

BMI CHART

Date: 10

Name: mm. shalini kumari Age: 32 yrs

Sex: M / F F

BP: 130/90 mmHg Height (cms): 167cm Weight(kgs): 86 kg BMI: _____
 SpO2 - 99%
 pulse - 94b/m

HEIGHT in/cm	WEIGHT lbs		kgs																																																			
	100	105	100	115	120	125	130	135	140	145	150	155	160	165	170	175	180	185	190	195	200	205	210	215																														
	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	41	42																													
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	42																													
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	41	42																												
5'4" - 162.5	17	18	18	19	20	21	22	23	24	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42																										
5'5" - 165.1	16	17	18	19	20	20	21	22	23	24	24	25	26	27	28	29	30	31	31	32	33	34	35	36	37	38	39	40	41	42																								
5'6" - 167.6	16	17	17	18	19	20	21	21	22	23	24	25	25	26	27	28	29	30	30	31	32	33	34	35	36	37	38	39	40	41	42																							
5'7" - 170.1	15	16	17	18	18	19	20	21	22	22	23	24	25	25	26	27	28	29	29	30	31	32	33	34	35	35	36	37	38	39	40	41	42																					
5'8" - 172.7	15	16	16	17	18	19	19	20	21	22	22	23	24	25	25	26	27	28	29	29	30	31	32	33	34	34	35	36	37	38	39	40	41	42																				
5'9" - 176.2	14	15	16	17	18	19	20	21	22	22	23	24	25	25	26	27	28	29	29	30	31	32	33	33	34	35	36	37	38	39	40	41	42																					
5'10" - 177.8	14	15	15	16	17	18	19	20	20	21	22	22	23	24	25	25	26	27	28	28	29	30	31	32	32	33	34	35	36	37	38	39	40	41	42																			
5'11" - 180.3	14	14	15	16	17	18	18	19	20	21	22	23	23	24	25	25	26	27	28	28	29	30	31	31	32	33	34	35	36	37	38	39	40	41	42																			
6'0" - 182.8	13	14	14	15	16	17	18	19	19	20	21	22	23	23	24	25	25	26	27	28	28	29	30	30	31	32	33	34	35	36	37	38	39	40	41	42																		
6'1" - 185.4	13	13	14	15	16	17	18	19	19	20	21	22	23	23	24	25	25	26	27	28	28	29	30	30	31	32	33	34	35	36	37	38	39	40	41	42																		
6'2" - 187.9	12	13	14	15	16	17	18	19	19	20	21	22	23	23	24	25	25	26	27	28	28	29	30	30	31	32	33	34	35	36	37	38	39	40	41	42																		
6'3" - 190.5	12	13	13	14	15	16	17	18	19	19	20	21	22	23	23	24	25	25	26	27	28	28	29	30	30	31	32	33	34	35	36	37	38	39	40	41	42																	
6'4" - 193.0	12	12	13	14	15	16	17	18	19	20	20	21	22	23	23	24	25	25	26	27	28	28	29	30	30	31	32	33	34	35	36	37	38	39	40	41	42																	

Doctors Notes:

Signature



UHID	12649694	Date	16/08/2023		
Name	Mrs. Shalini Kumari	Sex	Female	Age	32
OPD	Pap Smear	Health Check-up			

Drug allergy:
 Sys illness:

Shalini Kumari | 27yrs msk 8yrs P221-04

prev @ ves.

clo: irregular menses.

LMP - 25/7/23. UMP - 6/8/23 → 26/6/23.

LMP₁ - 3/5/23 → 23/5/23.

UMP₂ 14/4/23.

cb clo:

PIH - MAD.

PIA - (S)
 NT

Adv

pls - cb - mild erosion
 vag (H).

follow up
 Report

→ counselling done
 regarding contraception
 & HPV vaccine.
 Adv

Hiranandani Healthcare Pvt. Ltd.
 Mini Sea Shore Road, Sector 10 -A, Vashi, Navi Mumbai - 400703
 Board Line: 022 - 39199222 | Fax: 022 - 39199220
 Emergency: 022 - 39199100 | Ambulance: 1255
 For Appointment: 022 - 39199222 | Health Checkup: 022 - 39199300
 www.fortishealthcare.com |
 CIN : U85100MH2005PTC154823
 GST IN: 27AABCH5894D1ZG | PAN NO: AABCH5894D



Hiranandani
HOSPITAL
 (A Fortis Network Hospital)

UHID	12649694	Date	16/08/2023		
Name	Mrs. Shalini Kumari	Sex	Female	Age	32
OPD	Ophthal 14	Health Check-up			

No n/o spectacle usage

Drug allergy:
 Sys illness:

NO

Unaided VP { 6/6
 6/6

UM < NG
 NG

Ref & Plano - 6/6

CV < 17/17
 17/17 (M)

Da

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Hiranandani
 HOSPITAL

(A Fortis Network Hospital)

UHID	12649694	Date	16/08/2023		
Name	Mrs. Shalini Kumari	Sex	Female	Age	32
OPD	Dental 12	7387696540			
		Health Check-up			

Drug allergy:
 Sys illness:

missing 6/6

stains 88
 calculus 8

Caries 8/7

Treatment

Adv implant 8/6

Adv filling 8/7

Adv Gel prophylaxis

Adv CBCT

Dr. Disha - kaka

PATIENT NAME : MRS.SHALINI KUMARI	REF. DOCTOR :	
CODE/NAME & ADDRESS : C000045507 FORTIS VASHI-CHC -SPLZD FORTIS HOSPITAL # VASHI, MUMBAI 440001	ACCESSION NO : 0022WH003744 PATIENT ID : FH.12649694 CLIENT PATIENT ID: UID:12649694 ABHA NO :	AGE/SEX : 32 Years Female DRAWN : 16/08/2023 09:56:00 RECEIVED : 16/08/2023 09:56:37 REPORTED : 16/08/2023 13:28:56


CLINICAL INFORMATION :

UID:12649694 REQNO-1560004
CORP-OPD
BILLNO-150123OPCR046177
BILLNO-150123OPCR046177

Test Report Status	Results	Biological Reference Interval	Units
Final			

HAEMATOLOGY - CBC**CBC-5, EDTA WHOLE BLOOD****BLOOD COUNTS, EDTA WHOLE BLOOD**

HEMOGLOBIN (HB) METHOD : SLS METHOD	11.8 Low	12.0 - 15.0	g/dL
RED BLOOD CELL (RBC) COUNT METHOD : HYDRODYNAMIC FOCUSING	5.02 High	3.8 - 4.8	mil/ μ L
WHITE BLOOD CELL (WBC) COUNT METHOD : FLUORESCENCE FLOW CYTOMETRY	5.56	4.0 - 10.0	thou/ μ L
PLATELET COUNT METHOD : HYDRODYNAMIC FOCUSING BY DC DETECTION	135 Low	150 - 410	thou/ μ L
RBC AND PLATELET INDICES			
HEMATOCRIT (PCV) METHOD : CUMULATIVE PULSE HEIGHT DETECTION METHOD	38.6	36.0 - 46.0	%
MEAN CORPUSCULAR VOLUME (MCV) METHOD : CALCULATED PARAMETER	76.9 Low	83.0 - 101.0	fL
MEAN CORPUSCULAR HEMOGLOBIN (MCH) METHOD : CALCULATED PARAMETER	23.5 Low	27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION(MCHC) METHOD : CALCULATED PARAMETER	30.6 Low	31.5 - 34.5	g/dL
RED CELL DISTRIBUTION WIDTH (RDW) METHOD : CALCULATED PARAMETER	15.3 High	11.6 - 14.0	%
MENTZER INDEX METHOD : CALCULATED PARAMETER	15.3		
WBC DIFFERENTIAL COUNT			
NEUTROPHILS METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING	61	40.0 - 80.0	%
LYMPHOCYTES METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING	27	20.0 - 40.0	%
MONOCYTES METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING	9	2.0 - 10.0	%
EOSINOPHILS METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING	3	1 - 6	%


Dr. Akshay Dhotre
Consultant Pathologist



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



Patient Ref. No. 22000000865196

PATIENT NAME : MRS.SHALINI KUMARI

REF. DOCTOR :

CODE/NAME & ADDRESS : C000045507

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

ACCESSION NO : 0022WH003744

PATIENT ID : FH.12649694
CLIENT PATIENT ID: UID:12649694
ABHA NO :

AGE/SEX : 32 Years Female
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CORP-OPD
BILLNO-150123OPCR046177
BILLNO-150123OPCR046177

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BASOPHILS		0	0 - 2	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING				
ABSOLUTE NEUTROPHIL COUNT		3.39	2.0 - 7.0	thou/ μ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE LYMPHOCYTE COUNT		1.50	1.0 - 3.0	thou/ μ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE MONOCYTE COUNT		0.50	0.2 - 1.0	thou/ μ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE EOSINOPHIL COUNT		0.17	0.02 - 0.50	thou/ μ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE BASOPHIL COUNT		0 Low	0.02 - 0.10	thou/ μ L
METHOD : CALCULATED PARAMETER				
NEUTROPHIL LYMPHOCYTE RATIO (NLR)		2.2		
METHOD : CALCULATED				
MORPHOLOGY				
RBC		MILD HYPOCHROMASIA, MILD MICROCYTOSIS, MILD ANISOCYTOSIS		
METHOD : MICROSCOPIC EXAMINATION				
WBC		NORMAL MORPHOLOGY		
METHOD : MICROSCOPIC EXAMINATION				
PLATELETS		ADEQUATE ON SMEAR. PLATELETS SEEN ON SMEAR~1,50,000-1,60,000/microliter.		
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.
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.



Dr.Akshay Dhotre
Consultant Pathologist

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REF. DOCTOR :

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HAEMATOLOGY

ERYTHROCYTE SEDIMENTATION RATE (ESR),WHOLE BLOOD

E.S.R	18	0 - 20	mm at 1 hr
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METHOD : WESTERGRN 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 (Paraproteinemia, 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, (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. AACCC Press, 7th edition. Edited by S. Soldin; 3. The reference for the adult reference range is "Practical Haematology by Dacie and Lewis, 10th edition.



Dr. Akshay Dhotre
Consultant Pathologist

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Patient Ref. No. 2200000865196

PATIENT NAME : MRS.SHALINI KUMARI		REF. DOCTOR :	
CODE/NAME & ADDRESS : C000045507		ACCESSION NO : 0022WH003744	
FORTIS VASHI-CHC -SPLZD		AGE/SEX : 32 Years Female	
FORTIS HOSPITAL # VASHI,		DRAWN : 16/08/2023 09:56:00	
MUMBAI 440001		RECEIVED : 16/08/2023 09:56:37	
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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	0.41	0.2 - 1.0	mg/dL
METHOD : JENDRASSIK AND GROFF			
BILIRUBIN, DIRECT	0.09	0.0 - 0.2	mg/dL
METHOD : JENDRASSIK AND GROFF			
BILIRUBIN, INDIRECT	0.32	0.1 - 1.0	mg/dL
METHOD : CALCULATED PARAMETER			
TOTAL PROTEIN	7.3	6.4 - 8.2	g/dL
METHOD : BIURET			
ALBUMIN	3.7	3.4 - 5.0	g/dL
METHOD : BCP DYE BINDING			
GLOBULIN	3.6	2.0 - 4.1	g/dL
METHOD : CALCULATED PARAMETER			
ALBUMIN/GLOBULIN RATIO	1.0	1.0 - 2.1	RATIO
METHOD : CALCULATED PARAMETER			
ASPARTATE AMINOTRANSFERASE(AST/SGOT)	25	15 - 37	U/L
METHOD : UV WITH PSP			
ALANINE AMINOTRANSFERASE (ALT/SGPT)	32	< 34.0	U/L
METHOD : UV WITH PSP			
ALKALINE PHOSPHATASE	133 High	30 - 120	U/L
METHOD : PNPP-ANP			
GAMMA GLUTAMYL TRANSFERASE (GGT)	28	5 - 55	U/L
METHOD : GAMMA GLUTAMYL CARBOXY 4-NITROANTILIDE			
LACTATE DEHYDROGENASE	167	81 - 234	U/L
METHOD : LACTATE -PYRUVATE			
GLUCOSE FASTING, FLUORIDE PLASMA			
FBS (FASTING BLOOD SUGAR)	94	Normal : < 100 Pre-diabetes: 100-125 Diabetes: >=126	mg/dL
METHOD : HEXOKINASE			

GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD


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HBA1C		5.5	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) METHOD : CALCULATED PARAMETER		111.2	< 116.0	mg/dL
KIDNEY PANEL - 1				
BLOOD UREA NITROGEN (BUN), SERUM				
BLOOD UREA NITROGEN METHOD : UREASE - UV		8	6 - 20	mg/dL
CREATININE EGFR- EPI				
CREATININE METHOD : ALKALINE PICRATE KINETIC JAFFES		0.61	0.60 - 1.10	mg/dL
AGE		32		years
GLOMERULAR FILTRATION RATE (FEMALE) METHOD : CALCULATED PARAMETER		121.74	Refer Interpretation Below	mL/min/1.73m ²
BUN/CREAT RATIO				
BUN/CREAT RATIO METHOD : CALCULATED PARAMETER		13.11	5.00 - 15.00	
URIC ACID, SERUM				
URIC ACID METHOD : URICASE UV		3.8	2.6 - 6.0	mg/dL
TOTAL PROTEIN, SERUM				
TOTAL PROTEIN METHOD : BIURET		7.3	6.4 - 8.2	g/dL
ALBUMIN, SERUM				
ALBUMIN METHOD : BCP DYE BINDING		3.7	3.4 - 5.0	g/dL
GLOBULIN				
GLOBULIN METHOD : CALCULATED PARAMETER		3.6	2.0 - 4.1	g/dL



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ELECTROLYTES (NA/K/CL), SERUM

SODIUM, SERUM		141	136 - 145	mmol/L
METHOD : ISE INDIRECT				
POTASSIUM, SERUM		4.05	3.50 - 5.10	mmol/L
METHOD : ISE INDIRECT				
CHLORIDE, SERUM		106	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.

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, 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.

Dr. Akshay Dhotre
 Consultant Pathologist



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



Patient Ref. No. 2200000865196

PATIENT NAME : MRS.SHALINI KUMARI
REF. DOCTOR :
CODE/NAME & ADDRESS : C000045507
**FORTIS VASHI-CHC -SPLZD
FORTIS HOSPITAL # VASHI,
MUMBAI 440001**
ACCESSION NO : 0022WH003744
PATIENT ID : FH.12649694
CLIENT PATIENT ID: UID:12649694
ABHA NO :
AGE/SEX : 32 Years Female
DRAWN : 16/08/2023 09:56:00
RECEIVED : 16/08/2023 09:56:37
REPORTED : 16/08/2023 13:28:56
CLINICAL INFORMATION :
UID:12649694 REQNO-1560004
CORP-OPD
BILLNO-150123OPCR046177
BILLNO-150123OPCR046177

Test Report Status	Final	Results	Biological Reference Interval	Units
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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 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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Consultant Pathologist

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Patient Ref. No. 22000000865196

PATIENT NAME : MRS.SHALINI KUMARI

REF. DOCTOR :

CODE/NAME & ADDRESS : C000045507

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

ACCESSION NO : 0022WH003744

PATIENT ID : FH.12649694

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ABHA NO :

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BIOCHEMISTRY - LIPID

LIPID PROFILE, SERUM

CHOLESTEROL, TOTAL	172	< 200 Desirable 200 - 239 Borderline High ≥ 240 High	mg/dL
METHOD : ENZYMATIC/COLORIMETRIC, CHOLESTEROL OXIDASE, ESTERASE, PEROXIDASE			
TRIGLYCERIDES	64	< 150 Normal 150 - 199 Borderline High 200 - 499 High ≥ 500 Very High	mg/dL
METHOD : ENZYMATIC ASSAY			
HDL CHOLESTEROL	63 High	< 40 Low ≥ 60 High	mg/dL
METHOD : DIRECT MEASURE - PEG			
LDL CHOLESTEROL, DIRECT	101	< 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	109	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	12.8	<= 30.0	mg/dL
METHOD : CALCULATED PARAMETER			
CHOL/HDL RATIO	2.7 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			
LDL/HDL RATIO	1.6	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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Interpretation(s)


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CLINICAL PATH - URINALYSIS

URINALYSIS

PHYSICAL EXAMINATION, URINE

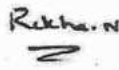
COLOR PALE YELLOW
 APPEARANCE SLIGHTLY HAZY

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, CONVERSION OF NITRATE TO NITRITE			
MICROSCOPIC EXAMINATION, URINE			
RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
PUS CELL (WBC'S)	5-7	0-5	/HPF
EPITHELIAL CELLS	2-3	0-5	/HPF
CASTS	NOT DETECTED		
CRYSTALS	NOT DETECTED		



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 Microbiologist

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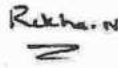
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 BILLNO-150123OPCR046177
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BACTERIA		DETECTED	NOT DETECTED	
METHOD : MICROSCOPIC EXAMINATION				
YEAST		NOT DETECTED	NOT DETECTED	
REMARKS		URINARY MICROSCOPIC EXAMINATION DONE ON URINARY CENTRIFUGED SEDIMENT		
METHOD : MICROSCOPIC EXAMINATION				
Interpretation(s)				



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SPECIALISED CHEMISTRY - HORMONE
THYROID PANEL, SERUM

T3	141.4	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 IMMUNOASSAY, COMPETITIVE PRINCIPLE			
T4	11.25	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 IMMUNOASSAY, COMPETITIVE PRINCIPLE			
TSH (ULTRASENSITIVE)	3.100	Non Pregnant Women 0.27 - 4.20 Pregnant Women 1st Trimester: 0.33 - 4.59 2nd Trimester: 0.35 - 4.10 3rd Trimester: 0.21 - 3.15	µIU/mL
METHOD : ELECTROCHEMILUMINESCENCE,SANDWICH IMMUNOASSAY			

Interpretation(s)
****End Of Report****

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

ACCESSION NO : 0022WH003805
PATIENT ID : FH.12649694
CLIENT PATIENT ID: UID:12649694
ABHA NO :
AGE/SEX : 32 Years Female
DRAWN : 16/08/2023 12:29:00
RECEIVED : 16/08/2023 12:30:26
REPORTED : 16/08/2023 14:20:43
CLINICAL INFORMATION :

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BIOCHEMISTRY
GLUCOSE, POST-PRANDIAL, PLASMA

PPBS(POST PRANDIAL BLOOD SUGAR) 99 70 - 140 mg/dL

METHOD : HEXOKINASE

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

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Patient Ref. No. 22000000865257

PATIENT NAME : MRS.SHALINI KUMARI		REF. DOCTOR :	
CODE/NAME & ADDRESS : C000045507		ACCESSION NO : 0022WH003870	
FORTIS VASHI-CHC -SPLZD		AGE/SEX : 32 Years Female	
FORTIS HOSPITAL # VASHI,		DRAWN : 16/08/2023 14:50:00	
MUMBAI 440001		RECEIVED : 16/08/2023 14:54:49	
		REPORTED : 16/08/2023 20:06:55	
		PATIENT ID : FH.12649694	
		CLIENT PATIENT ID: UID:12649694	
		ABHA NO :	

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Test Report Status **Final**

Units

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 FEW POLYMORPHS.

NEGATIVE FOR INTRAEPITHELIAL LESION OR MALIGNANCY

INTERPRETATION / RESULT

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Patient Ref. No. 2200000865322

12649694
32 Years

MRS SHALINI KUMARI
Female

ML

Rate 87 . Sinus rhythm.....normal P axis, V-rate 50- 99

PR 125
QRSD 87
QT 337
QTc 406

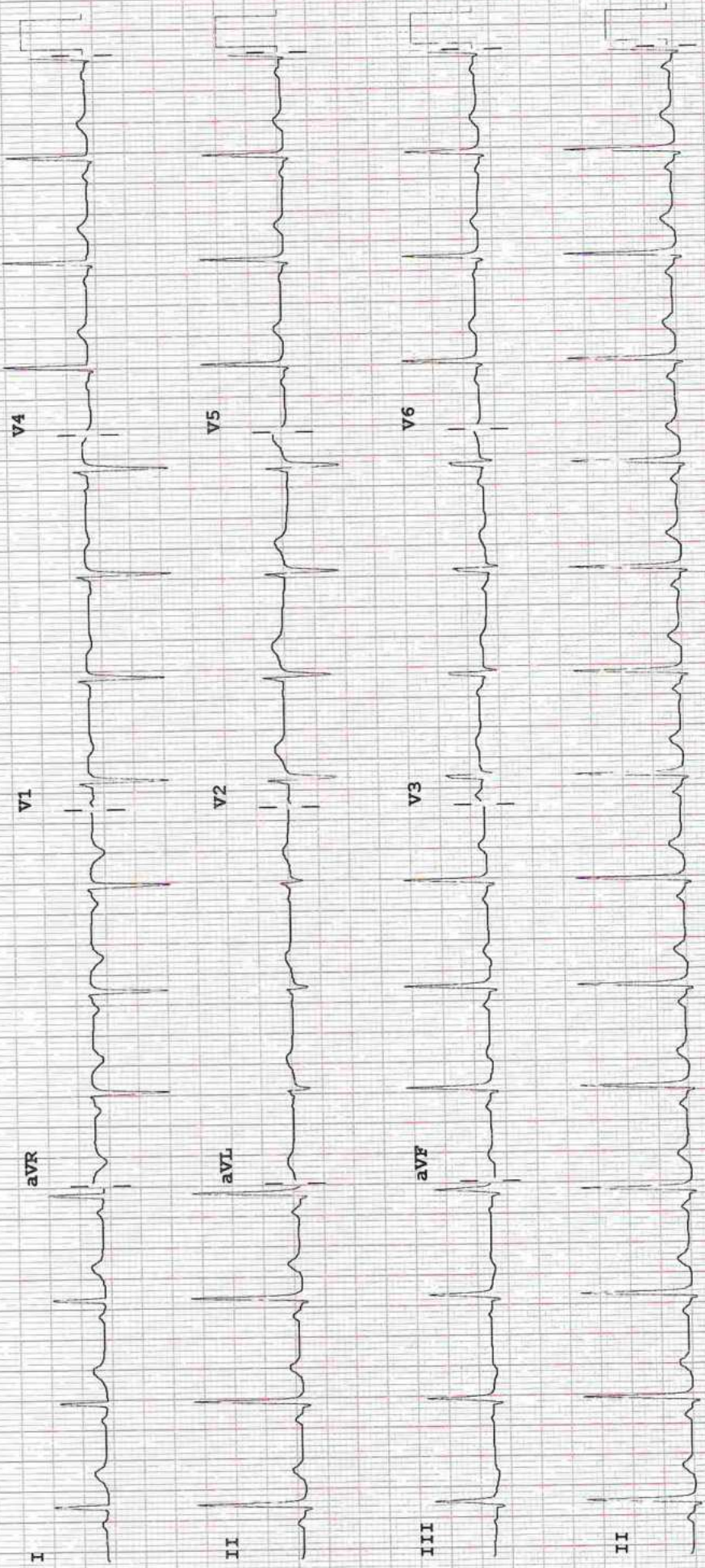
--AXIS--

P 51
QRS 62
T 17

12 Lead; Standard Placement

- NORMAL ECG -

Unconfirmed Diagnosis



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

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



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CIN: U85100MH2005PTC 154823
GST IN : 27AABCH5894D1ZG
PAN NO : AABCH5894D

(For Billing/Reports & Discharge Summary only)

Date: 17/Aug/2023

DEPARTMENT OF NIC

Name: Mrs. Shalini Kumari
Age | Sex: 32 YEAR(S) | Female
Order Station : FO-OPD
Bed Name :

UHID | Episode No : 12649694 | 46880/23/1501
Order No | Order Date: 1501/PN/OP/2308/97587 | 16-Aug-2023
Admitted On | Reporting Date : 17-Aug-2023 09:24:56
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	35	mm
AO Root	29	mm
AO CUSP SEP	18	mm
LVID (s)	31	mm
LVID (d)	43	mm
IVS (d)	09	mm
LVPW (d)	10	mm
RVID (d)	29	mm
RA	28	mm
LVEF	60	%

DOPPLER STUDY:



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DEPARTMENT OF NIC

Date: 17/Aug/2023

Name: Mrs. Shalini Kumari

UHID | Episode No : 12649694 | 46880/23/1501

Age | Sex: 32 YEAR(S) | Female

Order No | Order Date: 1501/PN/OP/2308/97587 | 16-Aug-2023

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Bed Name :

Order Doctor Name : Dr.SELF .

E WAVE VELOCITY: 0.9 m/sec.

A WAVE VELOCITY:0.5 m/sec

E/A RATIO:1.4

	PEAK (mmHg)	MEAN (mmHg)	V max (m/sec)	GRADE OF REGURGITATION
MITRAL VALVE	N			Nil
AORTIC VALVE	05			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)

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CIN: U85100MH2005PTC 154823

GST IN : 27AABCH5894D1ZG

PAN NO : AABCH5894D



DEPARTMENT OF RADIOLOGY

Date: 16/Aug/2023

Name: Mrs. Shalini Kumari

UHID | Episode No : 12649694 | 46880/23/1501

Age | Sex: 32 YEAR(S) | Female

Order No | Order Date: 1501/PN/OP/2308/97587 | 16-Aug-2023

Order Station : FO-OPD

Admitted On | Reporting Date : 16-Aug-2023 15:41:52

Bed Name :

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 are unremarkable.

DR. CHETAN KHADKE
M.D. (Radiologist)



(For Billing/Reports & Discharge Summary only)

DEPARTMENT OF RADIOLOGY

Date: 16/Aug/2023

Name: Mrs. Shalini Kumari

UHID | Episode No : 12649694 | 46880/23/1501

Age | Sex: 32 YEAR(S) | Female

Order No | Order Date: 1501/PN/OP/2308/97587 | 16-Aug-2023

Order Station : FO-OPD

Admitted On | Reporting Date : 16-Aug-2023 14:06:53

Bed Name :

Order Doctor Name : Dr.SELF .

US-WHOLE ABDOMEN

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

GALL BLADDER is physiologically distended and shows a calculus of size 17 mm in neck region. Gall bladder reveals normal wall thickness. 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 10.7 x 4.5 cm. Left kidney measures 10.1 x 5.3 cm.

PANCREAS: Head and body of pancreas is visualised and appears normal. Rest of the pancreas is obscured.

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

UTERUS is normal in size & anteverted, measuring 7.7 x 4.9 x 3.0 cm. Endometrium measures 7.5 mm in thickness.

Both ovaries are normal.

Right ovary measures 3.5 x 1.8 cm. Left ovary measures 4.7 x 2.1 cm.

No evidence of ascites.

Impression:

- Cholelithiasis without changes of cholecystitis.

DR. CHETAN KHADKE

M.D. (Radiologist)