

BMI CHART

Date: 09/12/22

Name: Chanchads Singh Age: 34 yrs Sex: M/F

BP: 180/100 mmHg Height (cms): 153cm Weight(kgs): 71.5kg BMI: _____

SpO2 - 100%
Pulse 94b/m

Repeat BP.

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	
kg	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	40		
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		
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		
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		
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		
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		
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		
5'9" - 175.2	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36		
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		
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		
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		
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		
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		
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		
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		

Doctors Notes:

Signature



UHID	12381878	Date	09/12/2023		
Name	Mrs.Chanchala Sinha	Sex	Female	Age	54
OPD	Dental 12	Health Check-up			

O/E stains +
- calculus +

Drug allergy:
Sys illness:

Treatment

- Ald - ① Scaling Grade I (Cleaning)
② OPG (Xray)

To pay -

⇒ OPG = Rs 1200

Dr. Tupti

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
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 GST IN: 27AABCH5894D1ZG | PAN NO: AABCH5894D



Hiranandani
HOSPITAL
 (A Fortis Network Hospital)

UHID	12381878	Date	09/12/2023		
Name	Mrs.Chanchala Sinha	Sex	Female	Age	54
OPD	Ophthal 14	Health Check-up			

Drug allergy:
 Sys illness:

O/C → Nil, sometimes itching

O/H → BP ∴ 26 times

(under medication)

OO → OD → -0.50X90
 OO → OS → -0.50X120
 Add! +2.25 DSph

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UHID	12381878	Date	09/12/2023		
Name	Mrs.Chanchala Sinha	Sex	Female	Age	54
OPD	Pap Smear	Health Check-up			

Drug allergy:
Sys illness:

PATIENT NAME : MRS.CHANCHALA SINHA

REF. DOCTOR :

CODE/NAME & ADDRESS : C000045507

ACCESSION NO : 0022WL001453

AGE/SEX : 54 Years Female

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

PATIENT ID : FH.12381878

DRAWN : 09/12/2023 10:37:00

CLIENT PATIENT ID: UID:12381878

RECEIVED : 09/12/2023 10:37:39

ABHA NO :

REPORTED : 09/12/2023 19:58:27

CLINICAL INFORMATION :

UID:12381878 REQNO-1635545
CORP-OPD
BILLNO-150123OPCR069341
BILLNO-150123OPCR069341

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HAEMATOLOGY - CBC

CBC-5, EDTA WHOLE BLOOD

BLOOD COUNTS, EDTA WHOLE BLOOD

HEMOGLOBIN (HB)	13.2	12.0 - 15.0	g/dL
METHOD : SLS METHOD			
RED BLOOD CELL (RBC) COUNT	4.68	3.8 - 4.8	mil/ μ L
METHOD : HYDRODYNAMIC FOCUSING			
WHITE BLOOD CELL (WBC) COUNT	5.71	4.0 - 10.0	thou/ μ L
METHOD : FLUORESCENCE FLOW CYTOMETRY			
PLATELET COUNT	239	150 - 410	thou/ μ L
METHOD : HYDRODYNAMIC FOCUSING BY DC DETECTION			

RBC AND PLATELET INDICES

HEMATOCRIT (PCV)	40.3	36.0 - 46.0	%
METHOD : CUMULATIVE PULSE HEIGHT DETECTION METHOD			
MEAN CORPUSCULAR VOLUME (MCV)	86.1	83.0 - 101.0	fL
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	28.2	27.0 - 32.0	pg
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION(MCHC)	32.8	31.5 - 34.5	g/dL
METHOD : CALCULATED PARAMETER			
RED CELL DISTRIBUTION WIDTH (RDW)	13.1	11.6 - 14.0	%
METHOD : CALCULATED PARAMETER			
MENTZER INDEX	18.4		
METHOD : CALCULATED PARAMETER			
MEAN PLATELET VOLUME (MPV)	9.3	6.8 - 10.9	fL
METHOD : CALCULATED PARAMETER			

WBC DIFFERENTIAL COUNT


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NEUTROPHILS		49	40.0 - 80.0	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING				
LYMPHOCYTES		41 High	20.0 - 40.0	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING				
MONOCYTES		5	2.0 - 10.0	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING				
EOSINOPHILS		5	1 - 6	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING				
BASOPHILS		0	0 - 2	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING				
ABSOLUTE NEUTROPHIL COUNT		2.80	2.0 - 7.0	thou/ μ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE LYMPHOCYTE COUNT		2.34	1.0 - 3.0	thou/ μ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE MONOCYTE COUNT		0.29	0.2 - 1.0	thou/ μ L
METHOD : CALCULATED PARAMETER				
ABSOLUTE EOSINOPHIL COUNT		0.29	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)		1.2		
METHOD : CALCULATED				

MORPHOLOGY

RBC

PREDOMINANTLY NORMOCYTIC NORMOCHROMIC

METHOD : MICROSCOPIC EXAMINATION

WBC

NORMAL MORPHOLOGY

METHOD : MICROSCOPIC EXAMINATION

PLATELETS

ADEQUATE

METHOD : MICROSCOPIC EXAMINATION



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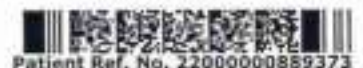
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CORP-OPD

BILLNO-150123OPCR069341

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Interpretation(s)

RBC AND PLATELET INDICES-Mentzer Index (MCV/PLC) is an automated cell-counter based calculated screen tool to differentiate cases of Iron deficiency anemia(>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, 45.1% 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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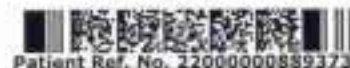
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HAEMATOLOGY

ERYTHROCYTE SEDIMENTATION RATE (ESR), EDTA BLOOD

E.S.R	16	0 - 20	mm at 1 hr
-------	----	--------	------------

METHOD : WESTERGRÉN 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


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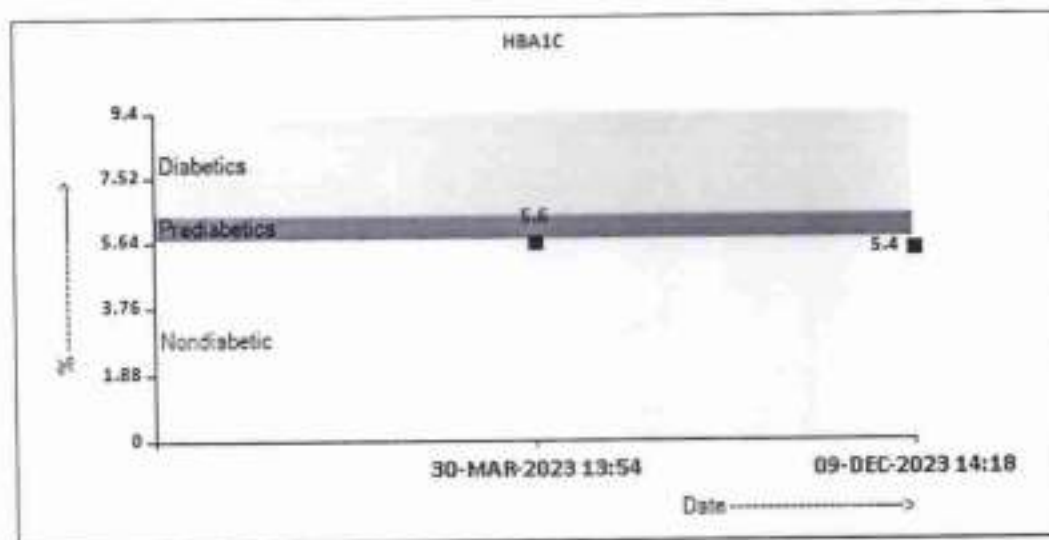
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Interpretation(s)

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 millimeters 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

Increased in: Infections, Vasculitis, Inflammatory arthritis, Renal disease, Anemia, Malignancies and plasma cell dyscrasia, Acute allergy Tissue injury, Pregnancy, Estrogen medication, Aging.

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

In pregnancy BIR in first trimester is 0-10 mm/hr (52 if anemic) and in second trimester (0-70 mm/hr) (55 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 : Folic acid, Sickle Cells, spherocytes, Microcytosis, Low fibrinogen, Very high WBC counts, Drugs (Quinine, salicylates)

REFERENCE :

1. Nathan and Oski's Hematology of Infancy and Childhood, 5th edition; 2. Paediatric reference intervals, AACCPress, 7th edition, Edited by S. Soldin; 3. The reference for

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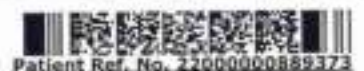
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The adult reference range is "Practical Haematology by Dacie and Lewis, 10th edition, GLYCOSYLATED HEMOGLOBIN(HbA1c), EDTA WHOLE BLOOD-Used Ferr

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

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.

1. eAG (Estimated average glucose) converts percentage HbA1c to mg/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, addition are reported to interfere with some assay methods, falsely increasing results.
4. Interference of hemoglobinopathies in HbA1c estimation is seen in

- a) Heterozygous 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



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IMMUNOHAEMATOLOGY

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP

TYPE AB

METHOD : TUBE AGGLUTINATION

RH TYPE

NEGATIVE

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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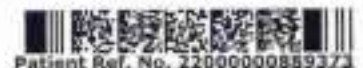
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BIOCHEMISTRY

LIVER FUNCTION PROFILE, SERUM

BILIRUBIN, TOTAL METHOD : JENDRASSIK AND GROFF	0.38	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	0.28	0.1 - 1.0	mg/dL
TOTAL PROTEIN METHOD : BIURET	7.4	6.4 - 8.2	g/dL
ALBUMIN METHOD : BCF DYE BINDING	4.1	3.4 - 5.0	g/dL
GLOBULIN METHOD : CALCULATED PARAMETER	3.3	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO METHOD : CALCULATED PARAMETER	1.2	1.0 - 2.1	RATIO
ASPARTATE AMINOTRANSFERASE(AST/SGOT) METHOD : UV WITH PSP	18	15 - 37	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT) METHOD : UV WITH PSP	32	< 34.0	U/L
ALKALINE PHOSPHATASE METHOD : PNPP-AMP	62	30 - 120	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT) METHOD : GAMMA GLUTAMYL CARBOXY 4-NITROANILIDE	22	5 - 55	U/L
LACTATE DEHYDROGENASE METHOD : LACTATE-P-HYDROLYTIC	141	81 - 234	U/L

GLUCOSE FASTING, FLUORIDE PLASMA

FBS (FASTING BLOOD SUGAR) METHOD : HEXOKINASE	95	Normal : < 100 Pre-diabetes: 100-125 Diabetes: >=126	mg/dL
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 CIN - U74899PB1995PLC045956
 Email : -


Patient Ref. No. 22000000889373

PATIENT NAME : MRS.CHANCHALA SINHA

REF. DOCTOR :

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

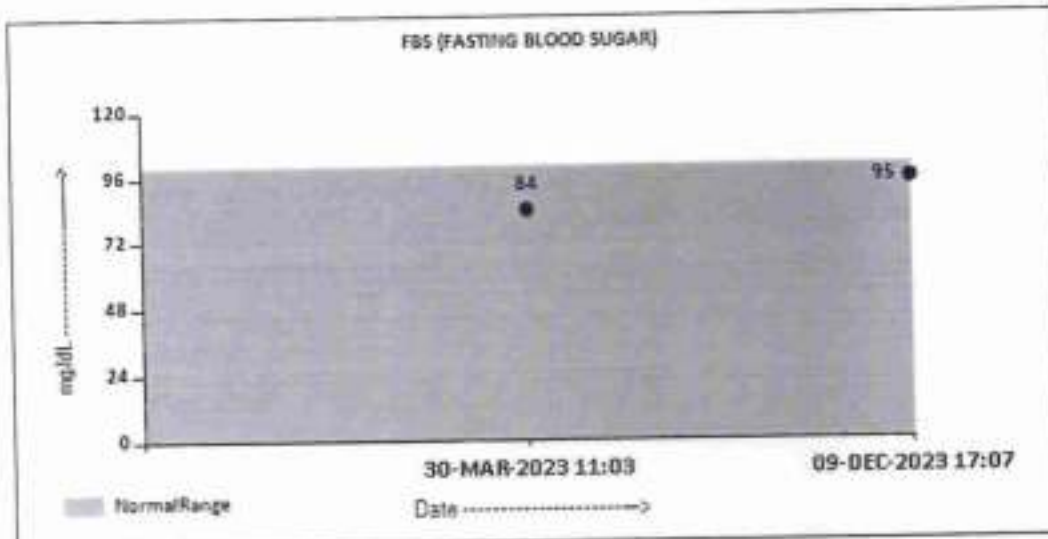
ACCESSION NO : 0022WL001453
PATIENT ID : FH.12381878
CLIENT PATIENT ID: UID:12381878
ABHA NO :

AGE/SEX : 54 Years Female
DRAWN : 09/12/2023 10:37:00
RECEIVED : 09/12/2023 10:37:39
REPORTED : 09/12/2023 19:58:27

CLINICAL INFORMATION :

UTD:12381878 REQNO-1635545
 CORP-OPD
 BILLNO-1501230PCR069341
 BILLNO-1501230PCR069341

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

BLOOD UREA NITROGEN (BUN), SERUM

BLOOD UREA NITROGEN 10 6 - 20 mg/dL
 METHOD : UREASE - UV

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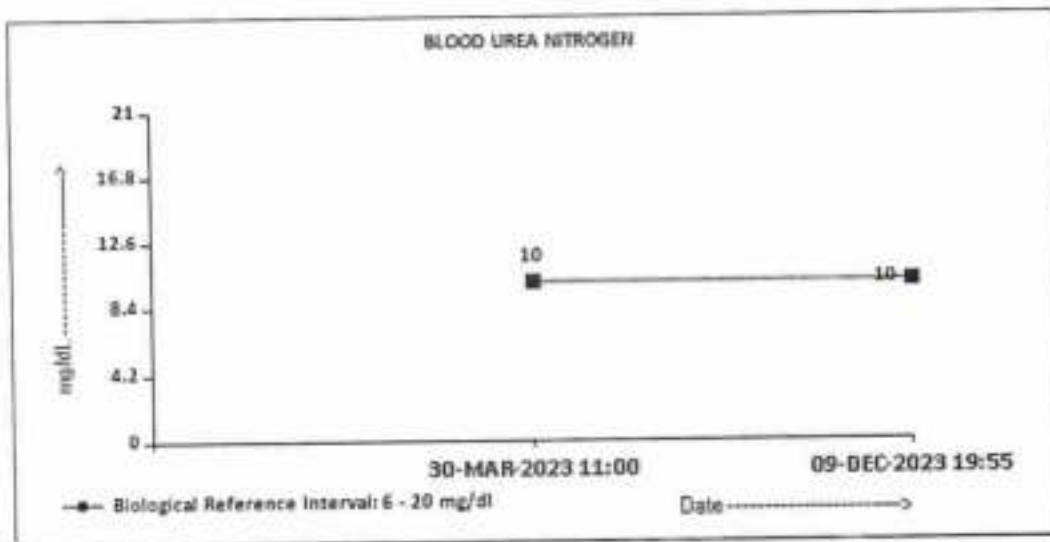


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CREATININE EGFR- EPI			
CREATININE	0.77	0.60 - 1.10	mg/dL
METHOD : ALKALINE PHOSPHATE KINETIC JAFFES			
AGE	54		years
GLOMERULAR FILTRATION RATE (FEMALE)	91.61	Refer Interpretation Below	mL/min/1.73m2
METHOD : CALCULATED PARAMETER			

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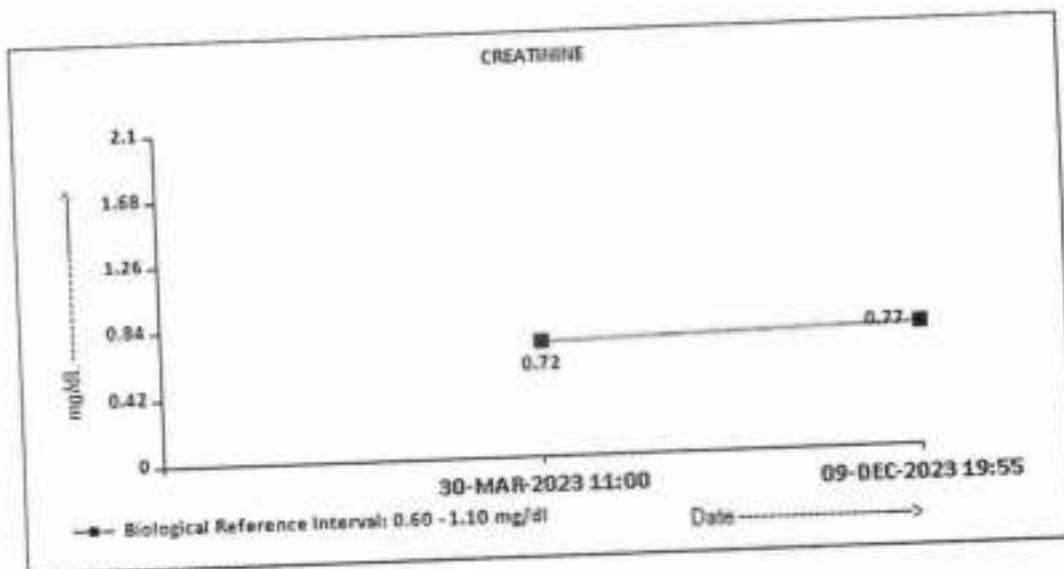


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BUN/CREAT RATIO	12.99	5.00 - 15.00	
BUN/CREAT RATIO			
METHOD : CALCULATED PARAMETER			
URIC ACID, SERUM	6.9 High	2.6 - 6.0	mg/dL
URIC ACID			
METHOD : URICASE UV			
TOTAL PROTEIN, SERUM	7.4	6.4 - 8.2	g/dL
TOTAL PROTEIN			
METHOD : BIURET			
ALBUMIN, SERUM			

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ALBUMIN		4.1	3.4 - 5.0	g/dL
METHOD : BCF DYE BINDING				
GLOBULIN		3.3	2.0 - 4.1	g/dL
METHOD : CALCULATED PARAMETER				
ELECTROLYTES (NA/K/CL), SERUM				
SODIUM, SERUM		141	136 - 145	mmol/L
METHOD : ISE INDIRECT				
POTASSIUM, SERUM		4.88	3.50 - 5.10	mmol/L
METHOD : ISE INDIRECT				
CHLORIDE, SERUM		105	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 haem 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 Hypoparathyroidism, 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


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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, Waldenström 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, hemodialysis, 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 Hypoglycemics & Insulin treatment, Renal Glycosuria, Glycemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc.

BLOOD UREA NITROGEN (BUN), SERUM-Causes of Increased levels include the renal (High protein diet, Increased protein catabolism, GI haemorrhage, Celiac), Dehydration, CHF Reveal), 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:

References:

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

Estimated GFR Calculated Using the CKD-EPI equation-<https://testguide.kidney.org/goldfile/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 Principles 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, Waldenström 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, hemodialysis, increased vascular permeability or decreased lymphatic clearance, malnutrition and wasting etc.

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Patient Ref. No. 22000000889373

PATIENT NAME : MRS.CHANCHALA SINHA

REF. DOCTOR :

CODE/NAME & ADDRESS : C000045507

ACCESSION NO : 0022WL001453

AGE/SEX : 54 Years Female

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FORTIS HOSPITAL # VASHI,
MUMBAI 440001

PATIENT ID : FH,12381878

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ABHA NO :

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CLINICAL INFORMATION :

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CORP-OPD
BILLNO-150123OPCR069341
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BIOCHEMISTRY - LIPID

LIPID PROFILE, SERUM

CHOLESTEROL, TOTAL	229 High	< 200 Desirable 200 - 239 Borderline High >= 240 High	mg/dL
METHOD : ENZYMATIC/COLORIMETRIC, CHOLESTEROL OXIDASE, ESTERASE, PEROXIDASE			
TRIGLYCERIDES	162 High	< 150 Normal 150 - 199 Borderline High 200 - 499 High >= 500 Very High	mg/dL
METHOD : ENZYMATIC ASSAY			
HDL CHOLESTEROL	48	< 40 Low >= 60 High	mg/dL
METHOD : DIRECT MEASURE - PEG			
LDL CHOLESTEROL, DIRECT	141 High	< 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	181 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			
VERY LOW DENSITY LIPOPROTEIN	32.4 High	<= 30.0	mg/dL
METHOD : CALCULATED PARAMETER			
CHOL/HDL RATIO	4.8 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			



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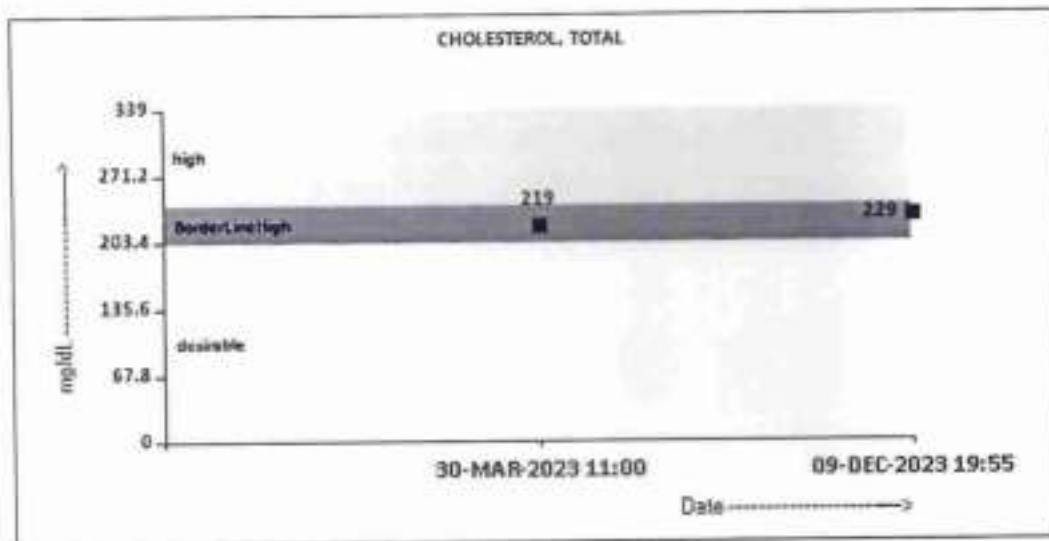
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LDL/HDL RATIO		2.9	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	

METHOD : CALCULATED PARAMETER.



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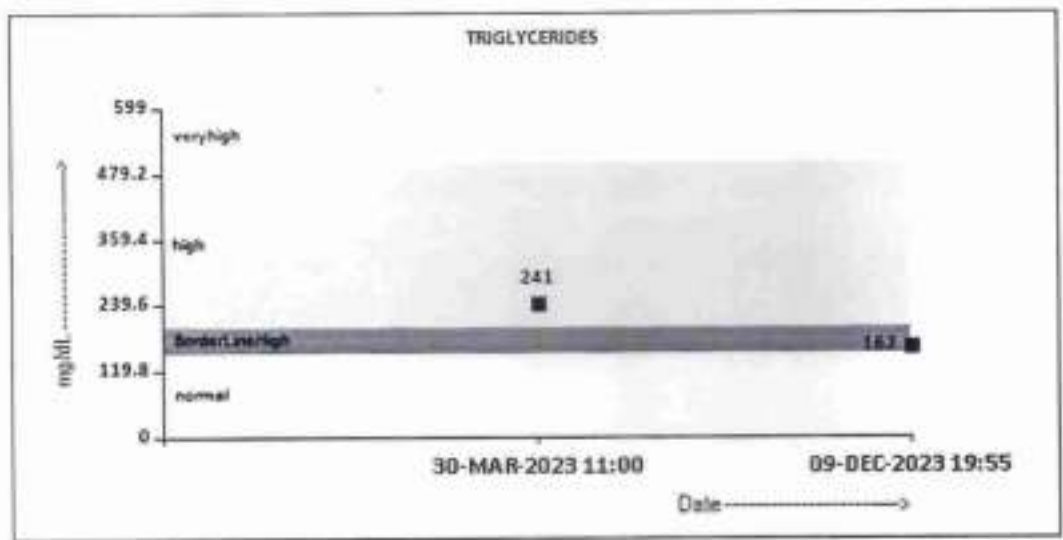


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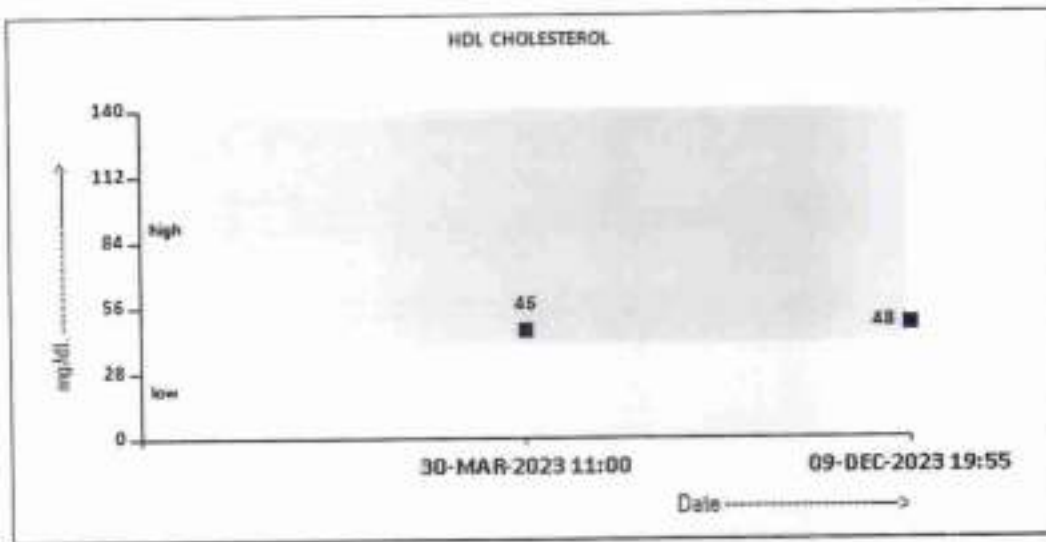


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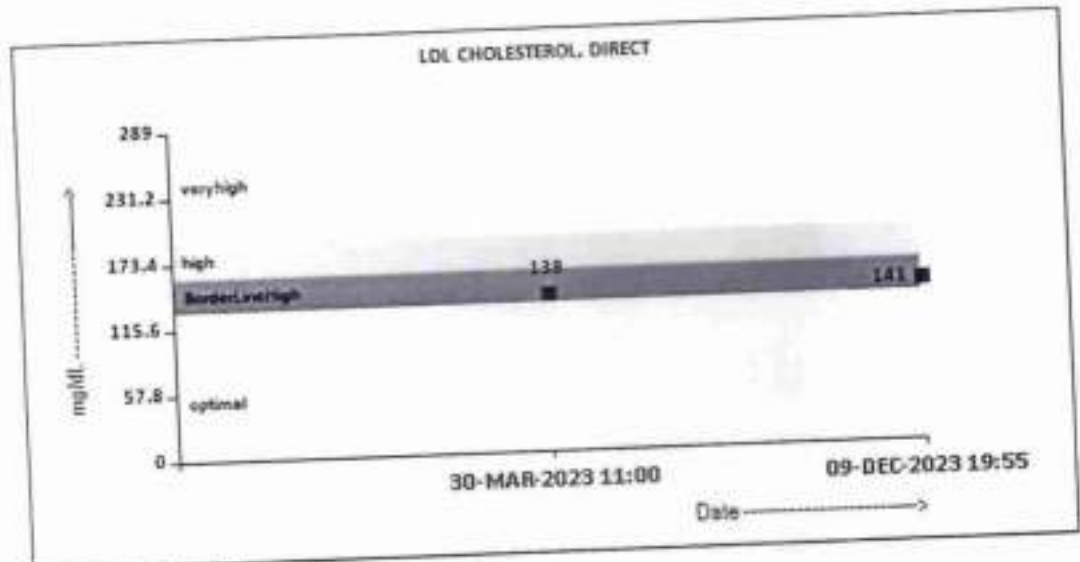


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 Consultant Pathologist



PERFORMED AT :
 Agilus Diagnostics Ltd.
 Hiranandani Hospital-Vashi, Mini Seashore Road, Sector 10,
 Navi Mumbai, 400703
 Maharashtra, India
 Tel : 022-39199222, 022-49723322,
 CIN - U74899MH1995PLC045956
 Email : -



PATIENT NAME : MRS.CHANCHALA SINHA		REF. DOCTOR :	
CODE/NAME & ADDRESS : C000045507	ACCESSION NO : 0022WL001453	AGE/SEX : 54 Years	Female
FORTIS VASHI-CHC -SPLZD	PATIENT ID : FH.12381878	DRAWN : 09/12/2023 10:37:00	
FORTIS HOSPITAL # VASHI,	CLIENT PATIENT ID: UID:12381878	RECEIVED : 09/12/2023 10:37:39	
MUMBAI 440001	ABHA NO :	REPORTED : 09/12/2023 19:58:27	

CLINICAL INFORMATION :

UID:12381878 REQNO-1635545
CORP-OPD
BILLNO-1501230PCRD69341
BILLNO-1501230PCRD69341

Test Report Status	Results	Biological Reference Interval	Units
Final			

CLINICAL PATH - URINALYSIS

KIDNEY PANEL - 1

PHYSICAL EXAMINATION, URINE

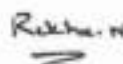
COLOR	PALE YELLOW
METHOD : PHYSICAL	
APPEARANCE	SLIGHTLY HAZY
METHOD : VISUAL	

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 PFA 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-GOO/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	DETECTED (+)	NOT DETECTED
METHOD : REFLECTANCE SPECTROPHOTOMETRY, ESTERASE HYDROLYSIS ACTIVITY		



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



Dr. Rekha Nair, MD
(Reg No. MMC 2001/06/2354)
Microbiologist



View Details



View Report

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Email : -



Patient Ref. No. 2200000889373

PATIENT NAME : MRS.CHANCHALA SINHA		REF. DOCTOR :
CODE/NAME & ADDRESS : C000045507 FORTIS VASHI-CHC -SPLZD FORTIS HOSPITAL # VASHI, MUMBAI 440001	ACCESSION NO : 0022WL001453 PATIENT ID : PH.12381878 CLIENT PATIENT ID: UID:12381878 ABHA NO :	AGE/SEX : 54 Years Female DRAWN : 09/12/2023 10:37:00 RECEIVED : 09/12/2023 10:37:39 REPORTED : 09/12/2023 19:58:27

CLINICAL INFORMATION :

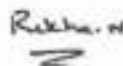
UID:12381878 REQNO-1635545
CORP-OPD
BILLNO-1501230PCRD69341
BILLNO-1501230PCRD69341

Test Report Status	Results	Biological Reference Interval	Units
Final			
MICROSCOPIC EXAMINATION, URINE			
RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
METHOD : MICROSCOPIC EXAMINATION			
PUS CELL (WBC'S)	15-20	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)



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



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Microbiologist



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Patient Ref. No. 2200000889373

PATIENT NAME : MRS.CHANCHALA SINHA

REF. DOCTOR :

CODE/NAME & ADDRESS : C000045507

ACCESSION NO : 0022WL001453

AGE/SEX : 54 Years Female

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

PATIENT ID : FH.12381878

DRAWN : 09/12/2023 10:37:00

CLIENT PATIENT ID: UID:12381878

RECEIVED : 09/12/2023 10:37:39

ABHA NO :

REPORTED : 09/12/2023 19:58:27

CLINICAL INFORMATION :

UID:12381878 REQNO-1635545
CORP-OPD
BILLNO-150123OPCR069341
BILLNO-150123OPCR069341

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

THYROID PANEL, SERUM

T3	149.2	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	8.24	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 (ULTRA SENSITIVE)	1.820	Non Pregnant Women 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	µIU/mL
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METHOD : ELECTROCHEMILUMINESCENCE, SANDWICH IMMUNOASSAY

Interpretation(s)

End Of Report

Please visit 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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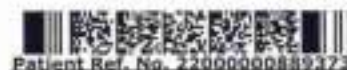


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Maharashtra, India
Tel : 022-39199222, 022-49723322,
CIN - U74899PB1995PLC045956
Email : -

Patient Ref. No. 22000000889173

PATIENT NAME : MRS.CHANCHALA SINHA

REF. DOCTOR :

CODE/NAME & ADDRESS : C000045507

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

ACCESSION NO : 0022WL001532

PATIENT ID : FH.12381878

CLIENT PATIENT ID: UID:12381878

ABHA NO :

AGE/SEX : 54 Years Female

DRAWN : 09/12/2023 13:53:00

RECEIVED : 09/12/2023 13:53:54

REPORTED : 09/12/2023 17:59:46

CLINICAL INFORMATION :

UID:12381878 REQNO-1635545

CORP-OPD

BILLNO-1501230PCR069341

BILLNO-1501230PCR069341

Test Report Status	Final	Results	Biological Reference Interval	Units
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BIOCHEMISTRY

GLUCOSE, POST-PRANDIAL, PLASMA

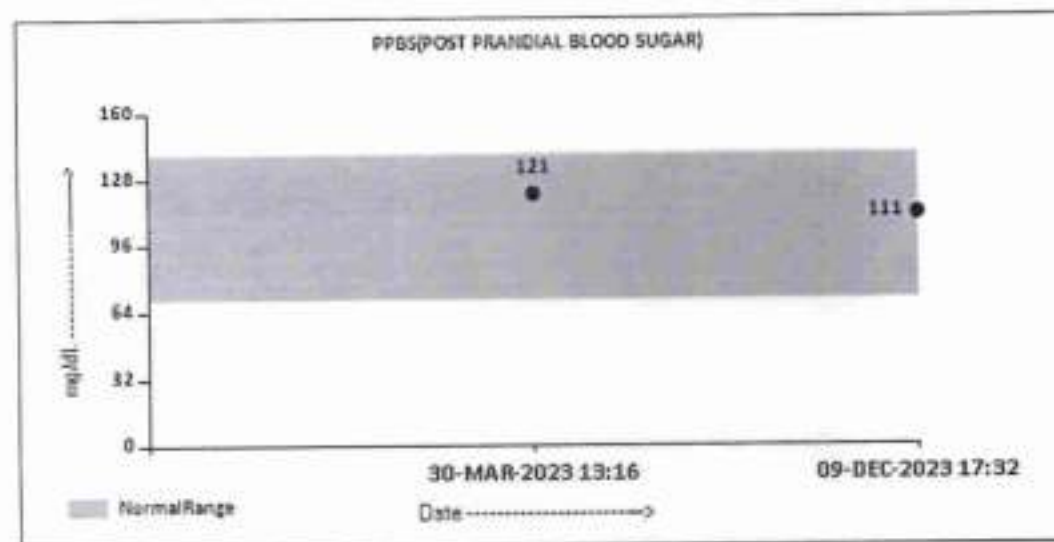
PPBS(POST PRANDIAL BLOOD SUGAR)

111

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, Glycemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc. Additional test HbA1c

End Of Report

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Dr. Akshay Dhotre, MD
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Email : -



Patient Ref. No. 22000000889452

12381878
54 Years

MRS. CHANCHALA SINHA
Female

12/9/2023 2:10:08 PM

Rate 95 . Sinus rhythm.normal P axis, V-rate 50- 99

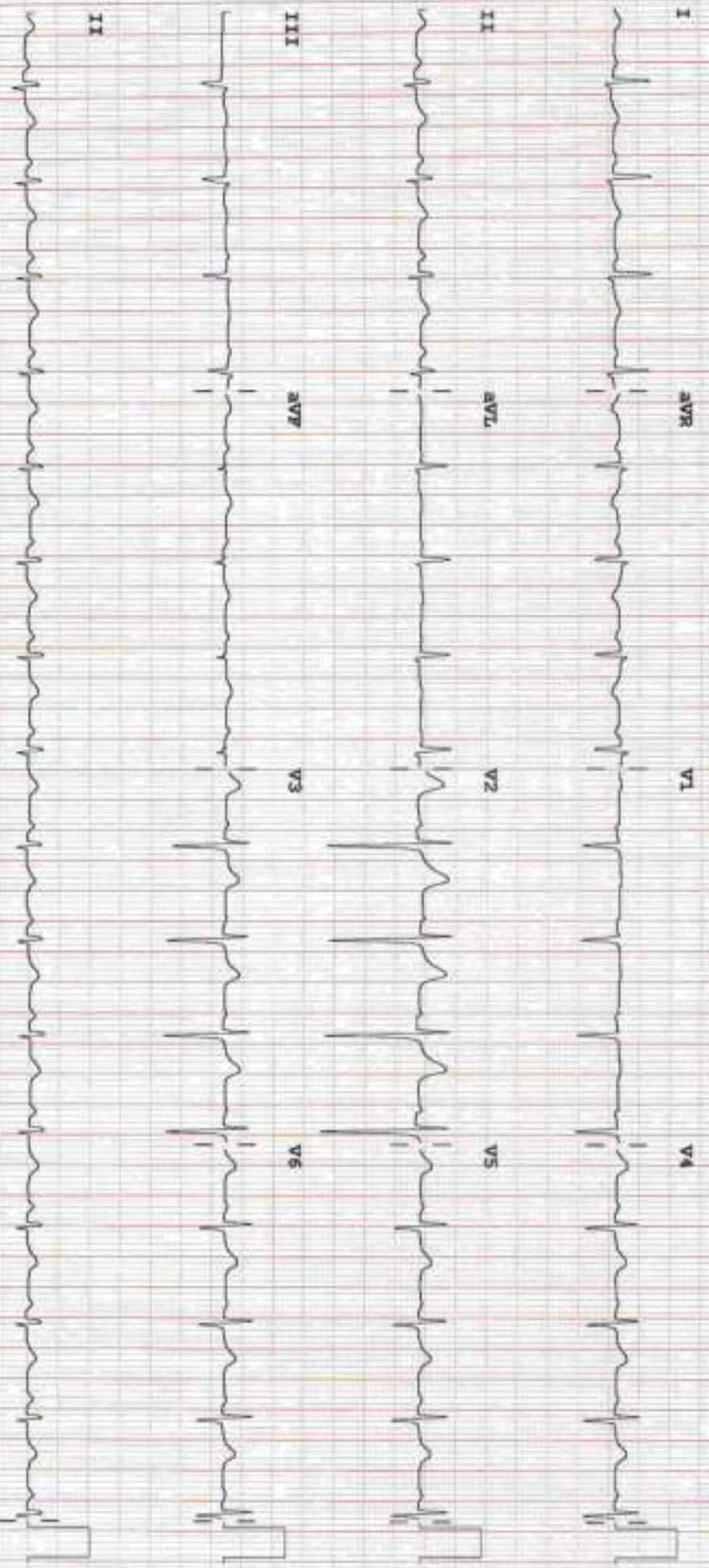
PR 137
QRSD 81
QT 361
QTc 454

--AXIS--
P 31
QRS -7
T 39

- NORMAL ECG -

12 lead; Standard Placement

Unconfirmed Diagnosis



*H/C
1st dx during
rel + Normal
P*

Device: Speed: 25 mm/sec Imp: 10 mm/mV Chest: 10.0 mm/mV F 50-0.50-100 Hz W 1000B CL P2



DEPARTMENT OF NIC

Date: 11/Dec/2023

Name: Mrs. Chaanchala Sinha
Age | Sex: 54 YEAR(S) | Female
Order Station : FO-OPD
Bed Name :

UHID | Episode No : 12381878 | 70566/23/1501
Order No | Order Date: 1501/PN/OP/2312/146480 | 09-Dec-2023
Admitted On | Reporting Date : 11-Dec-2023 12:11:36
Order Doctor Name : Dr.SELF.

ECHOCARDIOGRAPHY TRANSTHORACIC

FINDINGS:

- No left ventricle regional wall motion abnormality at rest.
- Normal left ventricle systolic function. LVEF = 60%.
- Grade I left ventricle diastolic dysfunction. No e/o raised LVEDP.
- No mitral regurgitation.
- No aortic regurgitation. No aortic stenosis.
- No tricuspid regurgitation. No pulmonary hypertension.
- Intact IVS and IAS.
- No left ventricle clot/vegetation/pericardial effusion.
- Normal right atrium and right ventricle dimension and function.
- Normal left atrium and left ventricle dimension.
- IVC measures 10 mm with normal inspiratory collapse.

M-MODE MEASUREMENTS:

LA	31	mm
AO Root	21	mm
AO CUSP SEP	18	mm
LVID (s)	22	mm
LVID (d)	44	mm
IVS (d)	10	mm
LVPW (d)	11	mm
RVID (d)	21	mm
RA	29	mm
LVEF	60	%



DEPARTMENT OF NIC

Date: 11/Dec/2023

Name: Mrs. Chanchala Sinha
Age | Sex: 54 YEAR(S) | Female
Order Station : FO-OPD
Bed Name :

UHID | Episode No : 12381878 | 70566/23/1501
Order No | Order Date: 1501/PN/OP/2312/146480 | 09-Dec-2023
Admitted On | Reporting Date : 11-Dec-2023 12:11:36
Order Doctor Name : Dr.SELF .

DOPPLER STUDY:

E WAVE VELOCITY: 0.9 m/sec.

A WAVE VELOCITY: 1.1 m/sec


E/A RATIO: 0.8

	PEAK (mmHg)	MEAN (mmHg)	V max (m/sec)	GRADE OF REGURGITATION
MITRAL VALVE	N			Nil
AORTIC VALVE	08			Nil
TRICUSPID VALVE	N			Nil
PULMONARY VALVE	04			Nil

Final Impression :

- No RWMA.
- Grade I LV diastolic dysfunction.
- Normal LV and RV systolic function.

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


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

Hiranandani Healthcare Pvt. Ltd.

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

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

Emergency: 022 - 39199100 | Ambulance: 1255

For Appointment: 022 - 39199200 | Health Checkup: 022 - 39199300

www.fortishealthcare.com | vashi@fortishealthcare.com

CIN: U85100MH2005PTC 154823

GST IN : 27AABCH5894D12G

PAN NO : AABCH5894D



DEPARTMENT OF RADIOLOGY

Date: 09/Dec/2023

Name: Mrs. Chanchala Sinha

Age | Sex: 54 YEAR(S) | Female

Order Station : FO-OPD

Bed Name :

UHID | Episode No : 12381878 | 70566/23/1501

Order No | Order Date: 1501/PN/OP/2312/146480 | 09-Dec-2023

Admitted On | Reporting Date : 09-Dec-2023 15:39:52

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	: Chanchala Sinha	Patient ID	: 12381878
Sex / Age	: F / 54Y 1M 28D	Accession No.	: PHC.7079793
Modality	: US	Scan DateTime	: 09-12-2023 12:16:36
IPID No	: 70566/23/1501	ReportDatetime	: 09-12-2023 16:25:51

USG – WHOLE ABDOMEN (TAS)

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 shows mildly raised cortical echogenicity with maintained corticomedullary differentiation. No evidence of calculi/hydronephrosis.

Right kidney measures 10.2 x 4.4 cm.

Left kidney measures 10.9 x 4.5 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 – post menopausal status.

Endometrium measures 3 mm in thickness.

Both ovaries are not visualised, however adnexae are clear.

No evidence of ascites.

Impression:

- Grade I fatty infiltration of liver.
- Mildly raised cortical echogenicity of both kidneys. *Recommended RFT correlation.*

DR. YOGESH PATHADE
M.D. (Radiologist)