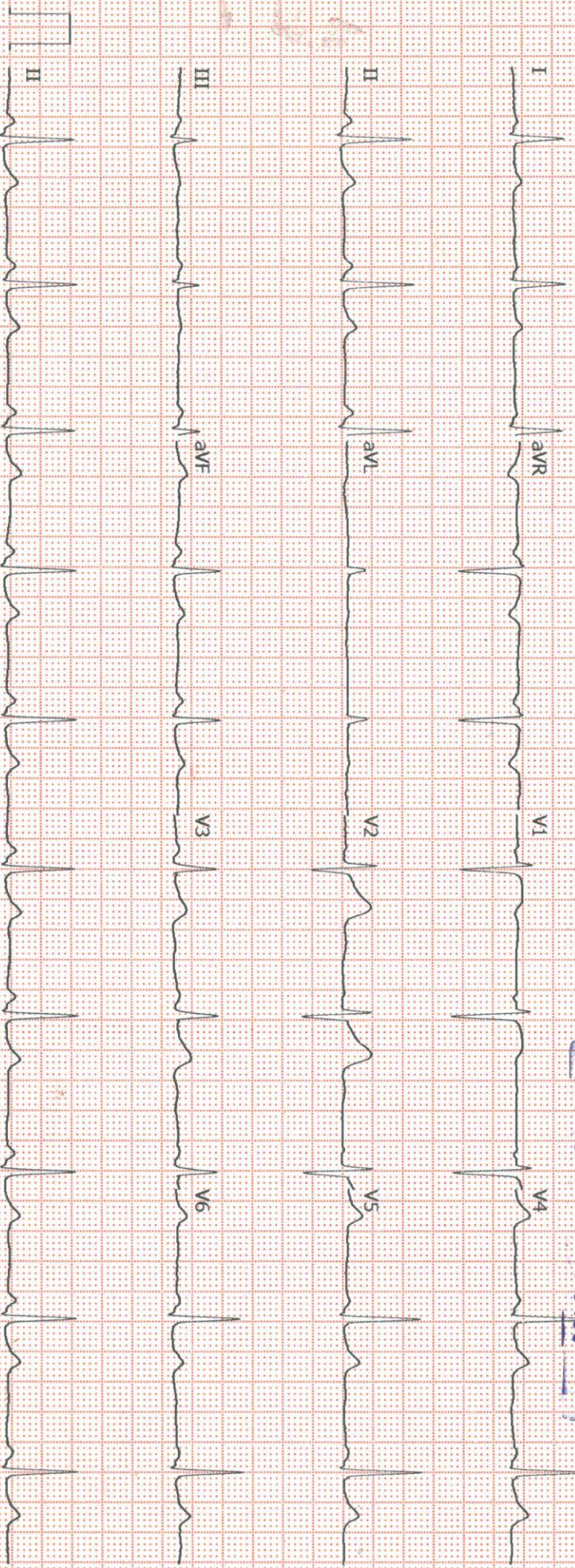


Female  
48 Years

10.08.2024 11:48:20  
TENET DIAGNOSTICS  
VIKRAMPURTI HUB  
HYDERABAD

Normal sinus rhythm  
Normal ECG

QRS	84 ms
QT / QTcbaz	408 / 410 ms
PR	134 ms
P	100 ms
RR / PP	988 / 983 ms
P / QRS / T	63 / 47 / 56 degrees



*Normal graph*

**D. (COL.) M. SITARAM**  
MD, DM, FICG, FCSI  
Sr. Consultant (Cardiology)  
Regd. No. APMC 5430

Unconfirmed



Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28058268
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 12:35 PM
Req.No	: BIL4574037	Reported on	: 10-Aug-2024 / 17:16 PM
		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	Pale Yellow	Straw to Yellow
Appearance Method:Physical	Slightly Cloudy	Clear
<b>Chemical Examination</b>		
Reaction and pH Method:Indicator	Acidic (6.0)	4.6-8.0
Specific gravity Method:Refractometry	1.004	1.000-1.035
Protein Method:Protein Error of pH indicators	Negative	Negative
Glucose Method:Glucose oxidase/Peroxidase	Negative	Negative
Blood Method:Peroxidase	<b>Positive (Trace)</b>	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	1-2	2 - 3 /hpf
Epithelial cells Method:Flow Digital Imaging/Microscopy	1-2	2 - 5 /hpf
RBC (erythrocytes) Method:Flow Digital Imaging/Microscopy	<b>2-3/hpf</b>	Absent
Casts Method:Flow Digital Imaging/Microscopy	Absent	Occasional hyaline casts may be seen



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





Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28056304
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 09:25 AM
Req.No	: BIL4574037	Reported on	: 11-Aug-2024 / 15:54 PM
		Reference	: Arcofemi Health Care Ltd -

TEST REPORT

DEPARTMENT OF CYTOPATHOLOGY

Pap Smear, Conventional

Cytology No	<b>C-8665/24</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 few parabasal cells. Moderate inflammation noted.
Organisms	Not present
Interpretation	<b>Negative for intraepithelial lesion or malignancy.</b>
.	<b>Inflammatory smear</b>
Note	Kindly correlate clinically

**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.ARATI CH** TID/SID : UMR1837799/ 28056166  
Age / Gender : 48 Years / Female Registered on : 10-Aug-2024 / 09:14 AM  
Ref.By : SELF Collected on : 10-Aug-2024 / 09:16 AM  
Req.No : BIL4574037 Reported on : 10-Aug-2024 / 14:43 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)	O
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.ARATI CH** TID/SID : UMR1837799/ 28056166  
Age / Gender : 48 Years / Female Registered on : 10-Aug-2024 / 09:14 AM  
Ref.By : SELF Collected on : 10-Aug-2024 / 09:16 AM  
Req.No : BIL4574037 Reported on : 10-Aug-2024 / 13:48 PM  
Reference : Arcofemi Health Care Ltd -

**TEST REPORT**

**DEPARTMENT OF HEMATOPATHOLOGY**

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

Investigation	Observed Value	Biological Reference Intervals
ESR 1st Hour Method:Westergren/Vesmatic	<b>17</b>	<=12 mm/hour

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

--- End Of Report ---



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





Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28056166
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 09:16 AM
Req.No	: BIL4574037	Reported on	: 10-Aug-2024 / 13:48 PM
		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**DEPARTMENT OF HEMATOPATHOLOGY**

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

Investigation	Observed Value	Biological Reference Intervals
Hemoglobin Method:Cyanide Free Lyse Hemoglobin	12.9	12.0-15.0 g/dL
PCV/HCT Method:Calculated	38.4	36.0-46.0 vol%
Total RBC Count Method:Electrical Impedance	4.30	3.80-4.80 mill /cu.mm
MCV Method:Calculated	89.4	83.0-101.0 fL
MCH Method:Calculated	30.1	27.0-32.0 pg
MCHC Method:Calculated	33.6	31.5-34.5 g/dL
RDW (CV) Method:Calculated	13.7	11.6-14.0 %
MPV Method:Calculated	8.0	7.0-10.0 fL
Total WBC Count Method:Electrical Impedance	6530	4000-10000 cells/cumm
Platelet Count Method:Electrical Impedance	2.82	1.50-4.10 lakhs/cumm
<b>Differential count</b>		
Neutrophils Method:Microscopy	57.3	40.0-80.0 %
Lymphocytes Method:Microscopy	32.1	20.0-40.0 %
Eosinophils	3.3	1.0-6.0 %
Monocytes	7.3	2.0-10.0 %
Basophils Method:Microscopy	0.0	< 1.0-2.0 %
Absolute Neutrophil Count Method:Calculated	3742	2000-7000 cells/cumm
Absolute Lymphocyte Count (ALC)	2096	1000-3000 cells/cumm
Absolute Eosinophil Count (AEC)	215	20-500 cells/cumm
Absolute Monocyte Count Method:Calculated	477	200-1000 cells/cumm



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Reference : Arcofemi Health Care Ltd -

**TEST REPORT**

Absolute Basophil Count **0** 20-100 cells/cumm  
Method:Calculated  
Neutrophil - Lymphocyte Ratio(NLR) **1.79** 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.ARATI CH	TID/SID	: UMR1837799/ 28056167
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 09:16 AM
Req.No	: BIL4574037	Reported on	: 10-Aug-2024 / 13:49 PM
		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	7	6-20 mg/dL
Urea. Method:Urease/UV	14.6	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

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

--- End Of Report ---

**Dr.Abdur Rehman Asif**  
Consultant Biochemist  
Reg.No - APMC/FMR/78102





Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28056167
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 09:16 AM
Req.No	: BIL4574037	Reported on	: 10-Aug-2024 / 13:49 PM
		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

**Creatinine, Serum**

Investigation	Observed Value	Biological Reference Interval
Creatinine. Method:Alkaline Picrate	0.57	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.

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

--- End Of Report ---

**Dr.Abdur Rehman Asif**  
Consultant Biochemist  
Reg.No - APMC/FMR/78102





Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28056168F
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 09:16 AM
Req.No	: BIL4574037	Reported on	: 10-Aug-2024 / 13:49 PM
		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

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

Investigation	Observed Value	Biological Reference Interval
Glucose Fasting Method:Hexokinase	89	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

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

--- End Of Report ---

**Dr.Abdur Rehman Asif**  
Consultant Biochemist  
Reg.No - APMC/FMR/78102





Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28056168P
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 12:07 PM
Req.No	: BIL4574037	Reported on	: 10-Aug-2024 / 14:48 PM
		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

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

Investigation	Observed Value	Biological Reference Interval
Glucose Post Prandial Method:Hexokinase	<b>85</b>	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

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

--- End Of Report ---

**Dr.Abdur Rehman Asif**  
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**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

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

Investigation	Observed Value	Biological Reference Interval
Glycosylated Hemoglobin (HbA1c) Method:High-Performance Liquid Chromatography	5.1	Non-diabetic: <= 5.6 % Pre-diabetic: 5.7 - 6.4 % Diabetic: >= 6.5 %
Estimated Average Glucose (eAG) Method:Calculated	100	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 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.

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

--- End Of Report ---

**Dr.Abdur Rehman Asif**  
Consultant Biochemist  
Reg.No - APMC/FMR/78102



PLEASE SCAN QR CODE  
TO VERIFY THE REPORT ONLINE



Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28056167
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 09:16 AM
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**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

**Lipid Profile, Serum**

Investigation	Observed Value	Biological Reference Interval
Total Cholesterol Method:Cholesterol Oxidase	153	Desirable: <200 mg/dL Borderline: 200-239 mg/dL High: >=240 mg/dL
HDL Cholesterol Method:Direct Measurement	57	Low: <40 mg/dL High: >=60 mg/dL
VLDL Cholesterol Method:Calculated	12.4	6.0-38.0 mg/dL
LDL Cholesterol Method:Calculated	83.6	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	62	Normal:<150 mg/dL Borderline: 150-199 mg/dL High: 200-499 mg/dL Very high: >=500 mg/dL
Chol/HDL Ratio Method:Calculated	2.7	Low Risk: 3.3-4.4 Average Risk: 4.5-7.1 Moderate Risk: 7.2-11.0
LDL Cholesterol/HDL Ratio Method:Calculated	1.47	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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Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28056167
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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	0.61	<1.2 mg/dL
Direct Bilirubin. Method:Diazo method	0.27	<0.30 mg/dL
Indirect Bilirubin. Method:Calculated	0.34	<0.9 mg/dL
Alanine Aminotransferase ,(ALT/SGPT) Method:UV wthout P5P	15	<34 U/L
Aspartate Aminotransferase,(AST/SGOT) Method:UV wthout P5P	20	<31 U/L
ALP (Alkaline Phosphatase). Method:PNPP-AMP Buffer	87	35-104 U/L
Gamma GT. Method:Gamma-Glutamyl - 3 - Carbossi - 4 - Nitroanilide (GCNA)	17	6-42 U/L
Total Protein. Method:Biuret	7.7	6.6-8.7 g/dL
Albumin. Method:Bromocresol Green (BCG)	4.2	3.5-5.2 g/dL
Globulin. Method:Calculated	3.5	1.8-3.8 g/dL
A/GRatio. Method:Calculated	1.2	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.Abdur Rehman Asif**  
Consultant Biochemist  
Reg.No - APMC/FMR/78102



Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28056167
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 09:16 AM
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		Reference	: Arcofemi Health Care Ltd -

**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

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

Investigation	Observed Value	Biological Reference Interval
Triiodothyronine Total (T3) Method:ECLIA	1.36	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	9.7	5.1-14.1 µg/dL
Thyroid Stimulating Hormone (TSH) Method:ECLIA	<b>4.45</b>	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

Note Kindly correlate clinically

**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.Abdur Rehman Asif**  
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Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28056167
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 09:16 AM
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**TEST REPORT**

**DEPARTMENT OF CLINICAL CHEMISTRY I**

**Uric Acid, Serum**

Investigation	Observed Value	Biological Reference Interval
Uric Acid. Method:Uricase	4.0	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 ---

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Consultant Biochemist  
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Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28056167
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 09:16 AM
Req.No	: BIL4574037	Reported on	: 10-Aug-2024 / 13:49 PM
		Reference	: Arcofemi Health Care Ltd -

TEST REPORT

DEPARTMENT OF CLINICAL CHEMISTRY I

Bun/Creatinine Ratio, Serum

Investigation	Observed Value	Reference Range
BUN/Creatinine Ratio	12	10-20
Method: Calculated		

**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. Abdur Rehman Asif**  
Consultant Biochemist  
Reg.No - APMC/FMR/78102

**R**



ARATI CH F BIL4574037 22096040 CHEST PA 10-08-2024  
TENET DIAGNOSTICS, VIKARAMPURI, SECUNDERABAD



Name	: MRS.ARATI CH	TID/SID	: UMR1837799/ 28056304
Age / Gender	: 48 Years / Female	Registered on	: 10-Aug-2024 / 09:14 AM
Ref.By	: SELF	Collected on	: 10-Aug-2024 / 09:25 AM
Req.No	: BIL4574037	Reported on	: 11-Aug-2024 / 15:54 PM
		Reference	: Arcofemi Health Care Ltd -

TEST REPORT

DEPARTMENT OF CYTOPATHOLOGY

Pap Smear, Conventional

Cytology No	<b>C-8665/24</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 few parabasal cells. Moderate inflammation noted.
Organisms	Not present
Interpretation	<b>Negative for intraepithelial lesion or malignancy.</b>
.	<b>Inflammatory smear</b>
Note	Kindly correlate clinically

**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





PLEASE SCAN QR CODE

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Name	: Mrs . ARATI CH	TID	: UMR1837799
Age/Gender	: 48 Years/Female	Registered On	: 10-Aug-2024 09:14 AM
Ref By	: Self	Reported On	: 10-Aug-2024 11:57 AM
Reg.No	: BIL4574037	Reference	: Arcofemi Health Care Ltd - Medi Whe

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DEPARTMENT OF MAMMOGRAPHY  
**Mammography Bilateral**

Mediolateral oblique and craniocaudal views was performed.

**BILATERAL MAMMOGRAPHY:**

Bilateral breasts show symmetrical fibroglandular parenchyma.

No evidence of focal soft tissue lesion.

No evidence of cluster microcalcification.

Subcutaneous fat deposition is within normal limits.

**IMPRESSION:**

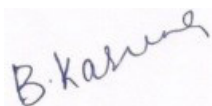
\* **Essentially normal study.**  
**BI-RADS CATEGORY - 1.**

**BI-RADS CLASSIFICATION**  
**CATEGORY 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%)<br>4B: Moderate suspicion for malignancy (10 - 49%)<br>4C : High suspicion for malignancy (50 - 94%)<br>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 \*\*\*

  
**Dr.KARUNA BELIDE**  
Consultant Radiologist



PLEASE SCAN QR CODE

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Name	: Mrs . ARATI CH	TID	: UMR1837799
Age/Gender	: 48 Years/Female	Registered On	: 10-Aug-2024 09:14 AM
Ref By	: Self	Reported On	: 10-Aug-2024 11:57 AM
Reg.No	: BIL4574037	Reference	: Arcofemi Health Care Ltd - Medi Whe

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PLEASE SCAN QR CODE

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Name	: Mrs . ARATI CH	TID	: UMR1837799
Age/Gender	: 48 Years/Female	Registered On	: 10-Aug-2024 09:14 AM
Ref By	: Self	Reported On	: 10-Aug-2024 12:53 PM
Reg.No	: BIL4574037	Reference	: Arcofemi Health Care Ltd - Medi Whe

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### Ultrasound Whole Abdomen

**LIVER** is normal shape, size ( 11.3 cms) and has uniform echopattern.  
No evidence of focal lesion. No intrahepatic biliary ductal dilatation.  
Hepatic and portal vein radicals are normal. Portal vein measures : 5.6 mm.

**GALL BLADDER** status post cholecystectomy.

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 ( 8.7 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 9.4 x 3.9 cms, Left kidney measures 9.9 x 4.0 cms.

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

**UTERUS** is anteverted. Mild bulky in size.  
Endometrial thickness measures 9.7 mm.  
Uterus measures 9.8 x 3.9 x 4.6 cms.

**OVARIES** are normal in size, shape and echotexture.  
Right ovary: 2.2 x 1.4 cms, Left ovary: 2.9 x 1.8 cms.

No evidence of free fluid in the abdomen and pelvis.

### IMPRESSION:

\* **Mildly bulky uterus.**

Suggested clinical correlation and follow up

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

**Dr.KARUNA BELIDE**  
Consultant Radiologist



PLEASE SCAN QR CODE

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Name : Mrs . ARATI CH  
Age/Gender : 48 Years/Female  
Ref By : Self  
Reg.No : BIL4574037

TID : UMR1837799  
Registered On : 10-Aug-2024 09:14 AM  
Reported On : 10-Aug-2024 12:53 PM  
Reference : Arcofemi Health Care Ltd  
- Medi Whe

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PLEASE SCAN QR CODE

Name	: Mrs . ARATI CH	TID	: UMR1837799
Age/Gender	: 48 Years/Female	Registered On	: 10-Aug-2024 09:14 AM
Ref By	: Self	Reported On	: 10-Aug-2024 02:03 PM
Reg.No	: BIL4574037	Reference	: Arcofemi Health Care Ltd - Medi Whe

**DEPARTMENT OF CARDIOLOGY**  
**2D Echo/Doppler Study**

MITRAL VALVE : Normal.

AORTIC VALVE : Normal.

TRICUSPID VALVE : Normal.

PULMONARY VALVE : Normal.

RIGHT ATRIUM : Normal.

RIGHT VENTRICLE : Normal.

LEFT ATRIUM : 2.8 cms.

LEFT VENTRICLE : EDD : 3.8 cm      IVS (d) : 0.7 cm      LVEF : 68 %  
 ESD : 2.3 cm      PW (d) : 0.7 cm      FS : 37 %  
 NO RWMA

IAS : Intact.

IVS : Intact.

AORTA : 2.1 cms.

PULMONARY ARTERY : Normal

PERICARDIUM : Normal.

IVC / SVC / CS : Normal.

PULMONARY VEINS : Normal.

INTRA - CARDIAC MASSES : No.

**DOPPLER STUDY**



PLEASE SCAN QR CODE

Name : Mrs . ARATI CH  
Age/Gender : 48 Years/Female  
Ref By : Self  
Reg.No : BIL4574037

TID : UMR1837799  
Registered On : 10-Aug-2024 09:14 AM  
Reported On : 10-Aug-2024 02:03 PM  
Reference : Arcofemi Health Care Ltd  
- Medi Whe

MITRAL FLOW : E: 0.7 m/s A: 0.7 m/s  
AORTIC FLOW : 1.4 m/s  
PULMONARY FLOW : 0.9 m/s  
TRICUSPID FLOW : E : 0.5 m/s A : 0.2 m/s

**COLOUR FLOW MAPPING**

MR : NIL  
AR : NIL  
TR : NIL  
PR : NIL

**IMPRESSION:**

- \* NORMAL LV SIZE & CONTRACTILITY
- \* NO RWMA
- \* GOOD LV / RV FUNCTION
- \* NO MR / AR / TR
- \* NO LA / LV CLOTS / NO PE

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

**Dr.M.Sitaram**  
MD DM FICC FCSI  
Sr. Consultant Cardiologist



PLEASE SCAN QR CODE

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Name	: Mrs . ARATI CH	TID	: UMR1837799
Age/Gender	: 48 Years/Female	Registered On	: 10-Aug-2024 09:14 AM
Ref By	: Self	Reported On	: 10-Aug-2024 05:04 PM
Reg.No	: BIL4574037	Reference	: Arcofemi Health Care Ltd - Medi Whe

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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:**

**\* Normal study.**

Suggested clinical correlation and follow up

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

**Dr.KARUNA BELIDE**  
Consultant Radiologist