

Name Mrs. Jyoti Bansal
UHID : 12362463 Date : 20/3/23
Age : 34 Gender : F

Nursing Assessment

Profile	
Height (cm) : <u>160 cm</u>	Waist Circumference (cm) : <u>32 inches</u>
Weight (Kg.) : <u>71 Kg</u>	Body Mass Index : <u>27.7 kg/m² / 118-231</u>
Occupation :	Marital Status <input type="checkbox"/> Single <input checked="" type="checkbox"/> Married << <u>59104</u>

Vital Signs	
Pulse Rate (/min) : <u>112 6/min</u>	Respiratory Rate (/min) : <u>SP0₂ - 98%</u>
Blood Pressure (mmHg) : <u>110/80 mmHg</u>	Temperature (if febrile) : <u>Afebrile</u>

Past History	
<input checked="" type="checkbox"/> Hypertension :	<input checked="" type="checkbox"/> Diabetes :
<input checked="" type="checkbox"/> Heart disease :	<input checked="" type="checkbox"/> Dyslipidemia :
<input checked="" type="checkbox"/> Asthma :	<input type="checkbox"/> Tuberculosis :
<input checked="" type="checkbox"/> Allergies :	
<input checked="" type="checkbox"/> Others :	

For Women	
LMP:	Last Pap smear done in
Menopause <input type="checkbox"/> Yes <input type="checkbox"/> No	Last Mammography done in
Consent for X-ray & Mammography	

Current Medications
<u>NA</u>

Signature, Name and Emp. ID of the Nurse : _____

Name: Mrs. Jyoti Bansal
 UHID: 12362463 Date: 20/3/23
 Age: 34 Gender: F

Internal Medicine Consultation

Relevant History:

- No complaints
- No medicine
- mild cough x 2d

Examination Findings:

- Rest WNL

B+ve

Investigations:

TSM - 4.790
 Hb - 10.4 (↓ per, mcv, MCH)
 - 50 mm
 FBS - 88 HbA1c - 5.5 %
 PP - 114
 Lipid (P) - n
 RFT | con
 2FT | Echo
 S.E | n
 stool Pk | n
 urine Pk | n
 ECG - n
 USK - fatty liver gr I.

Diagnosis: 1. subclinical Hypothyroidism.
 2. Anemia
 3. Fatty liver gr I

25-OH-D 4. OVERWEIGHT

Serum iron studies / Ferritin / B-12 / folate
 3. ? MRI

Advice / Treatment Plan:

- Regular Exercise
- Tab. Zentle stat
- Tab. ROTHERA 4 - (WS)
(empty stomach)
- Tab. Azithromycin 500 - WS
x 5d
- Acnesol topical - BS
- weight Reduction

WILCO R4L
 2 - 1

Dr. MANJEET SINGH TILHAN
 MBS (MBBS)
 Additional Director - Internal Medicine (FMC)
 Fortis Hospital, Mohali (Pb.)
 Mobile No. 9814104609
 Reg. No. PMC 24797

Signature and stamp of the Consultant

Name Mrs. Jyoti Bansal
 UHID : 12362463 Date : 20/3/23
 Age : 34 Gender : F

Ophthalmology Consultation

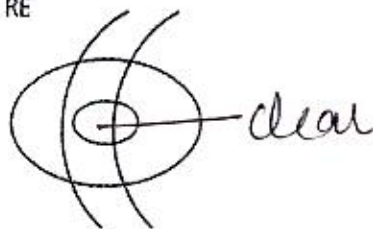
History: NIL

Examination findings:

Visual acuity $\left\{ \begin{array}{l} R \ 9 \\ L \ 6/6 \end{array} \right.$ Visual acuity with glasses $\left\{ \begin{array}{l} R \\ L \end{array} \right.$ Colour Vision $\left\{ \begin{array}{l} R \ 6/6 \\ L \ 6/6 \end{array} \right.$

Slit Lamp Examination

RE

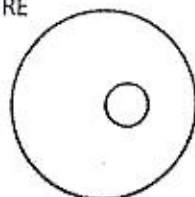


LE

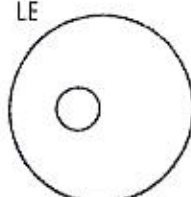


Fundus Examination

RE



LE



Diagnosis: NAD BE

Treatment: G. Refresh tears 000

Spectacle prescription:

Right eye

	SPH	CYL	AXIS	VA
Distance				6/6P
Near	Plano			N/G

Left eye

	SPH	CYL	AXIS	VA
Distance				6/6
Near	Plano			N/G

Signature and stamp of the Ophthalmologist : _____

Name Mr. Jyoti Baneal.
UHID : 12368463 Date : 20/3/2023
Age : 34 years

20/3/23

Gynecology Consultation

Symptoms: wants check up

Diagnosis:

Obstetric History:

OH = 2 FTND 1 abortion 1/2 yr
20 = 8 yr old Not taking any OCPs

Advice / Treatment Plan:

Menstrual History:

MH = 5 days
30-35 dup = 16 Feb

Past & Family History:

Not taking any medication for illness,
No sig family history.

Examination Findings:

• Breast:

• P/A: Left no palpable mass
no tenderness anywhere

• P/S: CO → mild erosion
vagn N

• PV: CO & ut ACAT r/s for

for free

Investigations:

Pap's Smear sent

Signature and stamp of the Consultant :

Dr. Santosh Yadav
MBBS, MS (Obs. & Gynae)
Empanelled Consultant-Gynaecology
Reg. No. RMC 7446
Mobile: 94140 45452
Fortis MEDCENTRE (A unit of Fortis Hospital, Mohali)
S.C.O. 11, Sector 11-D, Chandigarh-160011 (INDIA)
Phone No 0172-5061722, 5055441

Female

20.03.2023 11:33:39
Fortis Med Centre
sector 11
Ghandgarh

Technician:
Ordering Ph:
Referring Ph:
Attending Ph:

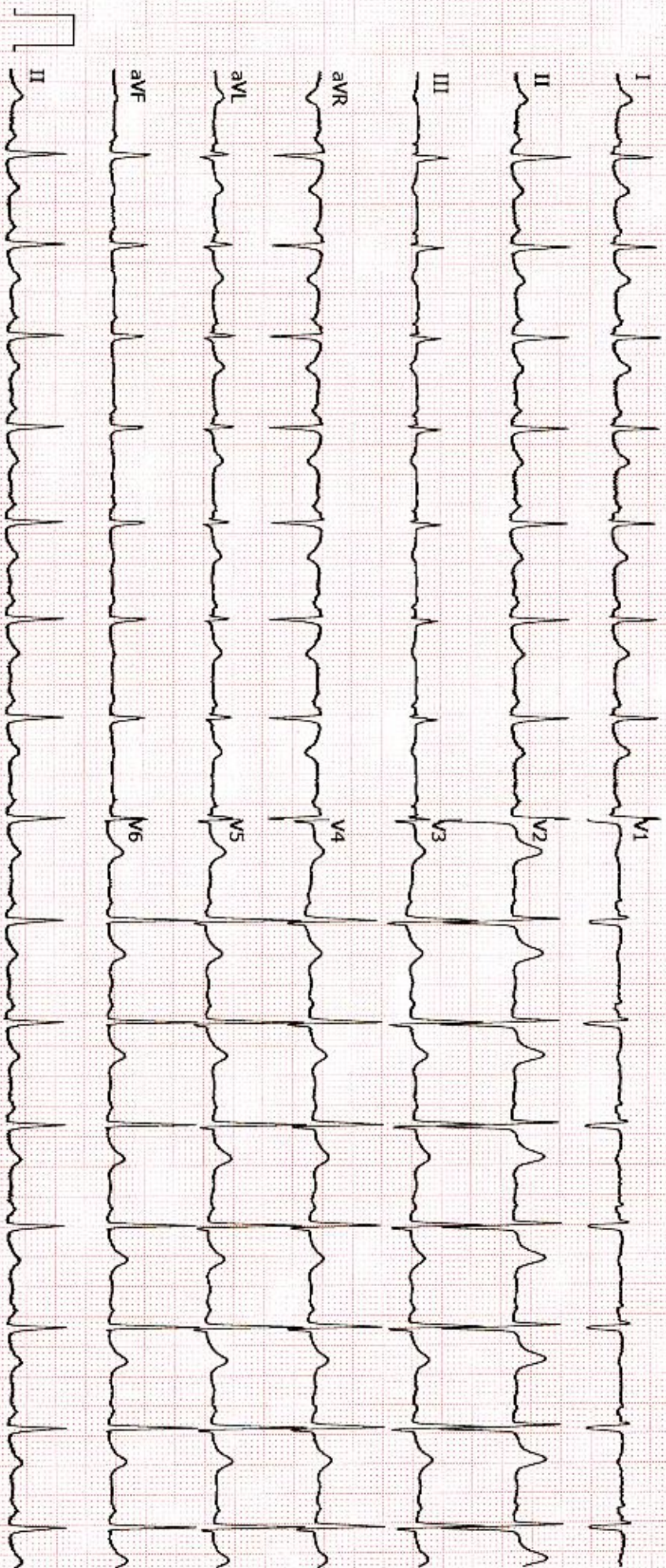
Location:
Order Number:
Visit:
Indication:
Medication 1:
Medication 2:
Medication 3:

Room:

92 bpm
--/-- mmHg

QRS : 70 ms
QT / QTcBaz : 346 / 427 ms
PR : 134 ms
P : 110 ms
RR / pp : 654 / 652 ms
P / QRS / T : 31 / 47 / 19 degrees

Normal sinus rhythm
Normal ECG



NAME: JOYTI BANSAL**AGE AND SEX: 34/F****UHID :12362463****DATE-20/03/2023****CHEST- PA**

Both the domes of diaphragm are normal.

Both costophrenic angles are normal.

Both lung fields are clear.

Cardiac size and silhouette are normal.

Both hila and mediastinum are normal.

Bony cage and soft tissues are normal.

IMPRESSION : NORMAL STUDY.**Please correlate clinically and with other relevant investigations.****DR NEHA CHHABRA**
CONSULTANT RADIOLOGIST

NAME: MRS. JYOTI BANSAL**AGE AND SEX:34 Y/F****UHID NO:12362463****DAT: 20/03/2023****ROI: WHOLE ABDOMEN**

Liver is normal in size, outline and shows mildly increased echogenicity. No focal lesion seen. IHBR's are not dilated. Portal vein and hepatic veins are normal.

Gall bladder is normally distended with anechoic lumen. Wall thickness is normal. No calculus / focal lesion seen. No pericholecystic fluid / collection seen. CBD is normal.

Pancreas is visualized in region of head and proximal body and is normal in size, shape, outline and echotexture. No focal lesion seen. Distal body and tail are obscured by bowel gases

Spleen is normal in size, outline and echotexture. No focal lesion

Right kidney is normal in size, outline and echogenicity. Cortico-medullary differentiation is maintained. No hydronephrosis / calculus is seen

Left kidney is normal in size, outline and echogenicity. Cortico-medullary differentiation is maintained. No hydronephrosis / calculus is seen.

Retroperitoneum is normal.

The urinary bladder is fully distended with normal outline and wall thickness. No calculus / SOL seen.

Uterus is normal in size, shape and outline. Endometrium measures 11.5 mm. No SOL seen.

Bilateral ovaries are normal (LO -11.5 cc, RO-8.5 cc) in size, shape and echotexture.

No free fluid is seen in POD.

Opinion: Fatty Liver Grade I**Suggested clinical correlation.**
Dr. NEHA CHHABRA.
Consultant Radiologist

JYOTI 34F

Study Date: 20/03/2023

Patient ID: 52031020230320

Accession #:

Alt ID:

DOB:

Age:

Gender: F

Ht:

Wt:

BSA:

Institution: Fortis MEDCENTRE, Chandigarh

Referring Physician:

Physician of Record:

Performed By:

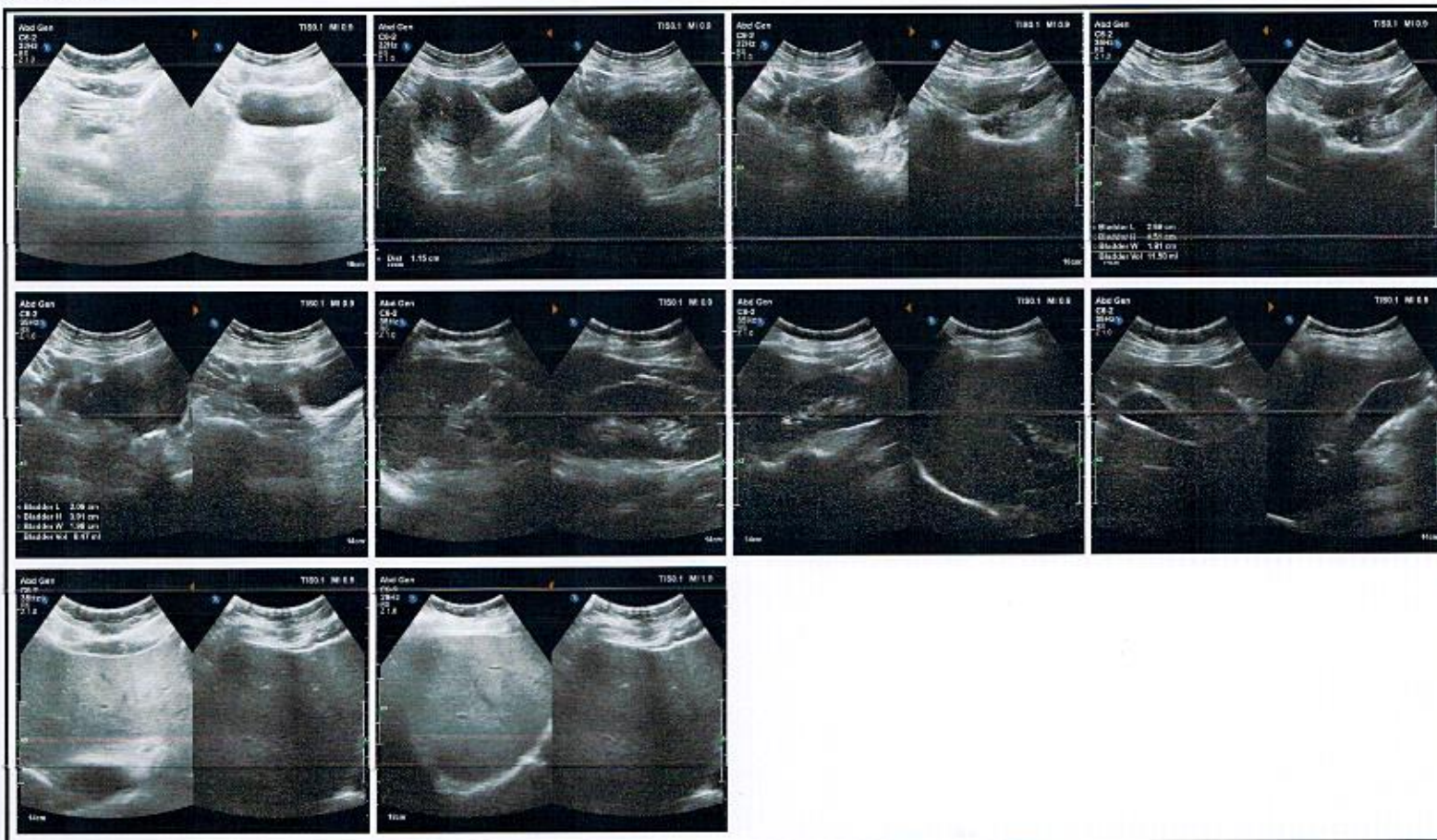
Comments:

Abdominal: Measurements and Calculations

2D Abdominal Organs and Vessels

Bladder Vol	7.92 ml	Bladder H	1.72 cm
Bladder L	4.44 cm	Bladder W	1.98 cm

Images



Signature

Signature:

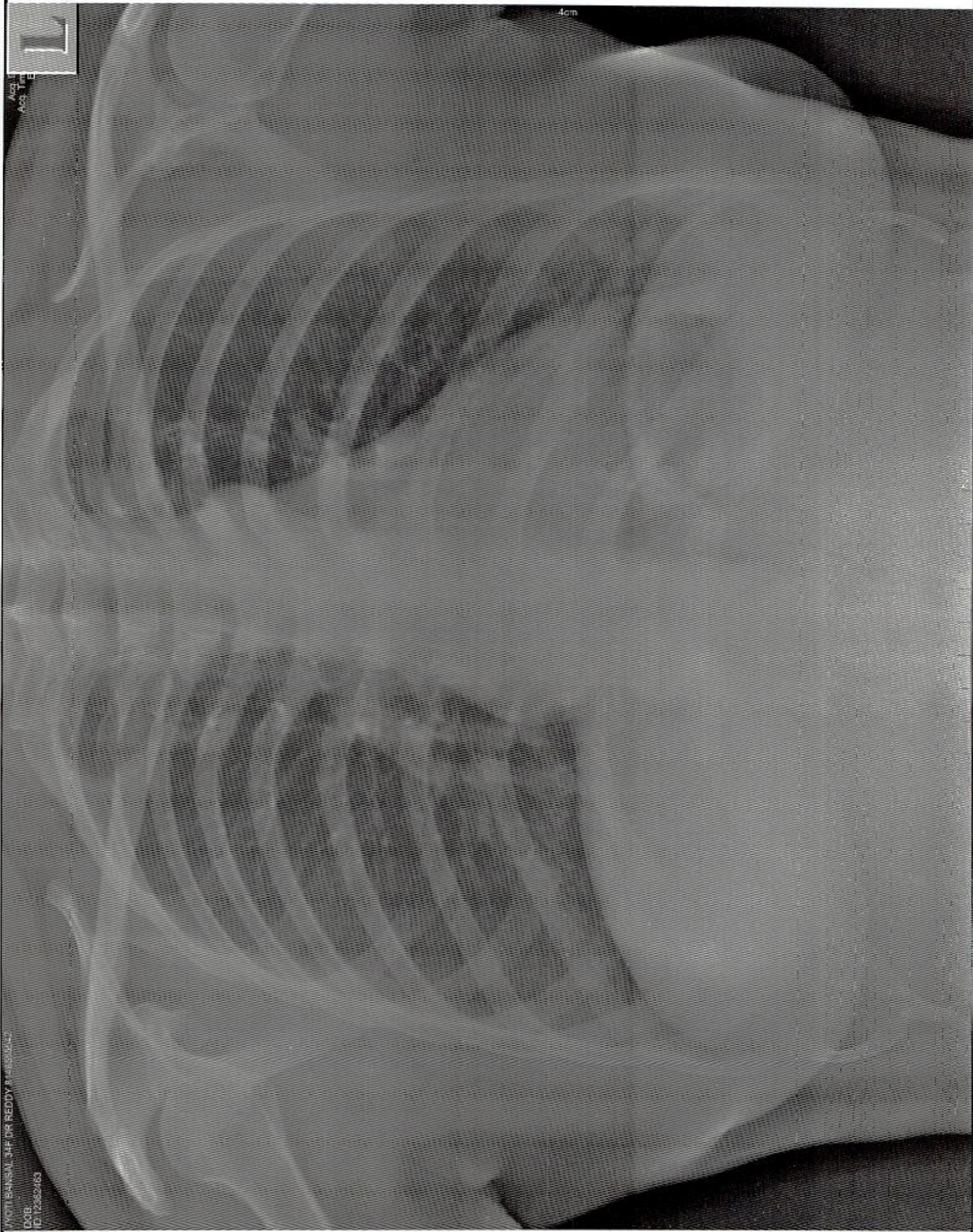
Name(Print):

Date:

L

4cm

JYOTI BANSAI, JAF DR REDDY #14593642
DOB: [REDACTED]
ID: 12382463



DEPARTMENT OF CARDIOLOGY
ECHOCARDIOGRAPHY LABORATORY
Phone 0172-5061222; Ext. 6422

Dated: 21 March 2023

Name: MRS JYOTI BANSAL Age: 34 Sex: FEMALE
FHL No 12362463 Lab No:
Clinical Diagnosis: R/O CAD
Ref By: FMC

MEASUREMENTS

Aortic Root Diameter	:	2.7	cm	Left Atrial dimension	2.7	cm
Aortic Valve Opening	:	---	cm	Right Ventricular dimension	1.2	cm
Left Ventricular ED dimension	:	3.7	cm	Left Ventricular ES dimension	2.5	cm
Interventricular Septal thickness	ED:	1.0	cm	ES:	1.1	cm
Left Ventricular PW thickness	ED:	0.8	cm	ES:	1.5	cm

INDICES OF LEFT VENTRICULAR FUNCTION:

LV Ejection Fraction : 64 %

IMAGING:

M mode examination revealed normal movement of both Mitral leaflets during diastole. No SAM or Mitral valve prolapse is seen. Aortic root is normal in size. Dimensions of left atrium and left ventricle are normal

2-D imaging in PLAX, SAX and apical views revealed normal sized left ventricle. Movement of anterior wall, septum, apex, inferior wall, posterior and lateral walls is normal. Mitral valve opening is normal. No evidence of Mitral valve prolapse is seen. Aortic valve has three cusps and its opening is not restricted. Pulmonary valve is normal. Interatrial and interventricular septa are intact. No intracardiac mass or thrombus is seen. No pericardial pathology is observed.

**DEPARTMENT OF CARDIOLOGY
ECHOCARDIOGRAPHY LABORATORY
Phone 0172-5061222; Ext. 6422****DOPPLER: PULSE WAVE; CONTINUOUS WAVE & COLOR FLOW MAPPING**

Mitral Valve : E= 76 A= 60 cm/sec; E > A; No MR
E wave Deceleration Time = 173 msec

Aortic Valve : 140 cm/sec No AR

Tricuspid Valve : No TR ; RVSP = + RAP mmHg

Pulmonary Valve : 93 cm/sec

FINAL DIAGNOSIS

- NO REGIONAL WALL MOTION ABNORMALITY OF LEFT VENTRICLE
- NORMAL LEFT VENTRICULAR SYSTOLIC FUNCTION (LVEF 64%)



Dr. MUKTI SHARMA
MD, DNB, FIAP, FCSI
Sr. Consultant
Fortis MEDCENTRE

PATIENT NAME : JYOTI BANSAL**REF. DOCTOR : SELF**

CODE/NAME & ADDRESS : C000138383

ACCESSION NO : **0080WC007764**

AGE/SEX : 34 Years Female

PROVISIONAL REPORT

PATIENT ID : JYOTF13088880

DRAWN :

CLIENT PATIENT ID:

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Test Report Status	Final	Results	Biological Reference Interval	Units
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MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE**THYROID PANEL, SERUM**

T3	124.3	80.00 - 200.00	ng/dL
METHOD : COMPETITIVE (ECLIA)			
T4	13.26	5.10 - 14.10	µg/dL
METHOD : COMPETITIVE (ECLIA)			
TSH (ULTRASENSITIVE)	4.790 High	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

Repeat x 1 month

METHOD : SANDWICH (ECLIA)

Interpretation(s)**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

MICROSCOPY

SMEARS SHOW ADEQUATE CELLULARITY COMPOSED PREDOMINANTLY OF INTERMEDIATE SQUAMOUS EPITHELIAL CELLS ALONG WITH FEW SUPERFICIAL SQUAMOUS EPITHELIAL CELLS IN A BACKGROUND OF POLYMORPHS AND BLOOD. ENDOCERVICAL CELLS SEEN. NO EVIDENCE OF MALIGNANCY SEEN.

INTERPRETATION / RESULT

NEGATIVE FOR INTRAEPITHELIAL LESION OR MALIGNANCY

REACTIVE CELLULAR CHANGES ASSOCIATED WITH INFLAMMATION.

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Tel : 9111591115,
CIN - U74899PB1995PLC045956

Patient Ref. No. 8000001394826

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LETTER

REQUEST LETTER

CX/191/2023



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HAEMATOLOGY - CBC

MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE

BLOOD COUNTS, EDTA WHOLE BLOOD

HEMOGLOBIN (HB)	10.4 Low	12.0 - 15.0	g/dL
METHOD : CYANMETHEMOGLOBIN METHOD			
RED BLOOD CELL (RBC) COUNT	3.89	3.8 - 4.8	mil/ μ L
WHITE BLOOD CELL (WBC) COUNT	5.60	4.0 - 10.0	thou/ μ L
PLATELET COUNT	259	150 - 410	thou/ μ L

RBC AND PLATELET INDICES

HEMATOCRIT (PCV)	31.8 Low	36.0 - 46.0	%
MEAN CORPUSCULAR VOLUME (MCV)	81.7 Low	83.0 - 101.0	fL
METHOD : DERIVED PARAMETER FROM RBC HISTOGRAM			
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	26.8 Low	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)	15.5 High	11.6 - 14.0	%
METHOD : CALCULATED PARAMETER			
MENTZER INDEX	21.0		
MEAN PLATELET VOLUME (MPV)	8.5	6.8 - 10.9	fL
METHOD : DERIVED PARAMETER FROM PLATELET HISTOGRAM			

WBC DIFFERENTIAL COUNT

NEUTROPHILS	67	40 - 80	%
METHOD : LIGHT ABSORBANCE OF CYTCHEMICAL STAINED CELLS IMPEDENCE			
LYMPHOCYTES	24	20 - 40	%
METHOD : LIGHT ABSORBANCE OF CYTCHEMICAL STAINED CELLS IMPEDENCE			
MONOCYTES	4	2.0 - 10.0	%
METHOD : LIGHT ABSORBANCE OF CYTCHEMICAL STAINED CELLS IMPEDENCE			
EOSINOPHILS	5	1.0 - 6.0	%
BASOPHILS	0	0 - 1	%
METHOD : LIGHT ABSORBANCE OF CYTCHEMICAL STAINED CELLS IMPEDENCE			
ABSOLUTE NEUTROPHIL COUNT	3.75	2.0 - 7.0	thou/ μ L
ABSOLUTE LYMPHOCYTE COUNT	1.34	1.0 - 3.0	thou/ μ L

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ABSOLUTE MONOCYTE COUNT		0.22	0.2 - 1.0	thou/ μ L
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ABSOLUTE EOSINOPHIL COUNT		0.28	0.02 - 0.50	thou/ μ L
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ABSOLUTE BASOPHIL COUNT		0.00 Low	0.02 - 0.10	thou/ μ L
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METHOD : CALCULATED PARAMETER

NEUTROPHIL LYMPHOCYTE RATIO (NLR)		2.8		
-----------------------------------	--	-----	--	--

METHOD : CALCULATED PARAMETER

Interpretation(s)

BLOOD COUNTS, EDTA WHOLE BLOOD-The cell morphology is well preserved for 24hrs. However after 24-48 hrs a progressive increase in MCV and HCT is observed leading to a decrease in MCHC. A direct smear is recommended for an accurate differential count and for examination of RBC morphology.

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.

(<1.5) in patients with microcytic anaemia. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HbA2 remains the gold standard for diagnosing a case of beta thalassaemia trait.

WBC DIFFERENTIAL COUNT-The optimal threshold of 3.3 for NLR showed a prognostic possibility of clinical symptoms to change from mild to severe in COVID positive patients. When age = 49.5 years old and NLR = 3.3, 46.1% COVID-19 patients with mild disease might become severe. By contrast, when age < 49.5 years old and NLR < 3.3, COVID-19 patients tend to show mild disease.

(Reference to - The diagnostic and predictive role of NLR, d-NLR and PLR in COVID-19 patients ; A.-P. Yang, et al.; International Immunopharmacology 84 (2020) 106504 This ratio element is a calculated parameter and out of NABL scope.

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HAEMATOLOGY

MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE

ERYTHROCYTE SEDIMENTATION RATE (ESR),WHOLE BLOOD

E.S.R	50 High	0 - 20	mm at 1 hr
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METHOD : MODIFIED WESTERGREN

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.

TESY INTERPRETATION

Increase in: Infections, Vasculitis, 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 0-45 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: Polikilocytosis, (Sickle Cells, spherocytes), Microcytosis, Low fibrinogen, Very high WBC counts, Drugs (Quinine, salicylates)

REFERENCE :

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

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IMMUNOHAEMATOLOGY

MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP

TYPE B

METHOD : SLIDE AGGLUTINATION

RH TYPE

POSITIVE

METHOD : SLIDE 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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CODE/NAME & ADDRESS : C000138383		ACCESSION NO : 0080WC007764	AGE/SEX : 34 Years Female
PROVISIONAL REPORT		PATIENT ID : JYOTF13088880	DRAWN :
		CLIENT PATIENT ID:	RECEIVED : 20/03/2023 08:46:05
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BIOCHEMISTRY

MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE

GLUCOSE FASTING, FLUORIDE PLASMA

FBS (FASTING BLOOD SUGAR) 88 74 - 106 mg/dL
METHOD : HEXOKINASE

GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD

HBA1C 5.5
 Non-diabetic Adult < 5.7 %
 Pre-diabetes 5.7 - 6.4
 Diabetes diagnosis: > or = 6.5
 Therapeutic goals: < 7.0
 Action suggested : > 8.0
 (ADA Guideline 2021)

ESTIMATED AVERAGE GLUCOSE(EAG) 111.2 < 116.0 mg/dL

GLUCOSE, POST-PRANDIAL, PLASMA

PPBS(POST PRANDIAL BLOOD SUGAR) 114 Non-Diabetes mg/dL
 70 - 140

LIPID PROFILE, SERUM

CHOLESTEROL, TOTAL 127 < 200 Desirable mg/dL
 200 - 239 Borderline High
 >/= 240 High

METHOD : CHOLESTEROL OXIDASE, ESTERASE, PEROXIDASE

TRIGLYCERIDES 73 < 150 Normal mg/dL
 150 - 199 Borderline High
 200 - 499 High
 >/= 500 Very High

METHOD : ENZYMATIC ASSAY

HDL CHOLESTEROL **32 Low** < 40 Low mg/dL
 >/=60 High

METHOD : DIRECT MEASURE - PEG



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SRL Ltd
 24 SCO, SECTOR 11 D
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 PUNJAB, INDIA
 Tel: 9111591115,
 CIN - U74899PB1995PLC045956

PATIENT NAME : JYOTI BANSAL

REF. DOCTOR : SELF

CODE/NAME & ADDRESS : C000138383

ACCESSION NO : 0080WC007764

AGE/SEX : 34 Years Female

PROVISIONAL REPORT

PATIENT ID : JYOTF13088880

DRAWN :

CLIENT PATIENT ID:

RECEIVED : 20/03/2023 08:46:05

ABHA NO :

REPORTED : 20/03/2023 16:19:33

Test Report Status	Final	Results	Biological Reference Interval	Units
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CHOLESTEROL LDL	80	< 100 Optimal 100 - 129 Near or above optimal 130 - 159 Borderline High 160 - 189 High >/= 190 Very High	mg/dL
-----------------	----	--	-------

METHOD : CHOLESTEROL OXIDASE, ESTERASE, PEROXIDASE

NON HDL CHOLESTEROL	95	Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL
---------------------	----	--	-------

METHOD : CALCULATED PARAMETER

VERY LOW DENSITY LIPOPROTEIN	14.6	Desirable value : 10 - 35	mg/dL
------------------------------	------	------------------------------	-------

METHOD : CALCULATED PARAMETER

CHOL/HDL RATIO	4.0	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	2.5	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	
---------------	-----	--	--

METHOD : CALCULATED PARAMETER

Interpretation(s)**LIVER FUNCTION PROFILE, SERUM**

BILIRUBIN, TOTAL	0.25	UPTO 1.2	mg/dL
------------------	------	----------	-------

METHOD : DIAZONIUM ION, BLANKED (ROCHE)

BILIRUBIN, DIRECT	0.10	0.00 - 0.30	mg/dL
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METHOD : DIAZOTIZATION

BILIRUBIN, INDIRECT	0.15	0.00 - 0.60	mg/dL
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METHOD : CALCULATED PARAMETER

TOTAL PROTEIN	7.3	6.6 - 8.7	g/dL
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METHOD : BIURET				
ALBUMIN		4.1	3.97 - 4.94	g/dL
METHOD : BROMOCRESOL GREEN				
GLOBULIN		3.2	2.0 - 4.0 Neonates - Pre Mature: 0.29 - 1.04	g/dL
METHOD : CALCULATED PARAMETER				
ALBUMIN/GLOBULIN RATIO		1.3	1.0 - 2.0	RATIO
METHOD : CALCULATED PARAMETER				
ASPARTATE AMINOTRANSFERASE (AST/SGOT)		15	0 - 32	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT)		10	0 - 31	U/L
METHOD : UV WITHOUT PYRIDOXAL-5 PHOSPHATE				
ALKALINE PHOSPHATASE		50	35 - 105	U/L
METHOD : PNPP - AMP BUFFER				
GAMMA GLUTAMYL TRANSFERASE (GGT)		11	5 - 36	U/L
METHOD : GAMMA GLUTAMYL CARBOXY 4-NITROANILIDE				
LACTATE DEHYDROGENASE		153	135 - 214	U/L
METHOD : LACTATE -PYRUVATE				
BLOOD UREA NITROGEN (BUN), SERUM				
BLOOD UREA NITROGEN		8	6 - 20	mg/dL
METHOD : UREASE - UV				
CREATININE, SERUM				
CREATININE		0.61	0.50 - 0.90	mg/dL
METHOD : ALKALINE PICRATE-KINETIC				
BUN/CREAT RATIO				
BUN/CREAT RATIO		13.11	5.00 - 15.00	
METHOD : CALCULATED PARAMETER				
URIC ACID, SERUM				
URIC ACID		4.0	2.4 - 5.7	mg/dL
METHOD : URICASE, COLORIMETRIC				
TOTAL PROTEIN, SERUM				
TOTAL PROTEIN		7.3	6.6 - 8.7	g/dL
METHOD : BIURET				
ALBUMIN, SERUM				

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ALBUMIN METHOD : BROMOCRESOL GREEN	4.1	3.97 - 4.94	g/dL
---------------------------------------	-----	-------------	------

GLOBULIN

GLOBULIN METHOD : CALCULATED PARAMETER	3.2	2.0 - 4.0 Neonates - Pre Mature: 0.29 - 1.04	g/dL
---	-----	---	------

ELECTROLYTES (NA/K/CL), SERUM

SODIUM, SERUM METHOD : ISE INDIRECT	136	136 - 145	mmol/L
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POTASSIUM, SERUM METHOD : ISE INDIRECT	4.55	3.5 - 5.1	mmol/L
---	------	-----------	--------

CHLORIDE, SERUM METHOD : ISE INDIRECT	104	98 - 107	mmol/L
--	-----	----------	--------

Interpretation(s)**Interpretation(s)****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 glycaemic 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 $eAG (mg/dl) = 28.7 * HbA1c - 46.7$

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

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anemia) will falsely lower HbA1c test results. Fructosamine is recommended in these patients which indicates diabetes control over 15 days.

(I) Vitamin C & E are reported to falsely lower test results, (possibly by inhibiting glycation of hemoglobin.

(II) 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.

(III) Interference of hemoglobinopathies in HbA1c estimation is seen in

- homozygous hemoglobinopathy. Fructosamine is recommended for testing of HbA1c.
- heterozygous state detected (D10 is corrected for HbS & HbC trait.)
- HbF > 25% on alternate platform (Soronate affinity chromatography) is recommended for testing of HbA1c. Abnormal Hemoglobin electrophoresis (HPLC method) is recommended for detecting a hemoglobinopathy

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

LIVER FUNCTION PROFILE, SERUM-LIVER FUNCTION PROFILE

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

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

ALP is a protein found in almost all body tissues. Tissues with higher amounts of ALP include the liver, bile ducts and bone. Elevated ALP levels are seen in Biliary obstruction, Osteoblastic bone tumors, osteomalacia, hepatitis, Hyperparathyroidism, Leukemia, Lymphoma, 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 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.

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 (Nephrolithiasis, Prostatism)

Causes of decreased level include Liver disease, SIADH.

CREATININE, SERUM-Higher than normal level may be due to:

- Blockage in the urinary tract
- Kidney problems, such as kidney damage or failure, infection, or reduced blood flow
- Loss of body fluid (dehydration)
- Muscle problems, such as breakdown of muscle fibers
- Problems during pregnancy, such as seizures (eclampsia), or high blood pressure caused by pregnancy (preeclampsia)

Lower than normal level may be due to:

- Myasthenia Gravis
- Muscular dystrophy

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, DCP, 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, 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.

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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Test Report Status **Final**

Results

Biological Reference Interval Units

CLINICAL PATH - URINALYSIS

MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE**PHYSICAL EXAMINATION, URINE**

COLOR PALE YELLOW
APPEARANCE SLIGHTLY HAZY

CHEMICAL EXAMINATION, URINE

PH	5.5	4.7 - 7.5	
METHOD : REFLECTANCE SPECTROPHOTOMETRY- DOUBLE INDICATOR METHOD			
SPECIFIC GRAVITY	1.025	1.003 - 1.035	
METHOD : REFLECTANCE SPECTROPHOTOMETRY (PKA CHANGE OF PRETREATED POLY ELECTROLYTES)			
PROTEIN	NOT DETECTED	NOT DETECTED	
METHOD : REFLECTANCE SPECTROPHOTOMETRY (PROTEIN-ERROR-OF-INDICATORS PRINCIPLE)			
GLUCOSE	NOT DETECTED	NOT DETECTED	
METHOD : REFLECTANCE SPECTROPHOTOMETRY (GLUCOSE OXIDASE/PEROXIDASE METHOD)			
KETONES	NOT DETECTED	NOT DETECTED	
METHOD : REFLECTANCE SPECTROPHOTOMETRY (SODIUM NITROPRUSSIDE REACTION)			
BLOOD	NOT DETECTED	NOT DETECTED	
METHOD : REFLECTANCE SPECTROPHOTOMETRY (PEROXIDASE METHOD)			
BILIRUBIN	NOT DETECTED	NOT DETECTED	
METHOD : REFLECTANCE SPECTROPHOTOMETRY (DIAZO REACTION)			
UROBILINOGEN	NORMAL	NORMAL	
METHOD : REFLECTANCE SPECTROPHOTOMETRY - EHRlich REACTION			
NITRITE	NOT DETECTED	NOT DETECTED	
METHOD : REFLECTANCE SPECTROPHOTOMETRY, CONVERSION OF NITRATE TO NITRITE			
LEUKOCYTE ESTERASE	NOT DETECTED	NOT DETECTED	
MICROSCOPIC EXAMINATION, URINE			
RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
METHOD : MICROSCOPIC EXAMINATION			
PUS CELL (WBC'S)	3-5	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION			
EPITHELIAL CELLS	3-5	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION			
CASTS	NOT DETECTED		
CRYSTALS	NOT DETECTED		

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METHOD : MICROSCOPIC EXAMINATION

BACTERIA

NOT DETECTED

NOT DETECTED

METHOD : MICROSCOPIC EXAMINATION

YEAST

NOT DETECTED

NOT DETECTED

Interpretation(s)



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CLINICAL PATH - STOOL ANALYSIS

MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE

PHYSICAL EXAMINATION,STOOL

COLOUR	BROWN		
CONSISTENCY	SEMI FORMED		
MUCUS	ABSENT	NOT DETECTED	
VISIBLE BLOOD	ABSENT	ABSENT	
ADULT PARASITE	NOT DETECTED		

METHOD : MICROSCOPIC EXAMINATION

CHEMICAL EXAMINATION,STOOL

STOOL PH 6.0

MICROSCOPIC EXAMINATION,STOOL

PUS CELLS	2-3		/hpf
RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
CYSTS	NOT DETECTED	NOT DETECTED	
OVA	NOT DETECTED		
LARVAE	NOT DETECTED	NOT DETECTED	
TROPHOZOITES	NOT DETECTED	NOT DETECTED	

Interpretation(s)

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