No intrahepatic biliary ductal dilatation. Hepatic and portal vein radicals are normal.

GALL BLADDER shows normal shape and has clear contents. Gall bladder wall is of normal thickness. CBD is of normal calibre.

PANCREAS has normal shape, size and uniform echopattern. No evidence of ductal dilatation or calcification.

SPLEEN shows normal shape, size and echopattern.

KIDNEYS move well with respiration and have normal shape, size and echopattern. Cortico-medullary differentiations are well madeout.

No evidence of calculus. No hydronephrosis in right kidney.

Prominent left renal pelvis, 16 mm in AP diameter.

Right kidney measures 9.7 x 4.5 cms, Left kidney measures 10.1 x 4.5 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 10.6 mm.

Uterus measures 8.4 x 4.7 x 5.7 cms.

OVARIES are normal in size, shape and echotexture.

Right ovary: $2.7 \times 2.0 \text{ cms}$, Left ovary: $2.7 \times 2.0 \text{ cms}$.

No evidence of free fluid in the abdomen and pelvis.

IMPRESSION:

* Prominent left renal pelvis.

Suggested clinical correlation and follow up

*** End Of Report ***

Dr.Abid yazden Consultant Radiologist





DI EASE SCAN OR CODE

Name : Mrs. MADHAVI N. TID : UMR2060615

Age/Gender : 40 Years/Female Registered On : 11-Oct-2024 08:56 AM

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- Medi Whe



: BIL4816279



PLEASE SCAN OR CODE

Name: Mrs. MADHAVI N. TID: UMR2060615

Age/Gender: 40 Years/FemaleRegistered On: 11-Oct-2024 08:56 AMRef By: SelfReported On: 11-Oct-2024 01:01 PM

Reference : Arcofemi Health Care Ltd

- Medi Whe

DEPARTMENT OF X-RAY X-Ray Chest PA View

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:

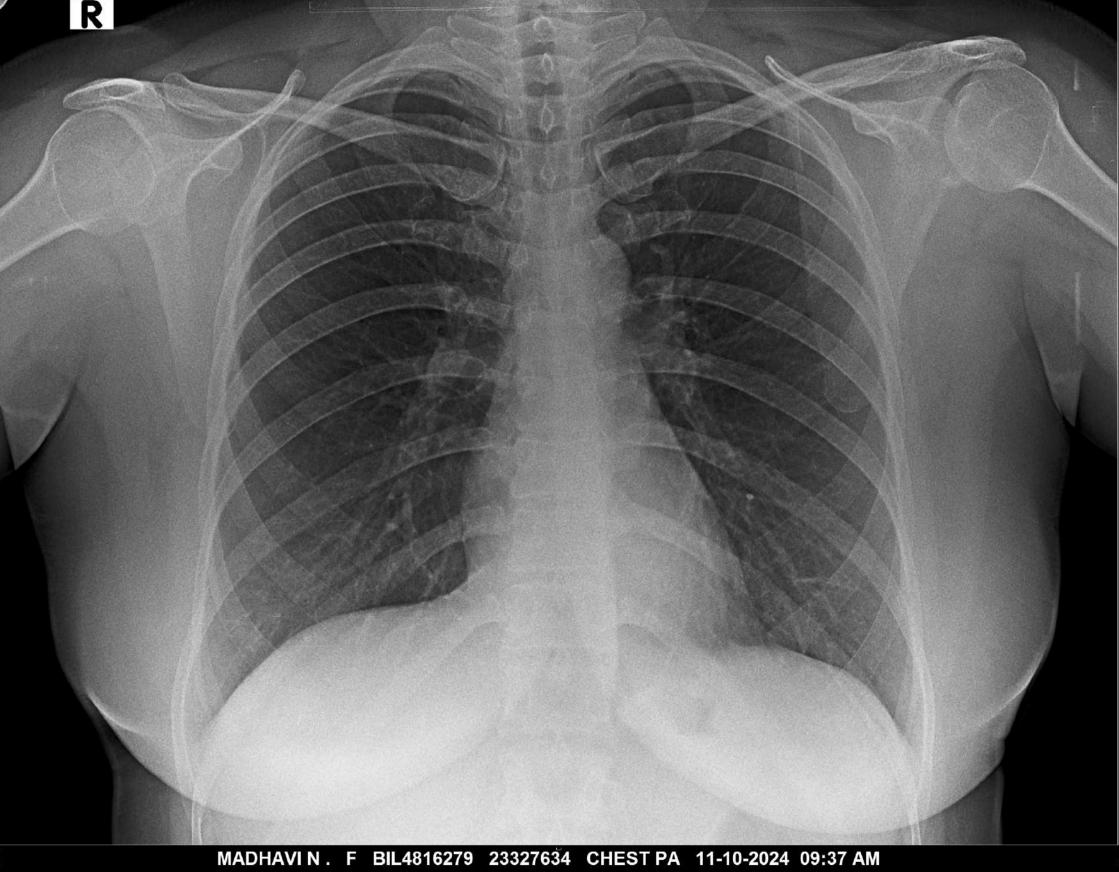
Reg.No

* Normal study.

Suggested clinical correlation and follow up.

*** End Of Report ***

Dr Revanth MD DNB



MADHAVIN. F BIL4816279 23327634 CHEST PA 11-10-2024 09:37 AM
TENET DIAGNOSTICS, VIKARAMPURI, SECUNDERABAD