



Hiranandani  
HOSPITAL

(A Fortis Network Hospital)

Hiranandani Fortis Hospital  
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Navi Mumbai - 400 703.  
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## BMI CHART

Date: 01/04/23

Name: Mrs Anil Age: 28 yrs Sex: M/F

BP: 120/80 Height (cms): 161 cm Weight(kgs): 88.8 BMI: 34

*mmHg*

WEIGHT lbs 100 105 110 115 120 125 130 135 140 145 150 155 160 165 170 175 180 185 190 195 200 205 210 215  
kgs 45.5 47.7 50.0 52.3 54.5 56.8 59.1 61.4 63.6 65.9 68.2 70.5 72.7 75.0 77.3 79.5 81.8 84.1 86.4 88.6 90.9 93.2 95.5 97.7

HEIGHT in/cm	Underweight				Healthy				Overweight				Obese				Extremely Obese							
	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42
5'0" - 152.4	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42
5'1" - 154.9	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41
5'2" - 157.4	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41
5'3" - 160.0	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40
5'4" - 162.5	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40
5'5" - 165.1	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39
5'6" - 167.6	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39
5'7" - 170.1	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38
5'8" - 172.7	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38
5'9" - 176.2	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37
5'10" - 177.8	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37
5'11" - 180.3	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37
6'0" - 182.8	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36
6'1" - 185.4	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36
6'2" - 187.9	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35
6'3" - 190.5	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35
6'4" - 193.0	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35

### Doctors Notes:

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UHID	11008739	Date	01/04/2023		
Name	Mrs.Aarti	Sex	Female	Age	28
OPD	Opthal 14	Health Check Up			

Drug allergy:  
 Sys illness:

Ref → RE → -0.50 Dm 6/6.  
 LG → -0.50 Dm 6/6.

MA → W6  
 W6

I.O.P. → RE → 13.8  
 LG → 12.7.

Antseg (WNL)

fundus (WNL)

(WNL)

NA

Rep

Aqualube eed

→ →


NA

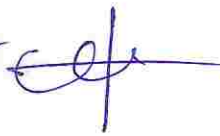


<b>UHID</b>	11008739	<b>Date</b>	01/04/2023		
<b>Name</b>	Mrs.Aarti	<b>Sex</b>	Female	<b>Age</b>	28
<b>OPD</b>	Dental 12 7387696540	<b>Health Check Up</b>			

Drug allergy:  
 Sys illness:

Impacted 

Caries 

stains ++ calculus ++ 


Treatment

Adv OPG.

Adv filling 

Adv surgical removal 

Adv oral prophylaxis

  
 Dr. Diksha Kataria



UHID	11008739	Date	01/04/2023		
Name	Mrs.Aarti	Sex	Female	Age	28
OPD	Pap Smear	Health Check Up			

Zyges/ P/L

Drug allergy:  
 Sys illness:

LMP: 22.3.23

Pmo: 3/3/21, RMP  
 -40

Psp - op/⊕ pap ✓  
 ng/⊕

Breast exam<sup>n</sup> ⊕

Adv

- Pap smear Zyges
- self breast exam<sup>n</sup> m/hly

Adv



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 FORTIS VASHI-CHC -SPLZD  
 FORTIS HOSPITAL # VASHI,  
 MUMBAI 440001

ACCESSION NO : 0022WD000117  
 PATIENT ID : FH.11008739  
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AGE/SEX : 28 Years Female  
 DRAWN : 01/04/2023 10:25:00  
 RECEIVED : 01/04/2023 10:24:54  
 REPORTED : 01/04/2023 13:59:52

CLINICAL INFORMATION :

UID:11008739 REQNO-1454619  
 CORP-OPD  
 BILLNO-150123OPCR019077  
 BILLNO-150123OPCR019077

Test Report Status	Final	Results	Biological Reference Interval	Units
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HAEMATOLOGY - CBC

CBC-5, EDTA WHOLE BLOOD

BLOOD COUNTS, EDTA WHOLE BLOOD

HEMOGLOBIN (HB)	12.8	12.0 - 15.0	g/dL
METHOD : SPECTROPHOTOMETRY			
RED BLOOD CELL (RBC) COUNT	4.87 High	3.8 - 4.8	mil/ $\mu$ L
METHOD : ELECTRICAL IMPEDANCE			
WHITE BLOOD CELL (WBC) COUNT	7.78	4.0 - 10.0	thou/ $\mu$ L
METHOD : DOUBLE HYDRODYNAMIC SEQUENTIAL SYSTEM(DHSS)CYTOMETRY			
PLATELET COUNT	373	150 - 410	thou/ $\mu$ L
METHOD : ELECTRICAL IMPEDANCE			
<b>RBC AND PLATELET INDICES</b>			
HEMATOCRIT (PCV)	37.9	36 - 46	%
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR VOLUME (MCV)	77.9 Low	83 - 101	fL
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	26.3 Low	27.0 - 32.0	pg
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION(MCHC)	33.7	31.5 - 34.5	g/dL
METHOD : CALCULATED PARAMETER			
RED CELL DISTRIBUTION WIDTH (RDW)	14.3 High	11.6 - 14.0	%
METHOD : CALCULATED PARAMETER			
MENTZER INDEX	16.0		
MEAN PLATELET VOLUME (MPV)	9.0	6.8 - 10.9	fL
METHOD : CALCULATED PARAMETER			
<b>WBC DIFFERENTIAL COUNT</b>			
NEUTROPHILS	55	40 - 80	%
METHOD : FLOWCYTOMETRY			
LYMPHOCYTES	38	20 - 40	%
METHOD : FLOWCYTOMETRY			

*Akta Dubey*

Dr.Akta Dubey  
 Consultant Pathologist



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MONOCYTES		5	2 - 10	%
METHOD : FLOWCYTOMETRY				
EOSINOPHILS		2	1 - 6	%
METHOD : FLOWCYTOMETRY				
BASOPHILS		00	0 - 2	%
METHOD : FLOWCYTOMETRY				
ABSOLUTE NEUTROPHIL COUNT		4.28	2.0 - 7.0	thou/ $\mu$ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE LYMPHOCYTE COUNT		2.96	1.0 - 3.0	thou/ $\mu$ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE MONOCYTE COUNT		0.39	0.2 - 1.0	thou/ $\mu$ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE EOSINOPHIL COUNT		0.16	0.02 - 0.50	thou/ $\mu$ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE BASOPHIL COUNT		0 Low	0.02 - 0.10	thou/ $\mu$ L
METHOD : CALCULATED PARAMETER				
NEUTROPHIL LYMPHOCYTE RATIO (NLR)		1.4		
METHOD : CALCULATED PARAMETER				
<b>MORPHOLOGY</b>				
RBC		PREDOMINANTLY NORMOCYTIC NORMOCHROMIC, MILD MICROCYTOSIS		
METHOD : MICROSCOPIC EXAMINATION				
WBC		NORMAL MORPHOLOGY		
METHOD : MICROSCOPIC EXAMINATION				
PLATELETS		ADEQUATE		
METHOD : MICROSCOPIC EXAMINATION				

Interpretation(s)

RBC AND PLATELET INDICES-Mentzer index (MCV/RBC) is an automated cell-counter based calculated screen tool to differentiate cases of Iron deficiency anaemia(>13) from Beta thalassaemia trait (<13) in patients with microcytic anaemia. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HbA2 remains the gold standard for diagnosing a case of beta thalassaemia trait.

Dr.Akta Dubey  
 Consultant Pathologist



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Patient Ref. No. 22000000838134

**LABORATORY REPORT**



**Fortis**

**SRL**  
Diagnostics

REF. DOCTOR : SELF

**PATIENT NAME : MRS.AARTI .**

**CODE/NAME & ADDRESS : C000045507 - FORTIS**  
FORTIS VASHI-CHC -SPLZD  
FORTIS HOSPITAL # VASHI,  
MUMBAI 440001

**ACCESSION NO : 0022WD000117**  
**PATIENT ID : FH.11008739**  
**CLIENT PATIENT ID: UID:11008739**  
**ABHA NO :**

**AGE/SEX : 28 Years Female**  
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Final			

WBC DIFFERENTIAL COUNT-The optimal threshold of 3.3 for NLR showed a prognostic possibility of clinical symptoms to change from mild to severe in COVID positive patients. When age = 49.5 years old and NLR = 3.3, 46.1% COVID-19 patients with mild disease might become severe. By contrast, when age < 49.5 years old and NLR < 3.3, COVID-19 patients tend to show mild disease.  
(Reference to - The diagnostic and predictive role of NLR, d-NLR and PLR in COVID-19 patients ; A.-P. Yang, et al.; International Immunopharmacology 84 (2020) 106504  
This ratio element is a calculated parameter and out of NABL scope.

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HAEMATOLOGY

ERYTHROCYTE SEDIMENTATION RATE (ESR), WHOLE BLOOD

E.S.R	12	0 - 20	mm at 1 hr
METHOD : WESTERGRÉN METHOD			

Interpretation(s)

ERYTHROCYTE SEDIMENTATION RATE (ESR), WHOLE BLOOD-TEST DESCRIPTION :-

Erythrocyte sedimentation rate (ESR) is a test that indirectly measures the degree of inflammation present in the body. The test actually measures the rate of fall (sedimentation) of erythrocytes in a sample of blood that has been placed into a tall, thin, vertical tube. Results are reported as the millimetres of clear fluid (plasma) that are present at the top portion of the tube after one hour. Nowadays fully automated instruments are available to measure ESR.

ESR is not diagnostic; it is a non-specific test that may be elevated in a number of different conditions. It provides general information about the presence of an inflammatory condition. CRP is superior to ESR because it is more sensitive and reflects a more rapid change.

TEST INTERPRETATION

**Increase** in: Infections, Vasculitides, Inflammatory arthritis, Renal disease, Anemia, Malignancies and plasma cell dyscrasias, Acute allergy Tissue injury, Pregnancy, Estrogen medication, Aging.

Finding a very accelerated ESR (>100 mm/hour) in patients with ill-defined symptoms directs the physician to search for a systemic disease (Paraproteinemias, Disseminated malignancies, connective tissue disease, severe infections such as bacterial endocarditis).

In pregnancy BRI in first trimester is 0-48 mm/hr (52 if anemic) and in second trimester (0-70 mm/hr (95 if anemic). ESR returns to normal 4th week post partum.

**Decreased** in: Polycythemia vera, Sickle cell anemia

LIMITATIONS

**False elevated ESR** : Increased fibrinogen, Drugs (Vitamin A, Dextran etc), Hypercholesterolemia  
**False Decreased** : Poikilocytosis, (Sickle Cells, spherocytes), Microcytosis, Low fibrinogen, Very high WBC counts, Drugs (Quinine, salicylates)

REFERENCE :

1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition; 2. Paediatric reference intervals. AACCPress, 7th edition. Edited by S. Soldin; 3. The reference for the adult reference range is "Practical Haematology by Dacie and Lewis, 10th edition.

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IMMUNOHAEMATOLOGY

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP TYPE B  
 METHOD : TUBE AGGLUTINATION  
 RH TYPE POSITIVE  
 METHOD : TUBE AGGLUTINATION

Interpretation(s)

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD-  
 Blood group is identified by antigens and antibodies present in the blood. Antigens are protein molecules found on the surface of red blood cells. Antibodies are found in plasma. To determine blood group, red cells are mixed with different antibody solutions to give A,B,O or AB.

Disclaimer: "Please note, as the results of previous ABO and Rh group (Blood Group) for pregnant women are not available, please check with the patient records for availability of the same."

The test is performed by both forward as well as reverse grouping methods.

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BIOCHEMISTRY

LIVER FUNCTION PROFILE, SERUM

BILIRUBIN, TOTAL METHOD : JENDRASSIK AND GROFF	0.48	0.2 - 1.0	mg/dL
BILIRUBIN, DIRECT METHOD : JENDRASSIK AND GROFF	0.10	0.0 - 0.2	mg/dL
BILIRUBIN, INDIRECT METHOD : CALCULATED PARAMETER	0.38	0.1 - 1.0	mg/dL
TOTAL PROTEIN METHOD : BIURET	8.0	6.4 - 8.2	g/dL
ALBUMIN METHOD : BCP DYE BINDING	4.5	3.4 - 5.0	g/dL
GLOBULIN METHOD : CALCULATED PARAMETER	3.5	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO METHOD : CALCULATED PARAMETER	1.3	1.0 - 2.1	RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT) METHOD : UV WITH PSP	16	15 - 37	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT) METHOD : UV WITH PSP	25	< 34.0	U/L
ALKALINE PHOSPHATASE METHOD : PNPP-ANP	112	30 - 120	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT) METHOD : GAMMA GLUTAMYL CARBOXY 4NITROANTILIDE	34	5 - 55	U/L
LACTATE DEHYDROGENASE METHOD : LACTATE -PYRUVATE	131	100 - 190	U/L

GLUCOSE FASTING, FLUORIDE PLASMA

FBS (FASTING BLOOD SUGAR) METHOD : HEXOKINASE	95	74 - 99	mg/dL
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GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD

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HBA1C		5.4	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 Therapeutic goals: < 7.0 Action suggested : > 8.0 (ADA Guideline 2021)	%
METHOD : HB VARIANT (HPLC)				
ESTIMATED AVERAGE GLUCOSE(EAG)		108.3	< 116.0	mg/dL
METHOD : CALCULATED PARAMETER				
<b>KIDNEY PANEL - 1</b>				
<b>BLOOD UREA NITROGEN (BUN), SERUM</b>				
BLOOD UREA NITROGEN		9	6 - 20	mg/dL
METHOD : UREASE - UV				
<b>CREATININE EGFR- EPI</b>				
CREATININE		0.65	0.60 - 1.10	mg/dL
METHOD : ALKALINE PICRATE KINETIC JAFFES				
AGE		28		years
GLOMERULAR FILTRATION RATE (FEMALE)		122.91	Refer Interpretation Below	mL/min/1.73m2
METHOD : CALCULATED PARAMETER				

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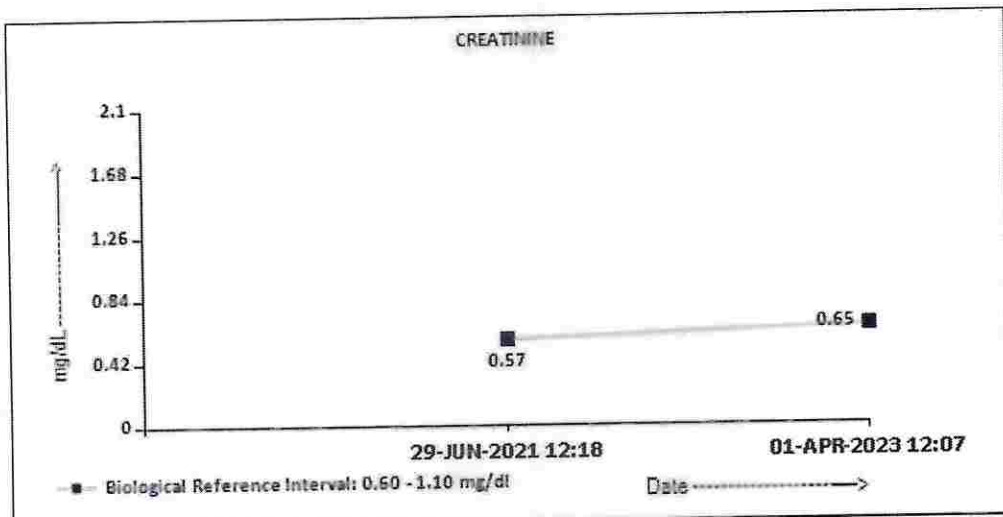
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 BILLNO-150123OPCR019077

Test Report Status	Results	Biological Reference Interval	Units
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BUN/CREAT RATIO

BUN/CREAT RATIO 13.85 5.00 - 15.00  
 METHOD : CALCULATED PARAMETER

URIC ACID, SERUM

URIC ACID 5.0 2.6 - 6.0 mg/dL  
 METHOD : URICASE UV

TOTAL PROTEIN, SERUM

TOTAL PROTEIN 8.0 6.4 - 8.2 g/dL  
 METHOD : BIURET

ALBUMIN, SERUM

ALBUMIN 4.5 3.4 - 5.0 g/dL  
 METHOD : BCP DYE BINDING

Dr. Akta Dubey  
 Consultant Pathologist



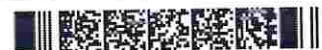
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 NAVI MUMBAI, 400703  
 MAHARASHTRA, INDIA  
 Tel : 022-39199222, 022-49723322,  
 CIN - U74899PB1995PLC045956  
 Email : -



Patient Ref. No. 22000000838134



REF. DOCTOR : SELF

PATIENT NAME : MRS.AARTI .

CODE/NAME & ADDRESS : C000045507 - FORTIS  
 FORTIS VASHI-CHC -SPLZD  
 FORTIS HOSPITAL # VASHI,  
 MUMBAI 440001

ACCESSION NO : 0022WD000117  
 PATIENT ID : FH.11008739  
 CLIENT PATIENT ID: UID:11008739  
 ABHA NO :

AGE/SEX : 28 Years Female  
 DRAWN : 01/04/2023 10:25:00  
 RECEIVED : 01/04/2023 10:24:54  
 REPORTED : 01/04/2023 13:59:52

CLINICAL INFORMATION :

UID:11008739 REQNO-1454619  
 CORP-OPD  
 BILLNO-150123OPCR019077  
 BILLNO-150123OPCR019077

Test Report Status	Final	Results	Biological Reference Interval	Units
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LOBULIN

LOBULIN

METHOD : CALCULATED PARAMETER

3.5

2.0 - 4.1

g/dL

ELECTROLYTES (NA/K/CL), SERUM

SODIUM, SERUM

METHOD : ISE INDIRECT

138

136 - 145

mmol/L

POTASSIUM, SERUM

METHOD : ISE INDIRECT

4.69

3.50 - 5.10

mmol/L

CHLORIDE, SERUM

METHOD : ISE INDIRECT

101

98 - 107

mmol/L

Interpretation(s)

Interpretation(s)

LIVER FUNCTION PROFILE, SERUM-  
 LIVER FUNCTION PROFILE

Bilirubin is a yellowish pigment found in bile and is a breakdown product of normal heme catabolism. Bilirubin is excreted in bile and urine, and elevated levels may give yellow discoloration in jaundice.

Elevated levels results from increased bilirubin production (eg, hemolysis and ineffective erythropoiesis), decreased bilirubin excretion (eg, obstruction and hepatitis), and abnormal bilirubin metabolism (eg, hereditary and neonatal jaundice). Conjugated (direct) bilirubin is elevated more than unconjugated (indirect) bilirubin in Viral hepatitis, Drug reactions, Alcoholic liver disease. Conjugated (direct) bilirubin is also elevated more than unconjugated (indirect) bilirubin when there is some kind of blockage of the bile ducts like in Gallstones getting into the bile ducts, tumors & Scarring of the bile ducts. Increased unconjugated (indirect) bilirubin may be a result of Hemolytic or pernicious anemia, Transfusion reaction & a common metabolic condition termed Gilbert syndrome, due to low levels of the enzyme that attaches sugar molecules to bilirubin.

AST is an enzyme found in various parts of the body. AST is found in the liver, heart, skeletal muscle, kidneys, brain, and red blood cells, and it is commonly measured clinically as a marker for liver health. AST levels increase during chronic viral hepatitis, blockage of the bile duct, cirrhosis of the liver, liver cancer, kidney failure, hemolytic anemia, pancreatitis, hemochromatosis. AST levels may also increase after a heart attack or strenuous activity. ALT test measures the amount of this enzyme in the blood. ALT is found mainly in the liver, but also in smaller amounts in the kidneys, heart, muscles, and pancreas. It is commonly measured as a part of a diagnostic evaluation of hepatocellular injury, to determine liver health. AST levels increase during acute hepatitis, sometimes due to a viral infection, ischemia to the liver, chronic hepatitis, obstruction of bile ducts, cirrhosis.

ALP is a protein found in almost all body tissues. Tissues with higher amounts of ALP include the liver, bile ducts and bone. Elevated ALP levels are seen in Biliary obstruction, Osteoblastic bone tumors, osteomalacia, hepatitis, Hyperparathyroidism, Leukemia, Lymphoma, Pagets disease, Rickets, Sarcoidosis etc. Lower-than-normal ALP levels seen in Hypophosphatasia, Malnutrition, Protein deficiency, Wilsons disease.

GGT is an enzyme found in cell membranes of many tissues mainly in the liver, kidney and pancreas. It is also found in other tissues including intestine, spleen, heart, brain and seminal vesicles. The highest concentration is in the kidney, but the liver is considered the source of normal enzyme activity. Serum GGT has been widely used as an index of liver dysfunction. Elevated serum GGT activity can be found in diseases of the liver, biliary system and pancreas. Conditions that increase serum GGT are obstructive liver disease, high alcohol consumption and use of enzyme-inducing drugs etc.

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 Consultant Pathologist



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**Total Protein** also known as total protein, is a biochemical test for measuring the total amount of protein in serum. Protein in the plasma is made up of albumin and globulin. Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenstroms disease. Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage), Burns, Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic syndrome, Protein-losing enteropathy etc.

**Albumin** is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by: Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance, malnutrition and wasting etc.

**GLUCOSE FASTING, FLUORIDE PLASMA-TEST DESCRIPTION**

Normally, the glucose concentration in extracellular fluid is closely regulated so that a source of energy is readily available to tissues and so that no glucose is excreted in the urine.

**Increased in:** Diabetes mellitus, Cushing's syndrome (10 – 15%), chronic pancreatitis (30%). Drugs: corticosteroids, phenytoin, estrogen, thiazides.

**Decreased in:** Pancreatic islet cell disease with increased insulin, insulinoma, adrenocortical insufficiency, hypopituitarism, diffuse liver disease, malignancy (adrenocortical, stomach, fibrosarcoma), infant of a diabetic mother, enzyme deficiency diseases (e.g. galactosemia), Drugs-insulin, ethanol, propranolol; sulfonylureas, tolbutamide, and other oral hypoglycemic agents.

**NOTE:** While random serum glucose levels correlate with home glucose monitoring results (weekly mean capillary glucose values), there is wide fluctuation within individuals. Thus, glycosylated hemoglobin (HbA1c) levels are favored to monitor glycemic control.

High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glycosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc.

**GLYCOSYLATED HEMOGLOBIN (HBA1C), EDTA WHOLE BLOOD-Used For:**

- Evaluating the long-term control of blood glucose concentrations in diabetic patients.
- Diagnosing diabetes.
- Identifying patients at increased risk for diabetes (prediabetes).

The ADA recommends measurement of HbA1c (typically 3-4 times per year for type 1 and poorly controlled type 2 diabetic patients, and 2 times per year for well-controlled type 2 diabetic patients) to determine whether a patient's metabolic control has remained continuously within the target range.

- eAG (Estimated average glucose) converts percentage HbA1c to mg/dl, to compare blood glucose levels.
- eAG gives an evaluation of blood glucose levels for the last couple of months.
- eAG is calculated as  $eAG (mg/dl) = 28.7 * HbA1c - 46.7$

**HbA1c Estimation can get affected due to :**

- Shortened Erythrocyte survival: Any condition that shortens erythrocyte survival or decreases mean erythrocyte age (e.g. recovery from acute blood loss, hemolytic anemia) will falsely lower HbA1c test results. Fructosamine is recommended in these patients which indicates diabetes control over 15 days.
- Vitamin C & E are reported to falsely lower test results (possibly by inhibiting glycation of hemoglobin).
- Iron deficiency anemia is reported to increase test results. Hypertriglyceridemia, uremia, hyperbilirubinemia, chronic alcoholism, chronic ingestion of salicylates & opiates addition are reported to interfere with some assay methods, falsely increasing results.
- Interference of hemoglobinopathies in HbA1c estimation is seen in

a) Homozygous hemoglobinopathy. Fructosamine is recommended for testing of HbA1c.

b) Heterozygous state detected (D10 is corrected for HbS & HbC trait.)

c) HbF > 25% on alternate platform (Boronate affinity chromatography) is recommended for testing of HbA1c. Abnormal Hemoglobin electrophoresis (HPLC method) is recommended for detecting a hemoglobinopathy.

**BLOOD UREA NITROGEN (BUN), SERUM-Causes of Increased levels include** Pre renal (High protein diet, Increased protein catabolism, GI haemorrhage, Cortisol, Dehydration, CHF Renal), Renal Failure, Post Renal (Malignancy, Nephrolithiasis, Prostatism)

**Causes of decreased level include** Liver disease, SIADH.

**CREATININE EGFR- EPI-GFR—** Glomerular filtration rate (GFR) is a measure of the function of the kidneys. The GFR is a calculation based on a serum creatinine test. Creatinine is a muscle waste product that is filtered from the blood by the kidneys and excreted into urine at a relatively steady rate. When kidney function decreases, less creatinine is excreted and concentrations increase in the blood. With the creatinine test, a reasonable estimate of the actual GFR can be determined.

A GFR of 60 or higher is in the normal range.

A GFR below 60 may mean kidney disease.

A GFR of 15 or lower may mean kidney failure.

Estimated GFR (eGFR) is the preferred method for identifying people with chronic kidney disease (CKD). In adults, eGFR calculated using the Modification of Diet in Renal Disease (MDRD) Study equation provides a more clinically useful measure of kidney function than serum creatinine alone.

The CKD-EPI creatinine equation is based on the same four variables as the MDRD Study equation, but uses a 2-slope spline to model the relationship between estimated GFR and serum creatinine, and a different relationship for age, sex and race. The equation was reported to perform better and with less bias than the MDRD Study equation, especially in patients with higher GFR. This results in reduced misclassification of CKD.

The CKD-EPI creatinine equation has not been validated in children & will only be reported for patients = 18 years of age. For pediatric and childrens, Schwartz Pediatric

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 Counsultant Pathologist



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Patient Ref. No. 22000000838134



REF. DOCTOR : SELF

PATIENT NAME : MRS.AARTI .

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 FORTIS VASHI-CHC -SPLZD  
 FORTIS HOSPITAL # VASHI,  
 MUMBAI 440001

ACCESSION NO : 0022WD000117  
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 ABHA NO :

AGE/SEX : 28 Years Female  
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Bedside eGFR (2009) formulae is used. This revised "bedside" pediatric eGFR requires only serum creatinine and height.  
 URIC ACID, SERUM-**Causes of Increased levels**:-Dietary(High Protein Intake,Prolonged Fasting,Rapid weight loss),Gout,Lesch nyhan syndrome,Type 2 DM,Metabolic syndrome **Causes of decreased levels**-Low Zinc Intake,OCP,Multiple Sclerosis  
 TOTAL PROTEIN, SERUM-is a biochemical test for measuring the total amount of protein in serum.Protein in the plasma is made up of albumin and globulin.  
**Higher-than-normal levels may be due to:** Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma,Waldenstroms disease.  
**Lower-than-normal levels may be due to:** Agammaglobulinemia, Bleeding (hemorrhage),Burns,Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic syndrome,Protein-losing enteropathy etc.  
 ALBUMIN, SERUM-  
 Human serum albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. **Low blood albumin levels (hypoalbuminemia) can be caused by:** Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance,malnutrition and wasting etc.

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Patient Ref. No. 2200000838134



<b>PATIENT NAME : MRS.AARTI .</b>		<b>REF. DOCTOR : SELF</b>	
<b>CODE/NAME &amp; ADDRESS :</b> C000045507 - FORTIS FORTIS VASHI-CHC -SPLZD FORTIS HOSPITAL # VASHI, MUMBAI 440001	<b>ACCESSION NO :</b> 0022WD000117	<b>AGE/SEX :</b> 28 Years Female	<b>DRAWN :</b> 01/04/2023 10:25:00
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**BIOCHEMISTRY - LIPID**

<b>LIPID PROFILE, SERUM</b>				
<b>CHOLESTEROL, TOTAL</b>	181	< 200 Desirable 200 - 239 Borderline High >/= 240 High		mg/dL
METHOD : ENZYMATIC/COLORIMETRIC, CHOLESTEROL OXIDASE, ESTERASE, PEROXIDASE				
<b>TRIGLYCERIDES</b>	146	< 150 Normal 150 - 199 Borderline High 200 - 499 High >/=500 Very High		mg/dL
METHOD : ENZYMATIC ASSAY				
<b>HDL CHOLESTEROL</b>	35 Low	< 40 Low >/=60 High		mg/dL
METHOD : DIRECT MEASURE - PEG				
<b>LDL CHOLESTEROL, DIRECT</b>	123	< 100 Optimal 100 - 129 Near or above optimal 130 - 159 Borderline High 160 - 189 High >/= 190 Very High		mg/dL
METHOD : DIRECT MEASURE WITHOUT SAMPLE PRETREATMENT				
<b>NON HDL CHOLESTEROL</b>	146 High	Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220		mg/dL
METHOD : CALCULATED PARAMETER				
<b>VERY LOW DENSITY LIPOPROTEIN</b>	29.2	</= 30.0		mg/dL
METHOD : CALCULATED PARAMETER				
<b>CHOL/HDL RATIO</b>	5.2 High	3.3 - 4.4 Low Risk 4.5 - 7.0 Average Risk 7.1 - 11.0 Moderate Risk > 11.0 High Risk		
METHOD : CALCULATED PARAMETER				
<b>LDL/HDL RATIO</b>	3.5 High	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk		
METHOD : CALCULATED PARAMETER				

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**Dr.Akta Dubey**  
 Consultant Pathologist



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Patient Ref. No. 22000000838134





MC-2275

**Fortis**

<b>PATIENT NAME : MRS.AARTI .</b>		<b>REF. DOCTOR : SELF</b>	
<b>CODE/NAME &amp; ADDRESS :</b> C000045507 - FORTIS FORTIS VASHI-CHC -SPLZD FORTIS HOSPITAL # VASHI, MUMBAI 440001	<b>ACCESSION NO :</b> <b>0022WD000117</b>	<b>AGE/SEX :</b> 28 Years Female	<b>DRAWN :</b> 01/04/2023 10:25:00
	<b>PATIENT ID :</b> FH.11008739	<b>RECEIVED :</b> 01/04/2023 10:24:54	<b>REPORTED :</b> 01/04/2023 13:59:52
	<b>CLIENT PATIENT ID:</b> UID:11008739		
	<b>ABHA NO :</b>		

**CLINICAL INFORMATION :**  
UID:11008739 REQNO-1454619  
CORP-OPD  
BILLNO-150123OPCR019077  
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<b>Final</b>			

**Interpretation(s)**

**Dr.Akta Dubey**  
Counsultant Pathologist



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**CLINICAL PATH - URINALYSIS**

**KIDNEY PANEL - 1**

**PHYSICAL EXAMINATION, URINE**

**COLOR** PALE YELLOW  
 METHOD : PHYSICAL

**APPEARANCE** CLEAR  
 METHOD : VISUAL

**CHEMICAL EXAMINATION, URINE**

**PH** 6.0 4.7 - 7.5  
 METHOD : REFLECTANCE SPECTROPHOTOMETRY- DOUBLE INDICATOR METHOD

**SPECIFIC GRAVITY** 1.020 1.003 - 1.035  
 METHOD : REFLECTANCE SPECTROPHOTOMETRY (APPARENT PKA CHANGE OF PRETREATED POLYELECTROLYTES IN RELATION TO IONIC CONCENTRATION)

**PROTEIN** NOT DETECTED NOT DETECTED  
 METHOD : REFLECTANCE SPECTROPHOTOMETRY - PROTEIN-ERROR-OF-INDICATOR PRINCIPLE

**GLUCOSE** NOT DETECTED NOT DETECTED  
 METHOD : REFLECTANCE SPECTROPHOTOMETRY, DOUBLE SEQUENTIAL ENZYME REACTION-GOD/POD

**KETONES** NOT DETECTED NOT DETECTED  
 METHOD : REFLECTANCE SPECTROPHOTOMETRY, ROTHERA'S PRINCIPLE

**BLOOD** NOT DETECTED NOT DETECTED  
 METHOD : REFLECTANCE SPECTROPHOTOMETRY, PEROXIDASE LIKE ACTIVITY OF HAEMOGLOBIN

**BILIRUBIN** NOT DETECTED NOT DETECTED  
 METHOD : REFLECTANCE SPECTROPHOTOMETRY, DIAZOTIZATION- COUPLING OF BILIRUBIN WITH DIAZOTIZED SALT

**UROBILINOGEN** NORMAL NORMAL  
 METHOD : REFLECTANCE SPECTROPHOTOMETRY (MODIFIED EHRlich REACTION)

**NITRITE** NOT DETECTED NOT DETECTED  
 METHOD : REFLECTANCE SPECTROPHOTOMETRY, CONVERSION OF NITRATE TO NITRITE

**LEUKOCYTE ESTERASE** NOT DETECTED NOT DETECTED  
 METHOD : REFLECTANCE SPECTROPHOTOMETRY, ESTERASE HYDROLYSIS ACTIVITY

**MICROSCOPIC EXAMINATION, URINE**

**RED BLOOD CELLS** NOT DETECTED NOT DETECTED /HPF  
 METHOD : MICROSCOPIC EXAMINATION

*Akta Dubey*  
**Dr. Akta Dubey**  
 Consultant Pathologist

*Rekha N*  
**Dr. Rekha Nair, MD**  
 Microbiologist



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Test Report Status	Final	Results	Biological Reference Interval	Units
PUS CELL (WBC'S)		2-3	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION				
EPITHELIAL CELLS		3-5	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION				
CASTS		NOT DETECTED		
METHOD : MICROSCOPIC EXAMINATION				
CRYSTALS		NOT DETECTED		
METHOD : MICROSCOPIC EXAMINATION				
BACTERIA		NOT DETECTED	NOT DETECTED	
METHOD : MICROSCOPIC EXAMINATION				
YEAST		NOT DETECTED	NOT DETECTED	
METHOD : MICROSCOPIC EXAMINATION				
REMARKS		URINARY MICROSCOPIC EXAMINATION DONE ON URINARY CENTRIFUGED SEDIMENT.		

Interpretation(s)

\*\*End Of Report\*\*

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Dr. Akta Dubey  
 Consultant Pathologist

Dr. Rekha Nair, MD  
 Microbiologist



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**SPECIALISED CHEMISTRY - HORMONE**

<b>THYROID PANEL, SERUM</b>				
T3	105.70	Non-Pregnant Women 80.0 - 200.0 Pregnant Women 1st Trimester:105.0 - 230.0 2nd Trimester:129.0 - 262.0 3rd Trimester:135.0 - 262.0	ng/dL	
METHOD : ELECTROCHEMILUMINESCENCE, COMPETITIVE IMMUNOASSAY				
T4	10.21	Non-Pregnant Women 5.10 - 14.10 Pregnant Women 1st Trimester: 7.33 - 14.80 2nd Trimester: 7.93 - 16.10 3rd Trimester: 6.95 - 15.70	µg/dL	
METHOD : ELECTROCHEMILUMINESCENCE, COMPETITIVE IMMUNOASSAY				
TSH (ULTRASENSITIVE)	1.690	0.270 - 4.200	µIU/mL	
METHOD : ELECTROCHEMILUMINESCENCE, COMPETITIVE IMMUNOASSAY				

**Interpretation(s)**

**\*\*End Of Report\*\***  
 Please visit [www.sriworld.com](http://www.sriworld.com) for related Test Information for this accession

*Signature*  
786

**Dr. Swapnil Sirmukaddam**  
 Consultant Pathologist



View Details



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**PERFORMED AT :**  
 SRL Ltd  
 BHOOMI TOWER, 1ST FLOOR, HALL NO.1, PLOT NO.28 SECTOR 4, KHARGHAR  
 NAVI MUMBAI, 410210  
 MAHARASHTRA, INDIA  
 Tel : 9111591115,  
 CIN - U74899PB1995PLC045956

**Patient Ref. No. 22000000838134**



<b>PATIENT NAME : MRS.AARTI .</b>		<b>REF. DOCTOR :</b>	
<b>CODE/NAME &amp; ADDRESS :</b> C000045507 - FORTIS FORTIS VASHI-CHC -SPLZD FORTIS HOSPITAL # VASHI, MUMBAI 440001	<b>ACCESSION NO :</b> 0022WD000183	<b>AGE/SEX :</b> 28 Years Female	<b>DRAWN :</b> 01/04/2023 13:28:00
	<b>PATIENT ID :</b> FH.11008739	<b>RECEIVED :</b> 01/04/2023 13:32:58	<b>REPORTED :</b> 01/04/2023 14:58:36
	<b>CLIENT PATIENT ID:</b> UID:11008739		
	<b>ABHA NO :</b>		

**CLINICAL INFORMATION :**  
 UID:11008739 REQNO-1454619  
 CORP-OPD  
 BILLNO-150123OPCR019077  
 BILLNO-150123OPCR019077

Test Report Status	Results	Biological Reference Interval	Units
Final			

**BIOCHEMISTRY**

<b>GLUCOSE, POST-PRANDIAL, PLASMA</b>			
PPBS(POST PRANDIAL BLOOD SUGAR)	91	70 - 139	mg/dL
METHOD : HEXOKINASE			

**Comments**  
 NOTE: - RECHECKED FOR POST PRANDIAL PLASMA GLUCOSE VALUE. TO BE CORRELATE WITH CLINICAL, DIETETIC AND THERAPEUTIC HISTORY.

**Interpretation(s)**  
 GLUCOSE, POST-PRANDIAL, PLASMA-High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glycosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc.Additional test HbA1c  
**\*\*End Of Report\*\***  
 Please visit [www.srlworld.com](http://www.srlworld.com) for related Test Information for this accession

*Akta Dubey*  
**Dr.Akta Dubey**  
 Consultant Pathologist



View Details



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**PERFORMED AT :**  
 SRL Ltd  
 HIRANANDANI HOSPITAL-VASHI, MINI SEASHORE ROAD, SECTOR 10,  
 NAVI MUMBAI, 400703  
 MAHARASHTRA, INDIA  
 Tel : 022-39199222,022-49723322,  
 CIN - U74899PB1995PLC045956  
 Email : -



Patient Ref. No. 22000000838200

<b>PATIENT NAME : MRS.AARTI .</b>		<b>REF. DOCTOR :</b>	
<b>CODE/NAME &amp; ADDRESS :</b> C000045507 - FORTIS FORTIS VASHI-CHC -SPLZD FORTIS HOSPITAL # VASHI, MUMBAI 440001	<b>ACCESSION NO :</b> 0022WD000243	<b>AGE/SEX :</b> 28 Years Female	<b>DRAWN :</b> 01/04/2023 15:44:00
	<b>PATIENT ID :</b> FH.11008739	<b>RECEIVED :</b> 01/04/2023 15:53:56	<b>REPORTED :</b> 03/04/2023 10:12:21
	<b>CLIENT PATIENT ID:</b> UID:11008739		
	<b>ABHA NO :</b>		

**CLINICAL INFORMATION :**

UID:11008739 REQNO-1454619  
CORP-OPD  
BILLNO-150123OPCR019077  
BILLNO-150123OPCR019077

<b>Test Report Status</b> <u>Final</u>	<b>Units</b>
--	--------------

**CYTOLOGY**

**PAPANICOLAOU SMEAR**

**PAPANICOLAOU SMEAR**

TEST METHOD

CONVENTIONAL GYNEC CYTOLOGY

SPECIMEN TYPE

TWO UNSTAINED CERVICAL SMEARS RECEIVED

REPORTING SYSTEM

2014 BETHESDA SYSTEM FOR REPORTING CERVICAL CYTOLOGY

SPECIMEN ADEQUACY

SATISFACTORY

METHOD : MICROSCOPIC EXAMINATION

MICROSCOPY

SMEARS STUDIED SHOW SUPERFICIAL SQUAMOUS CELLS, INTERMEDIATE SQUAMOUS CELLS, OCCASIONAL SQUAMOUS METAPLASTIC CELLS, OCCASIONAL CLUSTERS OF ENDOCERVICAL CELLS IN THE BACKGROUND OF MODERATE POLYMORPHS.

INTERPRETATION / RESULT

NEGATIVE FOR INTRAEPITHELIAL LESION OR MALIGNANCY

**Comments**

PLEASE NOTE PAPANICOLAOU SMEAR STUDY IS A SCREENING PROCEDURE FOR CERVICAL CANCER WITH INHERENT FALSE NEGATIVE RESULTS, HENCE SHOULD BE INTERPRETED WITH CAUTION.

NO CYTOLOGICAL EVIDENCE OF HPV INFECTION IN THE SMEARS STUDIED.

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

Please visit [www.srlworld.com](http://www.srlworld.com) for related Test Information for this accession



**Dr.Akta Dubey**  
Consultant Pathologist



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**PERFORMED AT :**

SRL Ltd  
HIRANANDANI HOSPITAL-VASHI, MINI SEASHORE ROAD, SECTOR 10,  
NAVI MUMBAI, 400703  
MAHARASHTRA, INDIA  
Tel : 022-39199222,022-49723322,  
CIN - U74899PB1995PLC045956  
Email : -



**Patient Ref. No. 2200000838260**

Rate 87 . Sinus rhythm.....normal P axis, V-rate 50-99  
 . Borderline T abnormalities, anterior leads.....T flat or neg, V2-V4  
 . Prolonged QT interval.....QTc >495ms

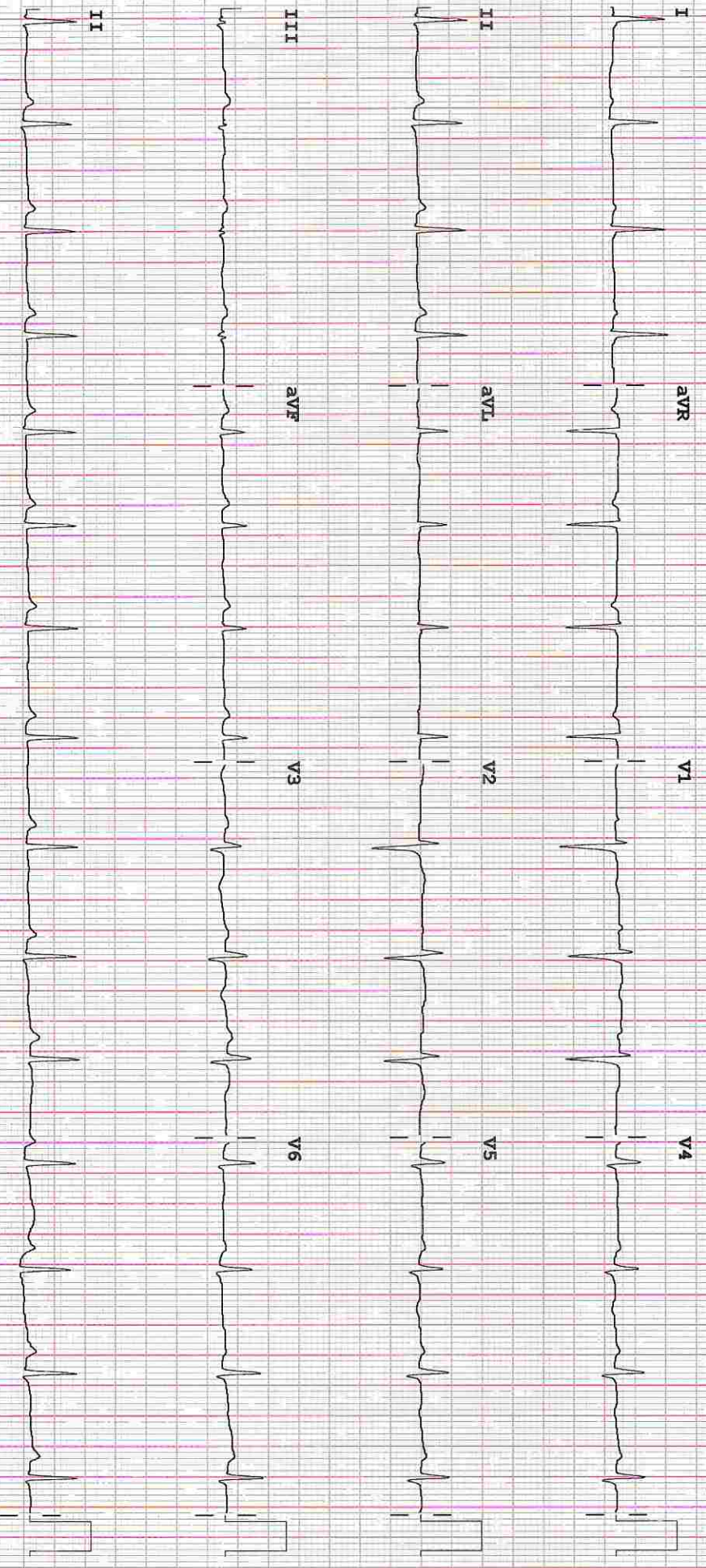
PR 173  
 QRSD 67  
 QT 436  
 QTc 525

--AXIS--  
 P 71  
 QRS 29  
 T -58

- ABNORMAL ECG -

12 Lead; Standard Placement

Unconfirmed Diagnosis



Device: Speed: 25 mm/sec Limb: 10 mm/mV Chest: 10.0 mm/mV

F 50~ 0.50-100 Hz W 100B CL P?

HC  
 NSR  
 Biphasic T in V2-V4  
 for



(For Billing/Reports &amp; Discharge Summary only)

## DEPARTMENT OF NIC

Date: 01/Apr/2023

Name: Mrs. Aarti .

UHID | Episode No : 11008739 | 19201/23/1501

Age | Sex: 28 YEAR(S) | Female

Order No | Order Date: 1501/PN/OP/2304/40283 | 01-Apr-2023

Order Station : FO-OPD

Admitted On | Reporting Date : 01-Apr-2023 13:51:04

Bed Name :

Order Doctor Name : Dr.SELF .

## ECHOCARDIOGRAPHY TRANSTHORACIC

**FINDINGS:**

- No left ventricle regional wall motion abnormality at rest.
- Normal left ventricle systolic function. LVEF = 60%.
- No left ventricle diastolic dysfunction.
- No left ventricle Hypertrophy. No left ventricle dilatation.
- Structurally normal valves.
- No mitral regurgitation.
- No aortic regurgitation. No aortic stenosis.
- No tricuspid regurgitation. No pulmonary hypertension.
- Intact IAS and IVS.
- No left ventricle clot/vegetation/pericardial effusion.
- Normal right atrium and right ventricle dimensions.
- Normal left atrium and left ventricle dimension.
- Normal right ventricle systolic function. No hepatic congestion

**M-MODE MEASUREMENTS:**

LA	31	mm
AO Root	25	mm
AO CUSP SEP	17	mm
LVID (s)	27	mm
LVID (d)	40	mm
IVS (d)	10	mm
LVPW (d)	09	mm
RVID (d)	29	mm
RA	31	mm
LVEF	60	%





(For Billing/Reports &amp; Discharge Summary only)

## DEPARTMENT OF NIC

Date: 01/Apr/2023

Name: Mrs. Aarti .

UHID | Episode No : 11008739 | 19201/23/1501

Age | Sex: 28 YEAR(S) | Female

Order No | Order Date: 1501/PN/OP/2304/40283 | 01-Apr-2023

Order Station : FO-OPD

Admitted On | Reporting Date : 01-Apr-2023 13:51:04

Bed Name :

Order Doctor Name : Dr.SELF .

**DOPPLER STUDY:**

E WAVE VELOCITY: 0.6 m/sec.

A WAVE VELOCITY:0.5 m/sec

E/A RATIO:1.2

	PEAK (mmHg)	MEAN (mmHg)	V max (m/sec)	GRADE OF REGURGITATION
MITRAL VALVE	N			Nil
AORTIC VALVE	06			Nil
TRICUSPID VALVE	N			Nil
PULMONARY VALVE	2.0			Nil

**Final Impression :**

- Normal 2 Dimensional and colour doppler echocardiography study.

DR. PRASHANT PAWAR  
 DNB(MED), DNB ( CARDIOLOGY)



DEPARTMENT OF RADIOLOGY

Date: 01/Apr/2023

Name: Mrs. Aarti .

Age | Sex: 28 YEAR(S) | Female

Order Station : FO-OPD

Bed Name :

UHID | Episode No : 11008739 | 19201/23/1501

Order No | Order Date: 1501/PN/OP/2304/40283 | 01-Apr-2023

Admitted On | Reporting Date : 01-Apr-2023 13:42:34

Order Doctor Name : Dr.SELF .

X-RAY-CHEST- PA

**Findings:**

Both lung fields are clear.

The cardiac shadow appears within normal limits.

Trachea and major bronchi appear normal.

Both costophrenic angles are well maintained.

Bony thorax appears unremarkable.

*Aditya*

**DR. ADITYA NALAWADE**

**M.D. (Radiologist)**



(For Billing/Reports & Discharge Summary only)

DEPARTMENT OF RADIOLOGY

Date: 01/Apr/2023

Name: Mrs. Aarti .

UHID | Episode No : 11008739 | 19201/23/1501

Age | Sex: 28 YEAR(S) | Female

Order No | Order Date: 1501/PN/OP/2304/40283 | 01-Apr-2023

Order Station : FO-OPD

Admitted On | Reporting Date : 01-Apr-2023 13:38:48

Bed Name :

Order Doctor Name : Dr.SELF .

US-WHOLE ABDOMEN

**LIVER** is normal in size and shows mildly raised echogenicity. No IHBR dilatation. No focal lesion is seen in liver. Portal vein appears normal in caliber.

**GALL BLADDER** is physiologically distended. Gall bladder reveals normal wall thickness. No evidence of calculi in gall bladder. No evidence of pericholecystic collection. **CBD** appears normal in caliber.

**SPLEEN** is normal in size and echogenicity.

**BOTH KIDNEYS** are normal in size and echogenicity. The central sinus complex is normal. No evidence of calculi/hydronephrosis. Right kidney measures 9.6 x 3.8 cm. Left kidney measures 9.9 x 4.2 cm.

**PANCREAS** is normal in size and morphology. No evidence of peripancreatic collection.

**URINARY BLADDER** is normal in capacity and contour. Bladder wall is normal in thickness. No evidence of intravesical calculi.

**UTERUS** is normal in size, measuring 5.5 x 4.6 x 3.0 cm. Endometrium measures 5.5 mm in thickness.

Both ovaries are normal. Right ovary measures 2.5 x 1.4 cm. Left ovary measures 2.1 x 1.4 cm.

No evidence of ascites.

**Impression:**

- Grade I fatty infiltration of liver.

  
DR. ADITYA NALAWADE  
M.D. (Radiologist)