

PATIENT NAME : MRS. MRS.UTPALPARNA BHATTACHARJEEPATIENT ID : **FH.12143432**

CLIENT PATIENT ID : UID:12143432

ACCESSION NO : **0022VK005774**

AGE : 29 Years

SEX : Female

ABHA NO :

DRAWN : 26/11/2022 09:00:00

RECEIVED : 26/11/2022 09:00:21

REPORTED : 26/11/2022 12:46:21

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR : SELF

CLINICAL INFORMATION :

UID:12143432 REQNO-1325842

CORP-OPD

BILLNO-150122OPCR059788

BILLNO-150122OPCR059788

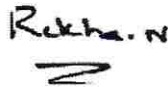
Test Report Status	Final	Results	Biological Reference Interval
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HbA1c Estimation can get affected due to :

- I.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.
- II.Vitamin C & E are reported to falsely lower test results.(possibly by inhibiting glycation of hemoglobin.
- III.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.
- IV.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

****End Of Report******Please visit www.srlworld.com for related Test Information for this accession**


Dr.Akta Dubey
Counsultant Pathologist



Dr. Rekha Nair, MD
Microbiologist



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CLIENT PATIENT ID : UID:12143432

ACCESSION NO : **0022VK005774**

AGE : 29 Years

SEX : Female

ABHA NO :

DRAWN : 26/11/2022 09:00:00

RECEIVED : 26/11/2022 09:00:21

REPORTED : 26/11/2022 16:30:11

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR : SELF

CLINICAL INFORMATION :

UID:12143432 REQNO-1325842

CORP-OPD

BILLNO-150122OPCR059788

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Test Report Status	Final	Results	Biological Reference Interval	Units
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SPECIALISED CHEMISTRY - HORMONE**THYROID PANEL, SERUM**

T3	109.8	80 - 200	ng/dL
METHOD : ELECTROCHEMILUMINESCENCE, COMPETITIVE IMMUNOASSAY			
T4	7.97	5.1 - 14.1	µg/dL
METHOD : ELECTROCHEMILUMINESCENCE, COMPETITIVE IMMUNOASSAY			
TSH (ULTRASENSITIVE)	1.590	0.270 - 4.200	µIU/mL
METHOD : ELECTROCHEMILUMINESCENCE, COMPETITIVE IMMUNOASSAY			

Interpretation(s)****End Of Report****Please visit www.srlworld.com for related Test Information for this accession


786

Dr. Swapnil Sirmukaddam
Consultant Pathologist



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PATIENT NAME : MRS. MRS.UTPALPARNA BHATTACHARJEEPATIENT ID : **FH.12143432**

CLIENT PATIENT ID : UID:12143432

ACCESSION NO : **0022VK005868**

AGE : 29 Years

SEX : Female

ABHA NO :

DRAWN : 26/11/2022 11:47:00

RECEIVED : 26/11/2022 12:04:49

REPORTED : 26/11/2022 13:39:27

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR :

CLINICAL INFORMATION :

UID:12143432 REQNO-1325842

CORP-OPD

BILLNO-150122OPCR059788

BILLNO-150122OPCR059788

Test Report Status	Final	Results	Biological Reference Interval	Units
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BIO CHEMISTRY**GLUCOSE, POST-PRANDIAL, PLASMA**

PPBS(POST PRANDIAL BLOOD SUGAR)

105

70 - 139

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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Dr.Akta Dubey

Consultant Pathologist

SRL Ltd

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



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Page 1 Of 1



Patient Ref. No. 2200000811301

12143432
29 Years

MRS Utpalparna bhattacharjee
Female

11/26/2022 11:14:13 AM

Rate 72 . Sinus rhythm.....normal P axis, V-rate 50- 99
Borderline short PR interval.....PR int <120ms

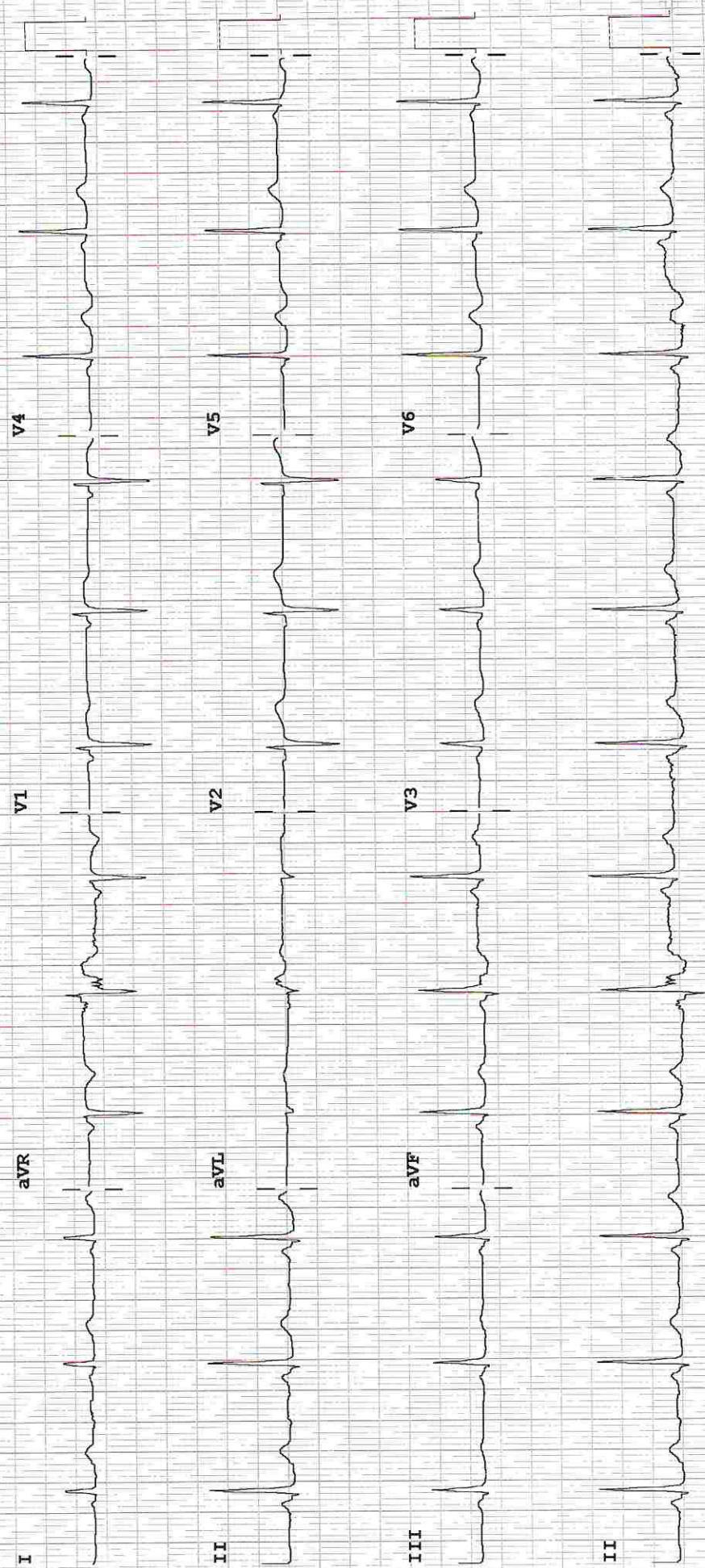
PR 116
QRSd 78
QT 364
QTc 399

--AXIS--
P 63
QRS 65
T 43

12 Lead; Standard Placement

- OTHERWISE NORMAL ECG -

Unconfirmed Diagnosis



Device:

Speed: 25 mm/sec

Limb: 10 mm/mV

Chest: 10.0 mm/mV

F 50~ 0.50-100 Hz W

100B CL

P?



DEPARTMENT OF NIC

Date: 26/Nov/2022

Name: Mrs. Utpalparna Bhattacharjee
Age | Sex: 29 YEAR(S) | Female
Order Station : FO-OPD
Bed Name :

UHID | Episode No : 12143432 | 59230/22/1501
Order No | Order Date: 1501/PN/OP/2211/125819 | 26-Nov-2022
Admitted On | Reporting Date : 26-Nov-2022 14:16:33
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	25	mm
AO Root	26	mm
AO CUSP SEP	18	mm
LVID (s)	24	mm
LVID (d)	33	mm
IVS (d)	08	mm
LVPW (d)	08	mm
RVID (d)	17	mm
RA	21	mm
LVEF	60	%

**DEPARTMENT OF NIC**

Date: 26/Nov/2022

Name: Mrs. Utpalparna Bhattacharjee

UHID | Episode No : 12143432 | 59230/22/1501

Age | Sex: 29 YEAR(S) | Female

Order No | Order Date: 1501/PN/OP/2211/125819 | 26-Nov-2022

Order Station : FO-OPD

Admitted On | Reporting Date : 26-Nov-2022 14:16:33

Bed Name :

Order Doctor Name : Dr.SELF.

DOPPLER STUDY:

E WAVE VELOCITY: 1.2 m/sec.

A WAVE VELOCITY:0.7 m/sec

E/A RATIO:1.7 , E/E'= 10

	PEAK (mmHg)	MEAN (mmHg)	V max (m/sec)	GRADE OF REGURGITATION
MITRAL VALVE	N			Nil
AORTIC VALVE	06			Nil
TRICUSPID VALVE	N			Nil
PULMONARY VALVE	01			Nil

Final Impression :

Normal 2 Dimensional and colour doppler echocardiography study.


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

Hiranandani Healthcare Pvt. Ltd.
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Emergency: 022 - 39199100 | Ambulance: 1255
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 RADIOLOGY

Date: 26/Nov/2022

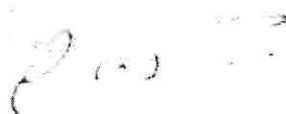
Name: Mrs. Utpalparna Bhattacharjee
Age | Sex: 29 YEAR(S) | Female
Order Station : FO-OPD
Bed Name :

UHID | Episode No : 12143432 | 59230/22/1501
Order No | Order Date: 1501/PN/OP/2211/125819 | 26-Nov-2022
Admitted On | Reporting Date : 26-Nov-2022 11:29:14
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)



DEPARTMENT OF RADIOLOGY

Date: 26/Nov/2022

Name: Mrs. Utpalparna Bhattacharjee
Age | Sex: 29 YEAR(S) | Female
Order Station : FO-OPD
Bed Name :

UHID | Episode No : 12143432 | 59230/22/1501
Order No | Order Date: 1501/PN/OP/2211/125819 | 26-Nov-2022
Admitted On | Reporting Date : 26-Nov-2022 11:43:09
Order Doctor Name : Dr.SELF .

US-WHOLE ABDOMEN

LIVER is normal in size (11.8 cm) and shows raised echogenicity. Intrahepatic portal and biliary systems are normal. No focal lesion is seen in liver. Portal vein appears normal.

GALL BLADDER is minimally distended.

SPLEEN is normal in size (11.0 cm) 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 3.8 cm.

Left kidney measures 11.2 x 3.3 cm.

PANCREAS: Head of pancreas appear unremarkable. Rest of the pancreas is obscured.

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 6.3 x 2.8 x 3.7 cm.

Endometrium measures 8.8 mm in thickness.

A fairly well-defined cyst of size 3.2 x 2.5 x 3.2 cm, volume 13.6 cc is noted in the right adnexa. Few internal echoes are noted within. Right ovary is not seen separately. No obvious vascularity is noted on color doppler evaluation.

Left ovary measures 2.6 x 1.3 x 2.5 cm, volume 4.8 cc.

No evidence of ascites.

IMPRESSION:

- Fatty infiltration of liver.
- Right adnexal cyst as described – possibility of hemorrhagic cyst is likely. Suggest: Clinical correlation / MRI pelvis for further evaluation if clinically indicated.

DR. YOGESH PATHADE
(MD Radio-diagnosis)



Cert. No. MC-2275



PATIENT NAME : MRS. UTPALPARNA BHATTACHARJEE

PATIENT ID : **FH.12143432**

CLIENT PATIENT ID : UID:12143432

ACCESSION NO : **0022VK005900**

AGE : 29 Years SEX : Female

ABHA NO :

DRAWN : 26/11/2022 13:13:00

RECEIVED : 26/11/2022 13:18:06

REPORTED : 28/11/2022 10:32:53

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR : SELF

CLINICAL INFORMATION :

UID:12143432 REQNO-1325842

CORP-OPD

BILLNO-150122OPCR059788

BILLNO-150122OPCR059788

Test Report Status **Final**

Units

CYTOLOGY

PAPANICOLAOU SMEAR

PAPANICOLAOU SMEAR

TEST METHOD

CONVENTIONAL GYNEC CYTOLOGY

SPECIMEN TYPE

TWO UNSTAINED CERVICAL SMEARS RECEIVED

REPORTING SYSTEM

2014 BETHESDA SYSTEM FOR REPORTING CERVICAL CYTOLOGY

SPECIMEN ADEQUACY

SATISFACTORY

METHOD : MICROSCOPIC EXAMINATION

MICROSCOPY

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

INTERPRETATION / RESULT

NEGATIVE FOR INTRAEPITHELIAL LESION OR MALIGNANCY

Comments

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

NO CYTOLOGICAL EVIDENCE OF HPV INFECTION IN THE SMEARS STUDIED.

****End Of Report****

Please visit www.srlworld.com for related Test Information for this accession

Dr.Akta Dubey

Consultant Pathologist

SRL Ltd
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NAVI MUMBAI, 400703
MAHARASHTRA, INDIA
Tel : 022-39199222,022-49723322,



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UHID	12143432	Date	26/11/2022		
Name	Mrs.Utpalparna Bhattacharjee	Sex	Female	Age	29
OPD	PAP	Health Check Up			

29yrs / Polo.

LMP: 5.11.22

PMC: 3/20d RMP

Drug allergy:
Sys illness:

Pap- cp / ng / (P) pap ✓

- Breast examⁿ (M)

Adv

- Fu c reports
- Pap smear Zysdy
- self breast examⁿ mthly

haha



UHID	12143432	Date	26/11/2022		
Name	Mrs. Utpalparna Bhattacharjee	Sex	Female	Age	29
OPD	Ophthal 141	Health Check Up			

Drug allergy:
 Sys illness:

O/C →
 O/H →
 M/H →

OPB ophthal.

No ocular c/o at present

fo: A/S; wrc

Immunus
 exam done

AVN, 6/6
 6/6.

Rx.

Vn	R →	-4.00 sph	6/6 6/6	Psychology PSCY/13/13 17/17 α/ooth.
	L →	-3.75 - 0.50 X 150		

A

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 GST IN: 27AABCH5894D1ZG | PAN NO: AABCH5894D



Hiranandani
 HOSPITAL

A Fortis Network Hospital

UHID	12143432	Date	26/11/2022	
Name	Mrs. Utpalparna Bhattacharjee	Sex	Female	Age 29
OPD	Dental 12	Health Check Up		

Drug allergy:
 Sys illness:

Caries $\frac{6}{7}$

Impacted $\frac{8}{8 \sqrt{8}}$

Stains 7+

Caries 7+

Treatment

Adv. surgical removal $\frac{8}{8 \sqrt{8}}$

Adv. filling $\frac{6}{7}$

Adv. oral prophylaxis

Adv. OPG

Dr. Diksha Kaka

PAP

PATIENT NAME : MRS. MRS.UTPALPARNA BHATTACHARJEE

PATIENT ID : **FH.12143432**

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CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR : SELF

CLINICAL INFORMATION :

UID:12143432 REQNO-1325842

CORP-OPD

BILLNO-150122OPCR059788

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KIDNEY PANEL - 1

BLOOD UREA NITROGEN (BUN), SERUM

BLOOD UREA NITROGEN 7 6 - 20 mg/dL

METHOD : UREASE - UV

CREATININE EGFR- EPI

CREATININE 0.63 0.60 - 1.10 mg/dL

METHOD : ALKALINE PICRATE KINETIC JAFFES

AGE 29 years

GLOMERULAR FILTRATION RATE (FEMALE) 123.07 Refer Interpretation Below mL/min/1.73m²

METHOD : CALCULATED PARAMETER

BUN/CREAT RATIO

BUN/CREAT RATIO 11.11 5.00 - 15.00

METHOD : CALCULATED PARAMETER

URIC ACID, SERUM

URIC ACID 4.3 2.6 - 6.0 mg/dL

METHOD : URICASE UV

TOTAL PROTEIN, SERUM

TOTAL PROTEIN 7.3 6.4 - 8.2 g/dL

METHOD : BIURET

ALBUMIN, SERUM

ALBUMIN 3.7 3.4 - 5.0 g/dL

METHOD : BCP DYE BINDING

GLOBULIN

GLOBULIN 3.6 2.0 - 4.1 g/dL

METHOD : CALCULATED PARAMETER

ELECTROLYTES (NA/K/CL), SERUM

SODIUM, SERUM 137 136 - 145 mmol/L

METHOD : ISE INDIRECT

POTASSIUM, SERUM 4.27 3.50 - 5.10 mmol/L

METHOD : ISE INDIRECT

CHLORIDE, SERUM 103 98 - 107 mmol/L

METHOD : ISE INDIRECT

Interpretation(s)

PHYSICAL EXAMINATION, URINE

SRL Ltd
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MAHARASHTRA, INDIA
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PATIENT ID : **FH.12143432** CLIENT PATIENT ID : UID:12143432
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COLOR		PALE YELLOW		
METHOD : PHYSICAL				
APPEARANCE		CLEAR		
METHOD : VISUAL				
CHEMICAL EXAMINATION, URINE				
PH		7.0	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				
MICROSCOPIC EXAMINATION, URINE				
RED BLOOD CELLS		NOT DETECTED	NOT DETECTED	/HPF
METHOD : MICROSCOPIC EXAMINATION				
PUS CELL (WBC'S)		0-1	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION				
EPITHELIAL CELLS		2-3	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION				
CASTS		NOT DETECTED		
METHOD : MICROSCOPIC EXAMINATION				
CRYSTALS		NOT DETECTED		
METHOD : MICROSCOPIC EXAMINATION				



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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 URINARY

Interpretation(s)**Interpretation(s)**

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-

Serum 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's 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.

SRL Ltd

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PATIENT NAME : MRS. MRS.UTPALPARNA BHATTACHARJEE

PATIENT ID : **FH.12143432** CLIENT PATIENT ID : UID:12143432
 ACCESSION NO : **0022VK005774** AGE : 29 Years SEX : Female ABHA NO :
 DRAWN : 26/11/2022 09:00:00 RECEIVED : 26/11/2022 09:00:21 REPORTED : 26/11/2022 12:46:21
 CLIENT NAME : **FORTIS VASHI-CHC -SPLZD** REFERRING DOCTOR : SELF

CLINICAL INFORMATION :

UID:12143432 REQNO-1325842
 CORP-OPD
 BILLNO-150122OPCR059788
 BILLNO-150122OPCR059788

Test Report Status	Final	Results	Biological Reference Interval
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HAEMATOLOGY

ERYTHROCYTE SEDIMENTATION RATE (ESR), WHOLE BLOOD

E.S.R **25** High 0 - 20 mm at 1 hr
 METHOD : WESTERGREN METHOD

CBC-5, EDTA WHOLE BLOOD

BLOOD COUNTS, EDTA WHOLE BLOOD

HEMOGLOBIN (HB) 12.9 12.0 - 15.0 g/dL
 METHOD : SPECTROPHOTOMETRY
 RED BLOOD CELL (RBC) COUNT 4.06 3.8 - 4.8 mil/ μ L
 METHOD : ELECTRICAL IMPEDANCE
 WHITE BLOOD CELL (WBC) COUNT 5.33 4.0 - 10.0 thou/ μ L
 METHOD : DOUBLE HYDRODYNAMIC SEQUENTIAL SYSTEM(DHSS)CYTOMETRY
 PLATELET COUNT **142** Low 150 - 410 thou/ μ L
 METHOD : ELECTRICAL IMPEDANCE

RBC AND PLATELET INDICES

HEMATOCRIT (PCV) 37.8 36 - 46 %
 METHOD : CALCULATED PARAMETER
 MEAN CORPUSCULAR VOLUME (MCV) 93.1 83 - 101 fL
 METHOD : CALCULATED PARAMETER
 MEAN CORPUSCULAR HEMOGLOBIN (MCH) 31.8 27.0 - 32.0 pg
 METHOD : CALCULATED PARAMETER
 MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION(MCHC) 34.2 31.5 - 34.5 g/dL
 METHOD : CALCULATED PARAMETER
 RED CELL DISTRIBUTION WIDTH (RDW) **14.7** High 11.6 - 14.0 %
 METHOD : CALCULATED PARAMETER
 MENTZER INDEX 22.9
 MEAN PLATELET VOLUME (MPV) 9.8 6.8 - 10.9 fL
 METHOD : CALCULATED PARAMETER

WBC DIFFERENTIAL COUNT

NEUTROPHILS **36** Low 40 - 80 %
 METHOD : FLOW CYTOMETRY
 LYMPHOCYTES **54** High 20 - 40 %
 METHOD : FLOW CYTOMETRY

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Patient Ref. No. 2200000081120

PATIENT NAME : MRS. MRS.UTPALPARNA BHATTACHARJEE

PATIENT ID : **FH.12143432** CLIENT PATIENT ID : UID:12143432
 ACCESSION NO : **0022VK005774** AGE : 29 Years SEX : Female ABHA NO :
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MONOCYTES		10	2 - 10 %
METHOD : FLOW CYTOMETRY			
EOSINOPHILS		0	Low 1 - 6 %
METHOD : FLOW CYTOMETRY			
BASOPHILS		0	0 - 2 %
METHOD : FLOW CYTOMETRY			
ABSOLUTE NEUTROPHIL COUNT		1.92	Low 2.0 - 7.0 thou/μL
METHOD : CALCULATED PARAMETER			
ABSOLUTE LYMPHOCYTE COUNT		2.88	1.0 - 3.0 thou/μL
METHOD : CALCULATED PARAMETER			
ABSOLUTE MONOCYTE COUNT		0.53	0.2 - 1.0 thou/μL
METHOD : CALCULATED PARAMETER			
ABSOLUTE EOSINOPHIL COUNT		0.00	Low 0.02 - 0.50 thou/μL
METHOD : CALCULATED PARAMETER			
ABSOLUTE BASOPHIL COUNT		0	Low 0.02 - 0.10 thou/μL
METHOD : CALCULATED PARAMETER			
NEUTROPHIL LYMPHOCYTE RATIO (NLR)		0.6	
METHOD : CALCULATED PARAMETER			
MORPHOLOGY			
RBC		PREDOMINANTLY NORMOCYTIC NORMOCHROMIC	
METHOD : MICROSCOPIC EXAMINATION			
WBC		NORMAL MORPHOLOGY	
METHOD : MICROSCOPIC EXAMINATION			
PLATELETS		REDUCED	
METHOD : MICROSCOPIC EXAMINATION			

Interpretation(s)

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

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

TEST INTERPRETATION

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

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

In pregnancy BRI in first trimester is 0-48 mm/hr(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

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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)

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.
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.

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

ABO GROUP

TYPE O

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.

BIO CHEMISTRY**LIVER FUNCTION PROFILE, SERUM**

BILIRUBIN, TOTAL	1.12	High 0.2 - 1.0	mg/dL
METHOD : JENDRASSIK AND GROFF			
BILIRUBIN, DIRECT	0.22	High 0.0 - 0.2	mg/dL
METHOD : JENDRASSIK AND GROFF			
BILIRUBIN, INDIRECT	0.90	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

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Patient Ref. No. 22000000811207

PATIENT NAME : MRS. MRS.UTPALPARNA BHATTACHARJEE

PATIENT ID : **FH.12143432**

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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)	28	15 - 37	U/L
METHOD : UV WITH PSP			
ALANINE AMINOTRANSFERASE (ALT/SGPT)	33	< 34.0	U/L
METHOD : UV WITH PSP			
ALKALINE PHOSPHATASE	83	30 - 120	U/L
METHOD : PNPP-ANP			
GAMMA GLUTAMYL TRANSFERASE (GGT)	21	5 - 55	U/L
METHOD : GAMMA GLUTAMYL CARBOXY 4NITROANILIDE			
LACTATE DEHYDROGENASE	217	High 100 - 190	U/L
METHOD : LACTATE -PYRUVATE			

LIPID PROFILE, SERUM

CHOLESTEROL, TOTAL	164	< 200 Desirable 200 - 239 Borderline High >= 240 High	mg/dL
METHOD : ENZYMATIC/COLORIMETRIC, CHOLESTEROL OXIDASE, ESTERASE, PEROXIDASE			
TRIGLYCERIDES	38	< 150 Normal 150 - 199 Borderline High 200 - 499 High >=500 Very High	mg/dL
METHOD : ENZYMATIC ASSAY			
HDL CHOLESTEROL	55	< 40 Low >=60 High	mg/dL
METHOD : DIRECT MEASURE - PEG			
LDL CHOLESTEROL, DIRECT	98	< 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			

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CHOL/HDL RATIO	3.0	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	1.8		0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk
METHOD : CALCULATED PARAMETER			
VERY LOW DENSITY LIPOPROTEIN	7.6		</= 30.0 mg/dL
METHOD : CALCULATED PARAMETER			

GLUCOSE FASTING, FLUORIDE PLASMA

FBS (FASTING BLOOD SUGAR)	89		74 - 99 mg/dL
METHOD : HEXOKINASE			

GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD

HBA1C	4.9		Non-diabetic: < 5.7 % Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 ADA Target: 7.0 Action suggested: > 8.0
METHOD : HB VARIANT (HPLC)			
ESTIMATED AVERAGE GLUCOSE(EAG)	93.9		< 116.0 mg/dL
METHOD : CALCULATED PARAMETER			

Interpretation(s)

LIVER FUNCTION PROFILE, SERUM-
LIVER FUNCTION PROFILE

Bilirubin is a yellowish pigment found in bile and is a breakdown product of normal heme catabolism. Bilirubin is excreted in bile and urine, and elevated levels may give yellow discoloration in jaundice. Elevated levels result 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, Paget's disease, Rickets, Sarcoidosis etc. Lower-than-normal ALP levels seen in Hypophosphatasia, Malnutrition, Protein deficiency, Wilson's 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

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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.Serum 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,Waldenstrom's disease.Lower-than-normal levels may be due to: Agammaglobulinemia,Bleeding (hemorrhage),Burns,Glomerulonephritis,Liver disease, Malabsorption,Malnutrition,Nephrotic syndrome,Protein-losing enteropathy etc.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

LIPID PROFILE, SERUM-Serum cholesterol is a blood test that can provide valuable information for the risk of coronary artery disease This test can help determine your risk of the build up of plaques in your arteries that can lead to narrowed or blocked arteries throughout your body (atherosclerosis). High cholesterol levels usually don't cause any signs or symptoms, so a cholesterol test is an important tool. High cholesterol levels often are a significant risk factor for heart disease and important for diagnosis of hyperlipoproteinemia, atherosclerosis, hepatic and thyroid diseases.

Serum Triglyceride are a type of fat in the blood. When you eat, your body converts any calories it doesn't need into triglycerides, which are stored in fat cells. High triglyceride levels are associated with several factors, including being overweight, eating too many sweets or drinking too much alcohol, smoking, being sedentary, or having diabetes with elevated blood sugar levels. Analysis has proven useful in the diagnosis and treatment of patients with diabetes mellitus, nephrosis, liver obstruction, other diseases involving lipid metabolism, and various endocrine disorders. In conjunction with high density lipoprotein and total serum cholesterol, a triglyceride determination provides valuable information for the assessment of coronary heart disease risk.It is done in fasting state.

High-density lipoprotein (HDL) cholesterol. This is sometimes called the "good" cholesterol because it helps carry away LDL cholesterol, thus keeping arteries open and blood flowing more freely.HDL cholesterol is inversely related to the risk for cardiovascular disease. It increases following regular exercise, moderate alcohol consumption and with oral estrogen therapy. Decreased levels are associated with obesity, stress, cigarette smoking and diabetes mellitus.

SERUM LDL The small dense LDL test can be used to determine cardiovascular risk in individuals with metabolic syndrome or established/progressing coronary artery disease, individuals with triglyceride levels between 70 and 140 mg/dL, as well as individuals with a diet high in trans-fat or carbohydrates. Elevated sdLDL levels are associated with metabolic syndrome and an 'atherogenic lipoprotein profile', and are a strong, independent predictor of cardiovascular disease. Elevated levels of LDL arise from multiple sources. A major factor is sedentary lifestyle with a diet high in saturated fat. Insulin-resistance and pre-diabetes have also been implicated, as has genetic predisposition. Measurement of sdLDL allows the clinician to get a more comprehensive picture of lipid risk factors and tailor treatment accordingly. Reducing LDL levels will reduce the risk of CVD and MI.

Non HDL Cholesterol - Adult treatment panel ATP III suggested the addition of Non-HDL Cholesterol as an indicator of all atherogenic lipoproteins (mainly LDL and VLDL). NICE guidelines recommend Non-HDL Cholesterol measurement before initiating lipid lowering therapy. It has also been shown to be a better marker of risk in both primary and secondary prevention studies.

Recommendations:
 Results of Lipids should always be interpreted in conjunction with the patient's medical history, clinical presentation and other findings.

NON FASTING LIPID PROFILE includes Total Cholesterol, HDL Cholesterol and calculated non-HDL Cholesterol. It does not include triglycerides and may be best used in patients for whom fasting is difficult.

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:
 Hypoglycemia is defined as a glucose of < 50 mg/dL in men and < 40 mg/dL in women. While random serum glucose levels correlate with home glucose monitoring results (weekly mean capillary glucose values), there is wide fluctuation within individuals.Thus, glycosylated hemoglobin(HbA1c) levels are favored to monitor glycemic control. High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glycosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc. **GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD-Used For:**

- 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 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

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