



Hiranandani  
HOSPITAL

(A Fortis Network Hospital)

Hiranandani Fortis Hospital  
Mini Seashore Road,  
Sector 10 - A, Vashi,  
Navi Mumbai - 400 703.  
Tel. : +91-22-3919 9222  
Fax : +91-22-3919 9220/21  
Email : vashi@vashihospital.com

## BMI CHART

Date: 5/3/24

Name: Amrita Bhorh Age: 33 yrs

Sex: M / F

BP: 110/80 mmHg Height (cms): 152 cm Weight(kgs): 53.7 kg BMI: \_\_\_\_\_

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							
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	
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		
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		
5'4" - 162.5	17	18	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37		
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		
5'6" - 167.6	16	17	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36		
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		
5'8" - 172.7	15	16	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35		
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		
5'10" - 177.8	14	15	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34		
5'11" - 180.3	14	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34		
6'0" - 182.8	13	14	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33		
6'1" - 185.4	13	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33		
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		
6'3" - 190.5	12	13	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32		
6'4" - 193.0	12	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32		

Doctors Notes:

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Signature



UHID	13012860	Date	05/03/2024	
Name	Mrs Amrita Bharti	Sex	F	Age 35
OPD	PAP	Health Check-Up		

SIB Dr. Shefali

Drug allergy: → No  
 Sys illness: → No

35/F P2L2 C Preu FIND

No fresh complaint  
 No Comorbidities.

P1L4 (F) → 4 1/2 yrs } FIND  
 P2L2 (M) → 6 months }

LMP → Lactational amenorrhoea.

No significant family history

PS → (x) / ug → (H)  
 Pap Smear taken

B/C breast → Soft  
 no lump / Swelling  
 Palpable

Adv  
 — Pap Smear taken

— Pap Smear every 3yrsly

— Counselled about HPV Vaccine [0, 2, 6 month]

— Flu & reports



UHID	13012860	Date	05/03/2024		
Name	Mrs Amrita Bharti	Sex	F	Age	35
OPD	Dental	Health Check-Up			

3/E - Stains +  
- Calculus +  
- proclined anterior

Drug allergy:  
Sys illness:

Treatment

Afd - Scaling Grade I

Dr. Supti

Hiranandani Healthcare Pvt. Ltd.  
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Board Line: 022 - 39199222 | Fax: 022 - 39199220  
Emergency: 022 - 39199100 | Ambulance: 1255  
For Appointment: 022 - 39199222 | Health Checkup: 022 - 39199300  
www.fortishhealthcare.com |  
DIN : U85100MH2005PTC154823  
GST IN: 27AABCH5894D1ZG | PAN NO: AABCH5894D



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Fortis Network Hospital

UHID	13012865	Date	05/01/2024		
Name	Mr.Rajendra Kumar	Sex	Male	Age	36
OPD	Dental 12	Health Check Up			

Drug allergy:  
Sys illness:

B/E - Stains +  
Calculus +

Treatment

Ad - Sealing Grade I

Dr. Trupti



<b>UHID</b>	<b>13012860</b>	<b>Date</b>	<b>05/03/2024</b>		
<b>Name</b>	<b>Mrs Amrita Bharti</b>	<b>Sex</b>	<b>F</b>	<b>Age</b>	<b>35</b>
<b>OPD</b>	<b>Ophthal</b>	<b>Health Check-Up</b>			

Drug allergy: → Not known.  
 Sys illness: → NO  
Habit: → NO.

Chc. No.

HC No.

Unilk → RG 6/60  
 → LG 6/60 [ Phure ]  
 NK → NG

Ref → RG - 2.25 / -0.75 X 90° 6/6.  
 → LG - 2.25 / -1.00 X 90° 6/6.

M → RG NG  
 → LG NG

JOP → RE → 14.8  
 → LE → 14.7

Abmp  
 Saw us P.M.P.

**PATIENT NAME : MRS.AMRITA BHARTI**

**REF. DOCTOR :**

**CODE/NAME & ADDRESS : C000045507**

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

**ACCESSION NO : 0022XC000879**

**PATIENT ID : FH.13012860**

**CLIENT PATIENT ID: UID:13012860**

**ABHA NO :**

**AGE/SEX :35 Years Female**

**DRAWN :05/03/2024 09:07:00**

**RECEIVED : 05/03/2024 09:09:10**

**REPORTED :05/03/2024 14:39:00**

**CLINICAL INFORMATION :**

UID:13012860 REQNO-1671444  
 CORP-OPD  
 BILLNO-150124OPCR012846  
 BILLNO-150124OPCR012846

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) METHOD : SLS METHOD	<b>11.7 Low</b>	12.0 - 15.0	g/dL
RED BLOOD CELL (RBC) COUNT METHOD : HYDRODYNAMIC FOCUSING	4.56	3.8 - 4.8	mil/ $\mu$ L
WHITE BLOOD CELL (WBC) COUNT METHOD : FLUORESCENCE FLOW CYTOMETRY	6.82	4.0 - 10.0	thou/ $\mu$ L
PLATELET COUNT METHOD : HYDRODYNAMIC FOCUSING BY DC DETECTION	307	150 - 410	thou/ $\mu$ L

**RBC AND PLATELET INDICES**

HEMATOCRIT (PCV) METHOD : CUMULATIVE PULSE HEIGHT DETECTION METHOD	38.8	36.0 - 46.0	%
MEAN CORPUSCULAR VOLUME (MCV) METHOD : CALCULATED PARAMETER	85.1	83.0 - 101.0	fL
MEAN CORPUSCULAR HEMOGLOBIN (MCH) METHOD : CALCULATED PARAMETER	<b>25.7 Low</b>	27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION(MCHC) METHOD : CALCULATED PARAMETER	<b>30.2 Low</b>	31.5 - 34.5	g/dL
RED CELL DISTRIBUTION WIDTH (RDW) METHOD : CALCULATED PARAMETER	13.0	11.6 - 14.0	%
MENTZER INDEX METHOD : CALCULATED PARAMETER	18.7		
MEAN PLATELET VOLUME (MPV) METHOD : CALCULATED PARAMETER	<b>11.0 High</b>	6.8 - 10.9	fL

**WBC DIFFERENTIAL COUNT**

**Dr. Akshay Dhotre, MD**  
 (Reg,no. MMC 2019/09/6377)  
 Consultant Pathologist



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NEUTROPHILS		66	40.0 - 80.0	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING				
LYMPHOCYTES		26	20.0 - 40.0	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING				
MONOCYTES		6	2.0 - 10.0	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING				
EOSINOPHILS		2	1 - 6	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING				
BASOPHILS		0	0 - 2	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING				
ABSOLUTE NEUTROPHIL COUNT		4.50	2.0 - 7.0	thou/ $\mu$ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE LYMPHOCYTE COUNT		1.77	1.0 - 3.0	thou/ $\mu$ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE MONOCYTE COUNT		0.41	0.2 - 1.0	thou/ $\mu$ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE EOSINOPHIL COUNT		0.14	0.02 - 0.50	thou/ $\mu$ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE BASOPHIL COUNT		<b>0.00 Low</b>	0.02 - 0.10	thou/ $\mu$ L
METHOD : CALCULATED PARAMETER				
NEUTROPHIL LYMPHOCYTE RATIO (NLR)		2.5		
METHOD : CALCULATED				

**MORPHOLOGY**

**RBC** MILD HYPOCHROMASIA, NORMOCYTIC  
 METHOD : MICROSCOPIC EXAMINATION  
**WBC** NORMAL MORPHOLOGY  
 METHOD : MICROSCOPIC EXAMINATION  
**PLATELETS** ADEQUATE  
 METHOD : MICROSCOPIC EXAMINATION

**Dr. Akshay Dhotre, MD**  
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**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.

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), EDTA BLOOD**

E.S.R	17	0 - 20	mm at 1 hr
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METHOD : WESTEREGREN METHOD

**GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD**

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)	%
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METHOD : HB VARIANT (HPLC)

ESTIMATED AVERAGE GLUCOSE(EAG)	108.3	< 116.0	mg/dL
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METHOD : CALCULATED PARAMETER

**Interpretation(s)**

**ERYTHROCYTE SEDIMENTATION RATE (ESR), EDTA 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(62 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,(SickleCells,spherocytes),Microcytosis, Low fibrinogen, Very high WBC counts, Drugs(Quinine, salicylates)

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**REFERENCE :**

1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition;2. Paediatric reference intervals. AACC Press, 7th edition. Edited by S. Soldin;3. The reference for the adult reference range is "Practical Haematology by Dacie and Lewis,10th edition. GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD-Used For:

1. Evaluating the long-term control of blood glucose concentrations in diabetic patients.
2. Diagnosing diabetes.
3. 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 patients metabolic control has remained continuously within the target range.

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

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

1. 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.
- 2.Vitamin C & E are reported to falsely lower test results.(possibly by inhibiting glycation of hemoglobin.
3. Iron deficiency anemia is reported to increase test results. Hypertriglyceridemia,uremia, hyperbilirubinemia, chronic alcoholism,chronic ingestion of salicylates & opiates addiction are reported to interfere with some assay methods,falsely increasing results.
4. 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 paltform (Boronate affinity chromatography) is recommended for testing of HbA1c.Abnormal Hemoglobin electrophoresis (HPLC method) is recommended for detecting a hemoglobinopathy

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**IMMUNOHAEMATOLOGY**

**ABO GROUP & RH TYPE, EDTA WHOLE BLOOD**

<b>ABO GROUP</b> METHOD : TUBE AGGLUTINATION	<b>TYPE B</b>
<b>RH TYPE</b> METHOD : TUBE AGGLUTINATION	<b>POSITIVE</b>

**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.

*Akshay*  
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## BIOCHEMISTRY


## LIVER FUNCTION PROFILE, SERUM

BILIRUBIN, TOTAL METHOD : JENDRASSIK AND GROFF	0.34	0.2 - 1.0	mg/dL
BILIRUBIN, DIRECT METHOD : JENDRASSIK AND GROFF	0.07	0.0 - 0.2	mg/dL
BILIRUBIN, INDIRECT METHOD : CALCULATED PARAMETER	0.27	0.1 - 1.0	mg/dL
TOTAL PROTEIN METHOD : BIURET	7.4	6.4 - 8.2	g/dL
ALBUMIN METHOD : BCP DYE BINDING	3.9	3.4 - 5.0	g/dL
GLOBULIN METHOD : CALCULATED PARAMETER	3.5	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO METHOD : CALCULATED PARAMETER	1.1	1.0 - 2.1	RATIO
ASPARTATE AMINOTRANSFERASE(AST/SGOT) METHOD : UV WITH P5P	14 Low	15 - 37	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT) METHOD : UV WITH P5P	25	< 34.0	U/L
ALKALINE PHOSPHATASE METHOD : PNPP-ANP	132 High	30 - 120	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT) METHOD : GAMMA GLUTAMYL CARBOXY 4NITROANILIDE	19	5 - 55	U/L
LACTATE DEHYDROGENASE METHOD : LACTATE -PYRUVATE	138	81 - 234	U/L

## GLUCOSE FASTING, FLUORIDE PLASMA

FBS (FASTING BLOOD SUGAR) METHOD : HEXOKINASE	87	Normal : < 100 Pre-diabetes: 100-125 Diabetes: >=126	mg/dL
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Dr. Akshay Dhotre, MD  
(Reg.no. MMC 2019/09/6377)  
Consultant Pathologist



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Email : -



Patient Ref. No. 2200000906786

**PATIENT NAME : MRS.AMRITA BHARTI**

**REF. DOCTOR :**

**CODE/NAME & ADDRESS : C000045507**

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

**ACCESSION NO : 0022XC000879**

**PATIENT ID : FH.13012860**

**CLIENT PATIENT ID: UID:13012860**

**ABHA NO :**

**AGE/SEX : 35 Years Female**

**DRAWN : 05/03/2024 09:07:00**

**RECEIVED : 05/03/2024 09:09:10**

**REPORTED : 05/03/2024 14:39:00**

**CLINICAL INFORMATION :**

UID:13012860 REQNO-1671444  
CORP-OPD  
BILLNO-150124OPCR012846  
BILLNO-150124OPCR012846

Test Report Status	Final	Results	Biological Reference Interval	Units
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**KIDNEY PANEL - 1**

**BLOOD UREA NITROGEN (BUN), SERUM**

**BLOOD UREA NITROGEN** 8 6 - 20 mg/dL  
METHOD : UREASE - UV

**CREATININE EGFR- EPI**

**CREATININE** 0.49 Low 0.60 - 1.10 mg/dL  
METHOD : ALKALINE PICRATE KINETIC JAFFES

**AGE** 35 years

**GLOMERULAR FILTRATION RATE (FEMALE)** 125.97 Refer Interpretation Below mL/min/1.73m2  
METHOD : CALCULATED PARAMETER

**BUN/CREAT RATIO**

**BUN/CREAT RATIO** 16.33 High 5.00 - 15.00  
METHOD : CALCULATED PARAMETER

**URIC ACID, SERUM**

**URIC ACID** 3.6 2.6 - 6.0 mg/dL  
METHOD : URICASE UV

**TOTAL PROTEIN, SERUM**

**TOTAL PROTEIN** 7.4 6.4 - 8.2 g/dL  
METHOD : BIURET

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## ALBUMIN, SERUM

ALBUMIN

3.9

3.4 - 5.0

g/dL

METHOD : BCP DYE BINDING

## GLOBULIN

GLOBULIN

3.5

2.0 - 4.1

g/dL

METHOD : CALCULATED PARAMETER

## ELECTROLYTES (NA/K/CL), SERUM

SODIUM, SERUM

140

136 - 145

mmol/L

METHOD : ISE INDIRECT

POTASSIUM, SERUM

4.37

3.50 - 5.10

mmol/L

METHOD : ISE INDIRECT

CHLORIDE, SERUM

104

98 - 107

mmol/L

METHOD : ISE INDIRECT

## Interpretation(s)

## Interpretation(s)

## LIVER FUNCTION PROFILE, SERUM-

**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.



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**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.

**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, sulfonyleureas, 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.

**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--** Kidney disease outcomes quality initiative (KDOQI) guidelines state that estimation of GFR is the best overall indices of the Kidney function.

- It gives a rough measure of number of functioning nephrons. Reduction in GFR implies progression of underlying disease.

- The GFR is a calculation based on serum creatinine test.

- Creatinine is mainly derived from the metabolism of creatine in muscle, and its generation is proportional to the total muscle mass. As a result, mean creatinine generation is higher in men than in women, in younger than in older individuals, and in blacks than in whites.

- Creatinine is filtered from the blood by the kidneys and excreted into urine at a relatively steady rate.

- When kidney function is compromised, excretion of creatinine decreases with a consequent increase in blood creatinine levels. With the creatinine test, a reasonable estimate of the actual GFR can be determined.

- This equation takes into account several factors that impact creatinine production, including age, gender, and race.

- CKD EPI (Chronic kidney disease epidemiology collaboration) equation performed better than MDRD equation especially when GFR is high (>60 ml/min per 1.73m2).. This formula has less bias and greater accuracy which helps in early diagnosis and also reduces the rate of false positive diagnosis of CKD.

**References:**

National Kidney Foundation (NKF) and the American Society of Nephrology (ASN).

Estimated GFR Calculated Using the CKD-EPI equation-<https://testguide.labmed.uw.edu/guideline/egfr>

Ghuman JK, et al. Impact of Removing Race Variable on CKD Classification Using the Creatinine-Based 2021 CKD-EPI Equation. Kidney Med 2022, 4:100471. 35756325

Harrison's Principle of Internal Medicine, 21st ed. pg 62 and 334

**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.

**Dr. Akshay Dhotre, MD**  
**(Reg.no. MMC 2019/09/6377)**  
**Consultant Pathologist**



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**Patient Ref. No. 2200000906786**

PATIENT NAME : MRS.AMRITA BHARTI

REF. DOCTOR :

CODE/NAME &amp; ADDRESS : C000045507

ACCESSION NO : 0022XC000879

AGE/SEX : 35 Years Female

FORTIS VASHI-CHC -SPLZD  
 FORTIS HOSPITAL # VASHI,  
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**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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**BIOCHEMISTRY - LIPID**

**LIPID PROFILE, SERUM**

CHOLESTEROL, TOTAL	149	< 200 Desirable 200 - 239 Borderline High >= 240 High	mg/dL
METHOD : ENZYMATIC/COLORIMETRIC, CHOLESTEROL OXIDASE, ESTERASE, PEROXIDASE			
TRIGLYCERIDES	54	< 150 Normal 150 - 199 Borderline High 200 - 499 High >=500 Very High	mg/dL
METHOD : ENZYMATIC ASSAY			
HDL CHOLESTEROL	54	< 40 Low >=60 High	mg/dL
METHOD : DIRECT MEASURE - PEG			
LDL CHOLESTEROL, DIRECT	81	< 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			
NON HDL CHOLESTEROL	95	Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL
METHOD : CALCULATED PARAMETER			
VERY LOW DENSITY LIPOPROTEIN	10.8	<= 30.0	mg/dL
METHOD : CALCULATED PARAMETER			
CHOL/HDL RATIO	2.8 Low	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			

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Patient Ref. No. 22000000906786



MC-5837

**PATIENT NAME : MRS.AMRITA BHARTI**

**REF. DOCTOR :**

**CODE/NAME & ADDRESS : C000045507**

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FORTIS HOSPITAL # VASHI,  
MUMBAI 440001

ACCESSION NO : **0022XC000879**

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LDL/HDL RATIO	1.5	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk		
METHOD : CALCULATED PARAMETER				

**Interpretation(s)**

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## CLINICAL PATH - URINALYSIS

## KIDNEY PANEL - 1

## PHYSICAL EXAMINATION, URINE

COLOR	PALE YELLOW
METHOD : PHYSICAL	
APPEARANCE	SLIGHTLY HAZY
METHOD : VISUAL	

## CHEMICAL EXAMINATION, URINE

PH	6.0	4.7 - 7.5
METHOD : REFLECTANCE SPECTROPHOTOMETRY- DOUBLE INDICATOR METHOD		
SPECIFIC GRAVITY	<=1.005	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		

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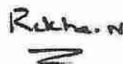
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<b>MICROSCOPIC EXAMINATION, URINE</b>				
RED BLOOD CELLS		NOT DETECTED	NOT DETECTED	/HPF
METHOD : MICROSCOPIC EXAMINATION				
PUS CELL (WBC'S)		5-7	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION				
EPITHELIAL CELLS		8-10	0-5	/HPF
METHOD : MICROSCOPIC 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)**

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**CODE/NAME & ADDRESS : C000045507**

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

**ACCESSION NO : 0022XC000879**

PATIENT ID : FH.13012860  
 CLIENT PATIENT ID: UID:13012860  
 ABHA NO :

AGE/SEX :35 Years Female

DRAWN :05/03/2024 09:07:00  
 RECEIVED :05/03/2024 09:09:10  
 REPORTED :05/03/2024 14:39:00

**CLINICAL INFORMATION :**

UID:13012860 REQNO-1671444  
 CORP-OPD  
 BILLNO-150124OPCR012846  
 BILLNO-150124OPCR012846

Test Report Status	Final	Results	Biological Reference Interval	Units
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**SPECIALISED CHEMISTRY - HORMONE**

**THYROID PANEL, SERUM**

T3 107.8 Non-Pregnant Women ng/dL  
 80.0 - 200.0  
 Pregnant Women  
 1st Trimester:105.0 - 230.0  
 2nd Trimester:129.0 - 262.0  
 3rd Trimester:135.0 - 262.0

METHOD : ELECTROCHEMILUMINESCENCE IMMUNOASSAY, COMPETITIVE PRINCIPLE

T4 6.03 Non-Pregnant Women µg/dL  
 5.10 - 14.10  
 Pregnant Women  
 1st Trimester: 7.33 - 14.80  
 2nd Trimester: 7.93 - 16.10  
 3rd Trimester: 6.95 - 15.70

METHOD : ELECTROCHEMILUMINESCENCE IMMUNOASSAY, COMPETITIVE PRINCIPLE

TSH (ULTRASENSITIVE) 3.470 Non Pregnant Women µIU/mL  
 0.27 - 4.20  
 Pregnant Women (As per  
 American Thyroid Association)  
 1st Trimester 0.100 - 2.500  
 2nd Trimester 0.200 - 3.000  
 3rd Trimester 0.300 - 3.000

METHOD : ELECTROCHEMILUMINESCENCE,SANDWICH IMMUNOASSAY

**Interpretation(s)**

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

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

**Dr. Akshay Dhotre, MD**  
 (Reg.no. MMC 2019/09/6377)  
 Consultant Pathologist



View Details



View Report

**PERFORMED AT :**

Agilus Diagnostics 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. 22000000906786

PATIENT NAME : MRS.AMRITA BHARTI

REF. DOCTOR :

CODE/NAME &amp; ADDRESS : C000045507

ACCESSION NO : 0022XC000943

AGE/SEX : 35 Years Female

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

PATIENT ID : FH.13012860

DRAWN : 05/03/2024 12:34:00

CLIENT PATIENT ID: UID:13012860

RECEIVED : 05/03/2024 12:42:05

ABHA NO :

REPORTED : 05/03/2024 13:37:22

## CLINICAL INFORMATION :

UID:13012860 REQNO-1671444  
CORP-OPD  
BILLNO-150124OPCR012846  
BILLNO-150124OPCR012846

Test Report Status	Final	Results	Biological Reference Interval	Units
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## BIOCHEMISTRY

## GLUCOSE, POST-PRANDIAL, PLASMA

PPBS(POST PRANDIAL BLOOD SUGAR)	71	70 - 140	mg/dL
METHOD : HEXOKINASE			

## Comments

NOTE : POST PRANDIAL PLASMA GLUCOSE VALUES. 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 &amp; Insulin treatment, Renal Glycosuria, Glycaemic index &amp; response to food consumed, Alimentary Hypoglycemia, Increased insulin response &amp; sensitivity etc.Additional test HbA1c

\*\*End Of Report\*\*

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


Dr. Akshay Dhotre, MD  
(Reg.no. MMC 2019/09/6377)  
Consultant Pathologist

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Maharashtra, India  
Tel : 022-39199222,022-49723322,  
CIN - U74899PB1995PLC045956  
Email : -



Patient Ref. No. 22000000906850

**PATIENT NAME : MRS.AMRITA BHARTI**

**REF. DOCTOR :**

**CODE/NAME & ADDRESS : C000045507**

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

**ACCESSION NO : 0022XC000979**

**PATIENT ID : FH.13012860**

**CLIENT PATIENT ID: UID:13012860**

**ABHA NO :**

**AGE/SEX : 35 Years Female**

**DRAWN : 05/03/2024 14:33:00**

**RECEIVED : 05/03/2024 14:37:56**

**REPORTED : 06/03/2024 13:15:34**

**CLINICAL INFORMATION :**

UID:13012860 REQNO-1671444  
 CORP-OPD  
 BILLNO-150124OPCR012846  
 BILLNO-150124OPCR012846

<b>Test Report Status</b>	<b>Final</b>	<b>Units</b>
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**CYTOLOGY**

**PAPANICOLAOU SMEAR**

**PAPANICOLAOU SMEAR**

**TEST METHOD**  
**SPECIMEN TYPE**  
**REPORTING SYSTEM**  
**SPECIMEN ADEQUACY**  
 METHOD : MICROSCOPIC EXAMINATION  
**MICROSCOPY**

CONVENTIONAL GYNEC CYTOLOGY  
 TWO UNSTAINED CERVICAL SMEARS RECEIVED  
 2014 BETHESDA SYSTEM FOR REPORTING CERVICAL CYTOLOGY  
 SATISFACTORY

SMEARS STUDIED SHOW SUPERFICIAL SQUAMOUS CELLS,  
 INTERMEDIATE SQUAMOUS CELLS, OCCASIONAL SQUAMOUS  
 METAPLASTIC CELLS, OCCASIONAL CLUSTERS OF ENDOCERVICAL CELLS  
 IN THE BACKGROUND OF PLENTY 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.agilusdiagnostics.com](http://www.agilusdiagnostics.com) for related Test Information for this accession

**Dr. Akshay Dhotre, MD**  
 (Reg.no. MMC 2019/09/6377)  
 Consultant Pathologist



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 Maharashtra, India  
 Tel : 022-39199222,022-49723322,  
 CIN - U74899PB1995PLC045956  
 Email : -



**Patient Ref. No. 22000000906886**

13012860  
35 Years

amrita bharti  
Male

01/01/2024 10:33:00 AM

He

Rate 59 Sinus rhythm.....normal P axis, V-rate 50-99  
 PR 116 Borderline short PR interval.....PR int <120ms  
 QRS 90 Baseline wander in lead(s) V6

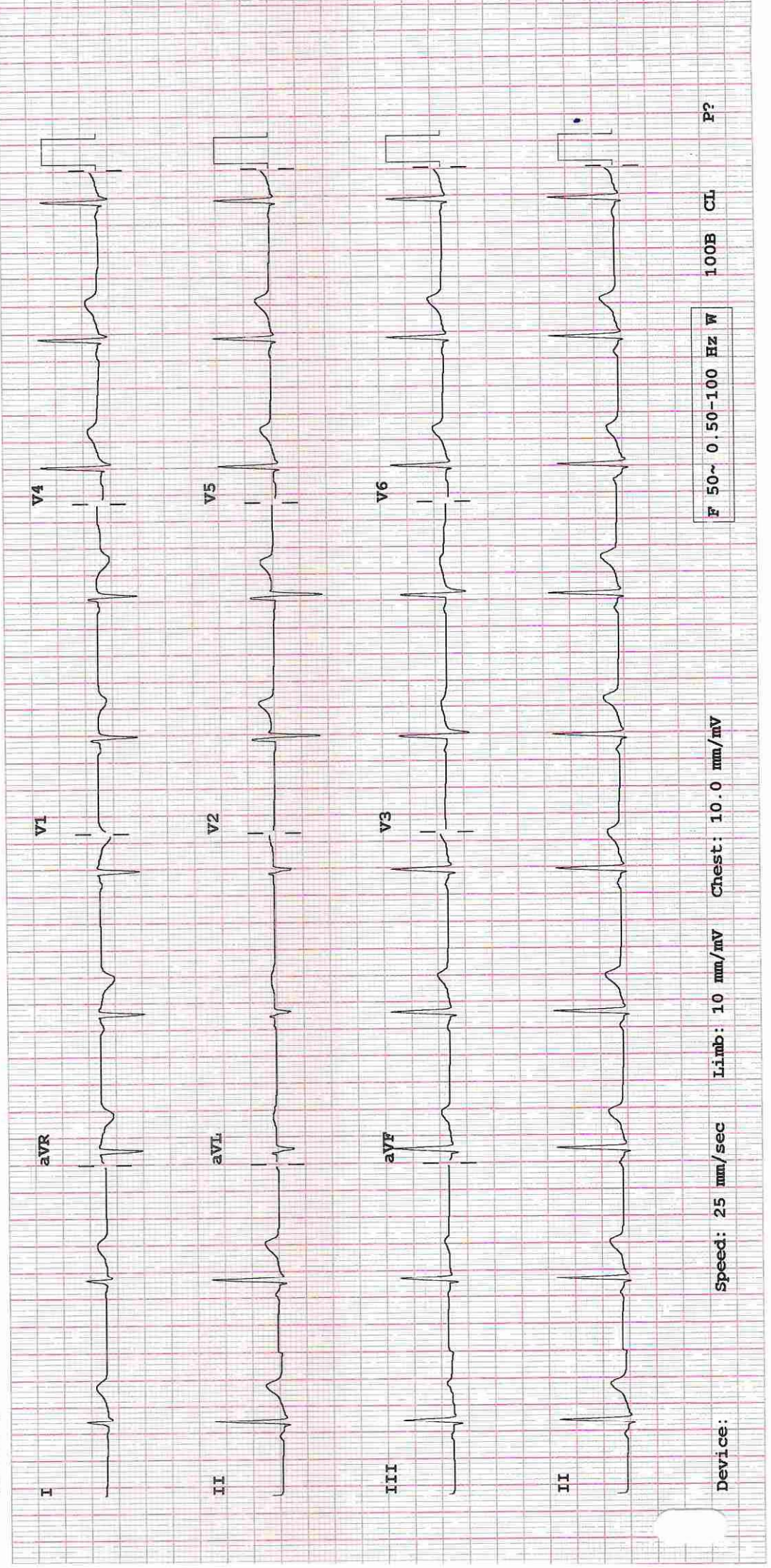
Normal *A*

--AXIS--  
 P 61  
 QRS 73  
 T 42

- OTHERWISE NORMAL ECG -

12 Lead; Standard Placement

Unconfirmed Diagnosis



Speed: 25 mm/sec    Limb: 10 mm/mV    Chest: 10.0 mm/mV    F 50~ 0.50-100 Hz W    100B CL    P?





DEPARTMENT OF NIC

Date: 05/Mar/2024

Name: Mrs. Amrita Bharti  
Age | Sex: 35 YEAR(S) | Female  
Order Station : FO-OPD  
Bed Name :

UHID | Episode No : 13012860 | 13169/24/1501  
Order No | Order Date: 1501/PN/OP/2403/27313 | 05-Mar-2024  
Admitted On | Reporting Date : 05-Mar-2024 18:06:25  
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 e/o raised LVEDP.
- Trivial mitral regurgitation.
- No aortic regurgitation. No aortic stenosis.
- Mild tricuspid regurgitation. Mild pulmonary hypertension.  
PASP = 36 mm of Hg.
- Intact IVS and IAS.
- No left ventricle clot/vegetation/pericardial effusion.
- Normal right atrium and right ventricle dimension.
- Normal left atrium and left ventricle dimension.
- Normal right ventricle systolic function. No hepatic congestion.
- IVC measures 11 mm with normal inspiratory collapse .

**M-MODE MEASUREMENTS:**

LA	24	mm
AO Root	18	mm
AO CUSP SEP	15	mm
LVID (s)	26	mm
LVID (d)	38	mm
IVS (d)	08	mm
LVPW (d)	09	mm
RVID (d)	24	mm
RA	28	mm
LVEF	60	%

Hiranandani Healthcare Pvt. Ltd.  
Mini Sea Shore Road, Sector 10-A, Vashi, Navi Mumbai - 400703.  
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For Appointment: 022 - 39199200 | Health Checkup: 022 - 39199300  
www.fortishealthcare.com | vashi@fortishealthcare.com  
CIN: U85100MH2005PTC 154823  
GST IN : 27AABCH5894D1ZG  
PAN NO : AABCH5894D



DEPARTMENT OF NIC

Date: 05/Mar/2024

Name: Mrs. Amrita Bharti  
Age | Sex: 35 YEAR(S) | Female  
Order Station : FO-OPD  
Bed Name :

UHID | Episode No : 13012860 | 13169/24/1501  
Order No | Order Date: 1501/PN/OP/2403/27313 | 05-Mar-2024  
Admitted On | Reporting Date : 05-Mar-2024 18:06:25  
Order Doctor Name : Dr.SELF.

**DOPPLER STUDY:**

E WAVE VELOCITY: 0.7 m/sec.  
A WAVE VELOCITY: 0.6 m/sec  
E/A RATIO: 1.2

	PEAK (mmHg)	MEAN (mmHg)	V max (m/sec)	GRADE OF REGURGITATION
MITRAL VALVE	N			Trivial
AORTIC VALVE	05			Nil
TRICUSPID VALVE	36			Mild
PULMONARY VALVE	2.0			Nil

**Final Impression :**

- No RWMA.
- Trivial MR and Mild TR. Mild PH.
- Normal LV and RV systolic function.

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

DR. AMIT SINGH,  
MD(MED), DM(CARD)

Hiranandani Healthcare Pvt. Ltd.

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CIN: U85100MH2005PTC 154823

GST IN : 27AABCH5894D1ZG

PAN NO : AABCH5894D



DEPARTMENT OF RADIOLOGY

Date: 05/Mar/2024

Name: Mrs. Amrita Bharti

Age | Sex: 35 YEAR(S) | Female

Order Station : FO-OPD

Bed Name :

UHID | Episode No : 13012860 | 13169/24/1501

Order No | Order Date: 1501/PN/OP/2403/27313 | 05-Mar-2024

Admitted On | Reporting Date : 05-Mar-2024 15:34:53

Order Doctor Name : Dr.SELF .

USG – BOTH BREAST

**Findings:**

Dilated ducts are seen in the left retroareolar region. The dilated ducts have collection (milk) within it, consistent with lactation breast.

Bilateral breast parenchyma appears normal.

No evidence of solid or cystic lesion.

The fibroglandular architecture is well maintained.

Retromammory soft tissues appear normal.

No evidence of axillary lymphadenopathy.

**Impression:**

- Dilated ducts in the left retroareolar region.

**DR. YOGINI SHAH**  
DMRD., DNB. (Radiologist)

Hiranandani Healthcare Pvt. Ltd.

Mini Sea Shore Road, Sector 10-A, Vashi, Navi Mumbai - 400703.

Board Line: 022 - 39199222 | Fax: 022 - 39133220

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CIN: U85100MH2005PTC 154823

GST IN : 27AABCH5894D1ZG

PAN NO : AABCH5894D



DEPARTMENT OF RADIOLOGY

Date: 05/Mar/2024

Name: Mrs. Amrita Bharti

Age | Sex: 35 YEAR(S) | Female

Order Station : FO-OPD

Bed Name :

UHID | Episode No : 13012860 | 13169/24/1501

Order No | Order Date: 1501/PN/OP/2403/27313 | 05-Mar-2024

Admitted On | Reporting Date : 05-Mar-2024 14:16:12

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 appears normal.

Both costophrenic angles are well maintained.

Bony thorax is unremarkable.

**DR. YOGINI SHAH**

**DMRD., DNB. (Radiologist)**



Patient Name	: Amrita Bharti	Patient ID	: 13012860
Sex / Age	: F / 35Y 6M 5D	Accession No.	: PHC.7612606
Modality	: US	Scan DateTime	: 05-03-2024 13:32:24
IPID No	: 13169/24/1501	ReportDatetime	: 05-03-2024 13:45:34

### US – WHOLE ABDOMEN

**LIVER** is normal in size and echogenicity. Intrahepatic portal and biliary systems are normal. No focal lesion is seen in liver. Portal vein is normal.

**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 11.2 x 4.6 cm.

Left kidney measures 10.7 x 4.6 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 mass/calculi.

**UTERUS** is normal in size, measuring 7.3 x 3.3 x 4.4 cm.

Endometrium measures 5.2 mm in thickness.

Both ovaries are normal.

Right ovary measures 3.3 x 2.2 x 2.0 cm, volume 8.1 cc.

Left ovary measures 3.7 x 3.2 x 1.4 cm, volume 8.9 cc.

*Minimal free fluid is seen in Pouch of Douglas.*

### Impression:

- No significant abnormality is detected.

**DR. CHETAN KHADKE**

(MD Radiologist)