

CHANDIGARH
(A unit of Fortis Hospital Mohali)
SCO 11, Sector 11-D, Chandigarh - 160011

Name	My. Alyry	kumar		
UHID	. 12268634	Date :		12023
Age	: 33	Gandar	Ma	10

Nursing Assessment

, NI	ursing Assessment
	Profile
Height (cm): 1800 m	Waist Circumference (cm): 32 Tuch
Weight (Kg.): 80 Ky	Body Mass Index :
Occupation: Grand Joh	Marital Status Single Married
	Vital Signs
Pulse Rate (/min): 75 B/m / SPa	2 100/ Respiratory Rate (/min): 23 b/w
Blood Pressure (mmHg): 110/60 m	m H! Temperature (if febrile): A F ebruli
	Past History
Hypertension:	Diabetes :
🗵 Heart disease :	Dyslipidemia :
Asthma :	Tuberculosis :
X Allergies :	
	For Women
LMP:	Last Pap smear done in
Menopause 🗌 Yes 🔡 No	Last Mammography done in
Consent for X-ray & Mammography	
Curr	ent Medications
	-Α-
	<u> </u>
Detail 28 19 200000	

Signature, Name and Emp. ID of the Nurse :



CHANDIGARH (A unit of Fortis Hospital Mohali) SCO 11, Sector 11-D, Chandigarh - 160011

Name	My. A Ja	y Kymay
UHID	:12268839	Date: 1/2/2023
Age	: 33	Gender: Mall

	Internal Medicine Consultation	1
	1	
Relevant History:	Diagnosis;	
		*
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was the second of the second o		
Examination Findings:	Advice / Treatment Plan:	
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8	All Valor	
	-	
Investigations:		
Tie .		

Signature and stamp of the Consultant :

Fortis MEDCENTRE

CHANDIGARH
(A unit of Fortis Hospital Mohali)
SCO 11, Sector 11-D, Chandigarh - 160011

Name My. A Jay kymay

UHID: 12268639 Date: 1/2/2013

Age: 33, Gender: Maly

Ophthalmology Consultation

History: NIL	
Examination findings:	-1 W/.
Visual acuity Visual acuity with glasses (R)	Colour Vision R WNL
Slit Lamp Examination	
RE //	LE //
deal	Clear
Fundus Examination	
RE	LE
Diagnosis: NAD BE	
Treatment"	
Spectacle prescription:	
Right eye	Left eye
Distance 1 AVIS VA	SPH CYL AXIS VA
Near land N6	Distance Plane
iveal []	Near W.6
Signature and stamp of the Ophthalmologist :	V -

CHANDIGARH

8729000261

Fortis Medcentre

SCO-11, Sector-11-D, Chandigarh - 160 011 (India)

Telephone : 0172 506 1222 / 505 5441

Fax : 0172-5055440

E-mail : contactus.fmc@fortishealthcare.com

Website : www.fortishealthcare.com

1/2/23

Mr. Ajay kumen 33 | Male

0 1E Dental Canc. 18, 28, 17, 47, 37

Impacted 48, 38

Stains ++ Calculus +>

du. Oral Prophylavis

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6Pg.

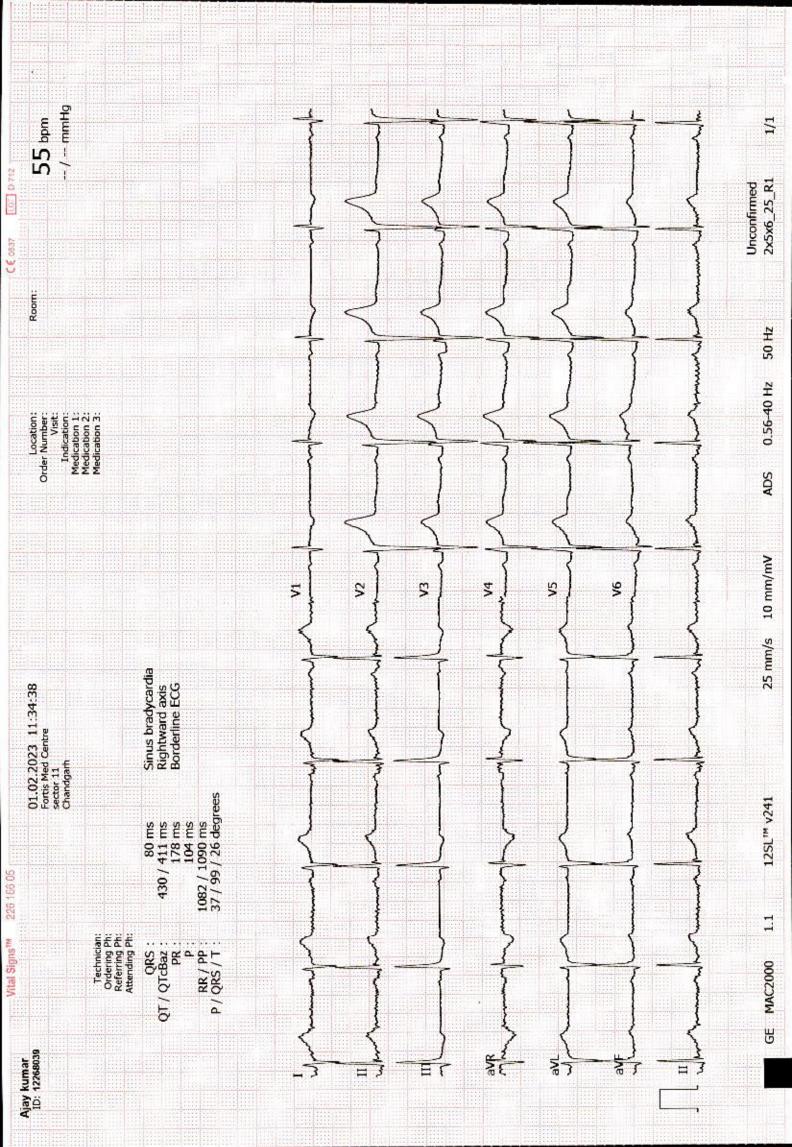
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Dr. Neetu B, Nerule
MD.BDS
Endodontist & Cosmetle Dehtist
Root Canal Specialist
Mobile:8729000281
Reg. No.2198-A
Forlic MEDCENTRE
S.C.C. 11, Sec. 11-D, Chandigerts
Ph. No.0172-5081222,5055141

For Dr newton.

A unit of FORTIS HOSPITAL MOHALI

Sector 62, Phase - VIII, Mohali - 160062, Punjab (India); Tel: +91 172 469 2222, 469 2250 Fax: +91 172 469 2221



4

Patient Name

: Ajay Kumar

UHID

: 12268039

Age / Gender

: 33 Year/ Male

Ward

Referred By

Diagnosis /

Clinical Information

Episode No.

: 0

Sample ID

: FHM23-R01530

Sample Drawn

Sample Received

: 01/Feb/2023 02:47 PM

Reported

: 01/Feb/2023 03:53 PM

Blood Group Report

Final Report

Sample Type

: EDTA

Method

: AUTOMATION

Forward Blood Group: B Rh Negative

Reverse Blood Group : B

Final Blood Group

: B Rh Negative

Remark

.

Tested By : hari om verma

Verified By : hari om verma

Approved By:

Dr. Apra Kalra Addi Director & Head Transfusion Medicine

Note: Blood group is identified by ABO antigens (forward grouping) present on red cell membrane And anti-ABO antibodies (reverse grouping) present in the plasma. A grouping discrepancy is when there is a mismatch in forward and reverse Blood grouping. Special methods need to be Performed to solve such discrepancies.

In case of Newborn/cord blood grouping, only forward blood grouping would be done as the anti-ABO antibodies (for reverse grouping) Are not present till 4 to 6 months of age. Thus new born grouping should be considered as provisional report and should be supplemented by re-blood grouping after 4 to 6 months of age/ or by more sensitive tests like molecular blood grouping.

"Blood grouping is done on the received sample. In case of any suspected discrepancy, Blood centre should be contacted,

*****End of Report ****

Reference:

Method section 2: Red cell typing; AABB technical manual 19th Ed Wong ECC, Punzalan RC. Neonatal and Pediatric Transfusion practice. Technical Manual, AABB, 19th Ed; p613-640



CHANDIGARH

NAME: MR. AJAY KUMAR

AGE AND SEX:33 Y/M UHID NO: 12268039 DATE: 01/02/2023

ROI: WHOLE ABDOMEN

Fortis Medcentre

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E-mail : contactus.fmc@fortishealthcare.com

Vebsite : www.fortishealthcare.com

Liver is normal in size, outline and 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 seen.

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 and is normal in outline and wall thickness. No calculi or growth seen

Prostate is normal in size, and shows normal outline and echopattern. No focal lesion seen.

No free fluid is seen.

Opinion: Normal study.

Suggested clinical correlation.

Dr. NEHA CHHABRA. Consultant Radiologist AJAY KUMAR 33/M

Accession #:

Study Date: 01/02/2023

Patient ID: 24391020230201

Alt ID:

DOB:

Age:

Gender:

Ht:

Wt:

BSA:

Institution: Fortis MEDCENTRE, Chandigarh

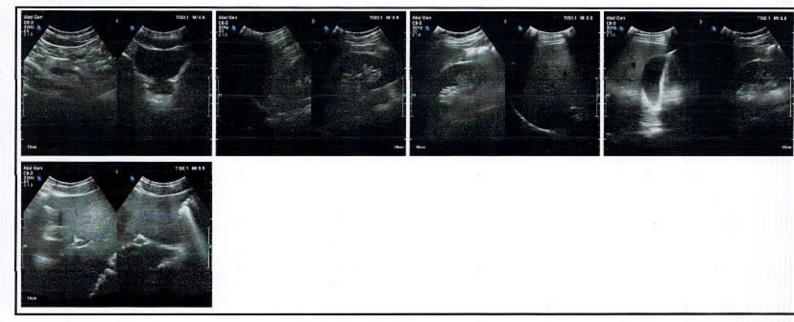
Referring Physician:

Physician of Record:

Performed By:

Comments:

Images



Signature

Signature:

Name(Print):

Date:



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Website : www.fortishealthcare.com

DEPARTMENT OF FMC-RADIOLOGY LAB

Date: 02/Fe1 / 11/12 »

" nels

Name: Mr. Ajay Kumar

Age | Sex: 33 YEAR(S) | Male

Order Station: FRONTOFFICE-FMC

Bed Name:

UHID | Episode No : 12268039 | 1122/23/10/21

Order No | Order Date: 10021/FN/OP/2302/2975 | 01-4 c - 2023

Admitted On | Reporting Date: 02-Feb-2 373 #9 34:44

Order Doctor Name : L . E F .

CHEST X-RAY (PA VIEW)

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.

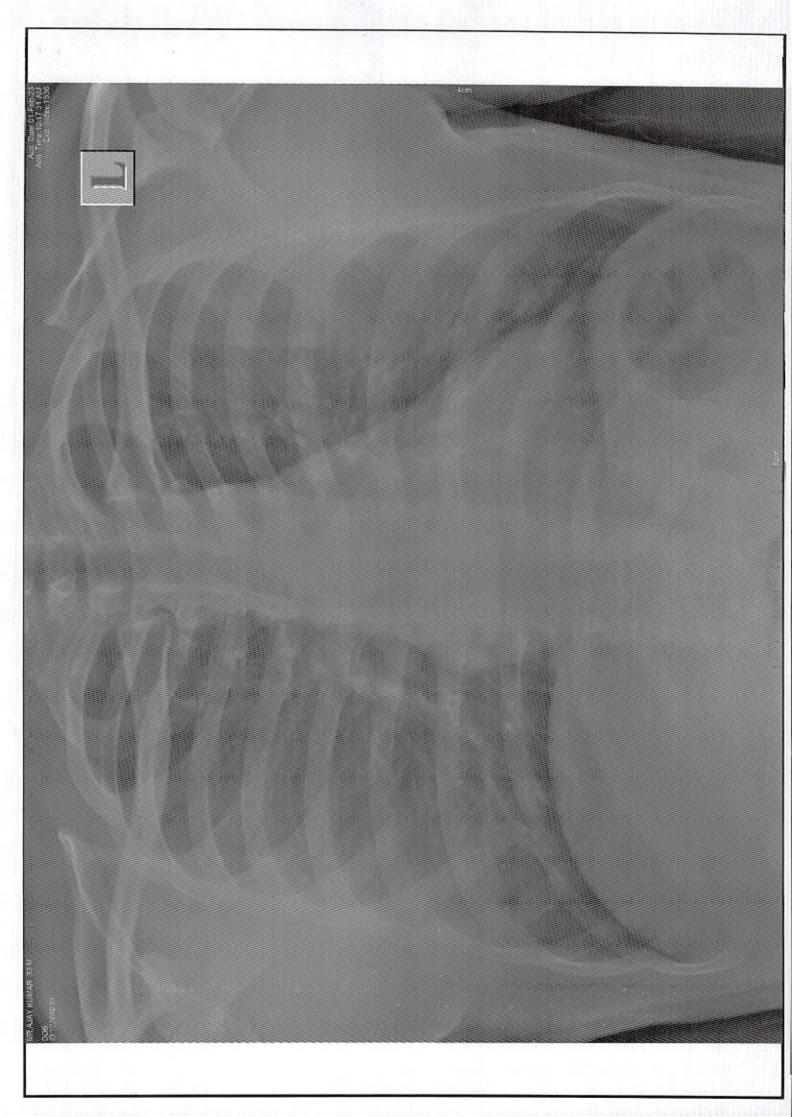
Bony cage and soft tissues are normal.

IMPRESSION: NORMAL STUDY.

Please correlate clinically and with other relevant investigations.

DR NEHA CHHABRA

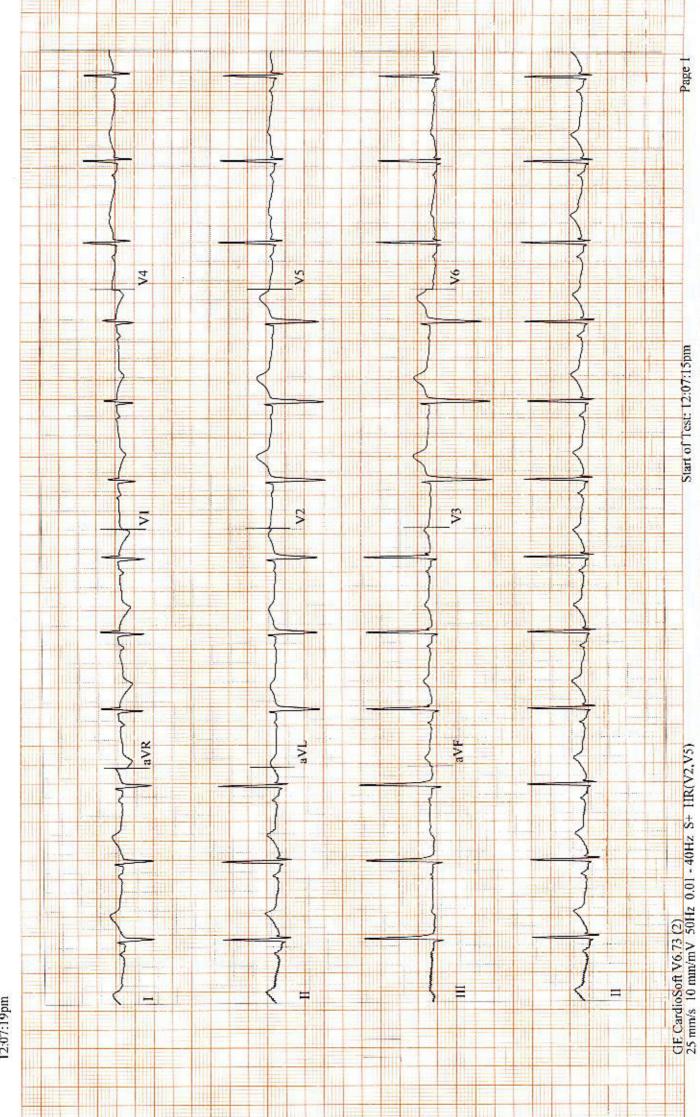
CONSULTANT RADIOLOGIST



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	STAGE 2 STAGE 3		03:00	5.50	14.00	109	130/80										
	STAGE 2 STAGE 3		03:00 02:24	5.50 6.80	14.00 16.00	109 131	130/80 140/80										
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he patient he resting naximal, agressure of unctional IR Respond Pressure In Respond Thest Pain: Arrhythmia	stage 2 stage 3 stage 4 exercised heart rate ge-predict 140/80 m on Resting Et Capacity: se to Exer se to Exer none. s: none. oression: 1	acco of 67 ed he mHg. CG: n norm reise:	o3:00 02:24 02:12 rding to the body rotal rate. The execution ormal repropriation or mail repropriation or ma	5.50 6.80 0.00 The BRU se to a m The resting testing Bl	14.00 16.00 8.90 CE for 1 naximal lang blood was sto	1:24 mir neart rate pressure pped due	130/80 140/80 110/80 h:s, achie e of 131 t e of 110/ e to Targe	pm. T 70 mm	his Hg	val , ro	ue re se to	pre:	sent naxi	s 7	0 %	ot	the
he patient he resting naximal, ag ressure of nterpretational IR Respond P Respond Chest Pain: Arrhythmia T Change Overall imp	stage 2 stage 3 stage 4 exercised heart rate ge-predict 140/80 m on Resting Et Capacity: se to Exer se to Exer none. s: none. oression: 1	acco of 67 ed he mHg. CG: n norm reise:	o3:00 02:24 02:12 rding to the body rotal rate. The execution ormal repropriation or mail repropriation or ma	5.50 6.80 0.00 The BRU se to a m The resting testing Bl	14.00 16.00 8.90 CE for 1 naximal lang blood was sto	1:24 mir neart rate pressure pped due	130/80 140/80 110/80 h:s, achie e of 131 t e of 110/ e to Targe	pm. T 70 mm	his Hg	val , ro	ue re se to	pre:	sent naxi	s 7	0 %	ot	the
he patient he resting naximal, ag ressure of nterpretatio ummary: I unctional IR Respon Chest Pain: Arrhythmia IT Change Overall imp	stage 2 stage 3 stage 4 exercised heart rate ge-predict 140/80 m on Resting Et Capacity: se to Exer se to Exer none. s: none. oression: 1	acco of 67 ed he mHg. CG: n norm reise:	o3:00 02:24 02:12 rding to the body rotal rate. The execution ormal repropriation or mail repropriation or ma	5.50 6.80 0.00 The BRU se to a m The resting testing Bl	14.00 16.00 8.90 CE for 1 naximal lang blood was sto	1:24 mir neart rate pressure pped due	130/80 140/80 110/80 h:s, achie e of 131 t e of 110/ e to Targe	pm. T 70 mm	his Hg	val , ro	ue re se to	pre:	sent naxi	s 7	0 %	ot	the
the patient the resting naximal, agressure of nterpretational IR Responding Presponding Change Overall implementation	stage 2 stage 3 stage 4 exercised heart rate ge-predict 140/80 m on Resting Et Capacity: se to Exer se to Exer none. s: none. oression: 1	acco of 67 ed he mHg. CG: n norm reise:	o3:00 02:24 02:12 rding to the body rotal rate. The execution ormal repropriation or mail repropriation or ma	5.50 6.80 0.00 The BRU se to a m The resting testing Bl	14.00 16.00 8.90 CE for 1 naximal lang blood was sto	1:24 mir neart rate pressure pped due	130/80 140/80 110/80 h:s, achie e of 131 t e of 110/ e to Targe	pm. T 70 mm	his Hg	val , ro	ue re se to	pre:	sent naxi	s 7	0 %	ot	the
the patient the resting naximal, agressure of nterpretational IR Responding Presponding Change Overall implementation	stage 2 stage 3 stage 4 exercised heart rate ge-predict 140/80 m on Resting Et Capacity: se to Exer se to Exer none. s: none. oression: 1	acco of 67 ed he mHg. CG: n norm reise:	o3:00 02:24 02:12 rding to the body rotal rate. The execution ormal repropriation or mail repropriation or ma	5.50 6.80 0.00 The BRU se to a m The resting testing Bl	14.00 16.00 8.90 CE for 1 naximal lang blood was sto	1:24 mir neart rate pressure pped due	130/80 140/80 110/80 h:s, achie e of 131 t e of 110/ e to Targe	pm. T 70 mm	his Hg	val , ro	ue re se to	pre:	sent naxi	s 7	0 %	ot	the
The patient The resting naximal, agressure of nterpretational IR Responsible Paint Arrhythmia T Change Overall implicational Improvement of the conclusion o	stage 2 stage 3 stage 4 exercised heart rate ge-predict 140/80 m on Resting Et Capacity: se to Exer se to Exer none. s: none. oression: 1	acco of 67 ed he mHg. CG: n norm reise:	o3:00 02:24 02:12 rding to the body rotal rate. The execution ormal repropriation or mail repropriation or ma	5.50 6.80 0.00 The BRU se to a m The resting testing Bl	14.00 16.00 8.90 CE for 1 naximal lang blood was sto	1:24 mir neart rate pressure pped due	130/80 140/80 110/80 h:s, achie e of 131 t e of 110/ e to Targe	pm. T 70 mm	his Hg	val , ro	ue re se to	pre:	sent naxi	s 7	0 %	ot	the
The patient The resting naximal, agressure of nterpretational IR Respondents Paint Arrhythmia T Change Overall implements on clusion	stage 2 stage 3 stage 4 exercised heart rate ge-predict 140/80 m on Resting Et Capacity: se to Exer se to Exer none. s: none. oression: 1	acco of 67 ed he mHg. CG: n norm reise:	o3:00 02:24 02:12 rding to the body rotal rate. The execution ormal repropriation or mail repropriation or ma	5.50 6.80 0.00 The BRU se to a m The resting testing Bl	14.00 16.00 8.90 CE for 1 naximal lang blood was sto	1:24 mir neart rate pressure pped due	130/80 140/80 110/80 h:s, achie e of 131 t e of 110/ e to Targe	pm. T 70 mm	his Hg	val , ro	ue re se to	pre:	sent naxi	s 7	0 %	ot	the
ECOVERY	stage 2 stage 3 stage 4 exercised heart rate ge-predict 140/80 m on Resting Et Capacity: se to Exer se to Exer none. s: none. oression: 1	acco of 67 ed he mHg. CG: n norm reise:	o3:00 02:24 02:12 rding to the body rotal rate. The execution ormal repropriation or mail repropriation or ma	5.50 6.80 0.00 The BRU se to a m The resting testing Bl	14.00 16.00 8.90 CE for 1 naximal lang blood was sto	1:24 mir neart rate pressure pped due	130/80 140/80 110/80 h:s, achie e of 131 t e of 110/ e to Targe	pm. T 70 mm	his Hg	val , ro	ue re se to	pre:	sent naxi	s 7	0 %	ot	the

AJAY, KUMAR Patient ID 12268039 01.02.2023 12:07:19pm

70 bpm



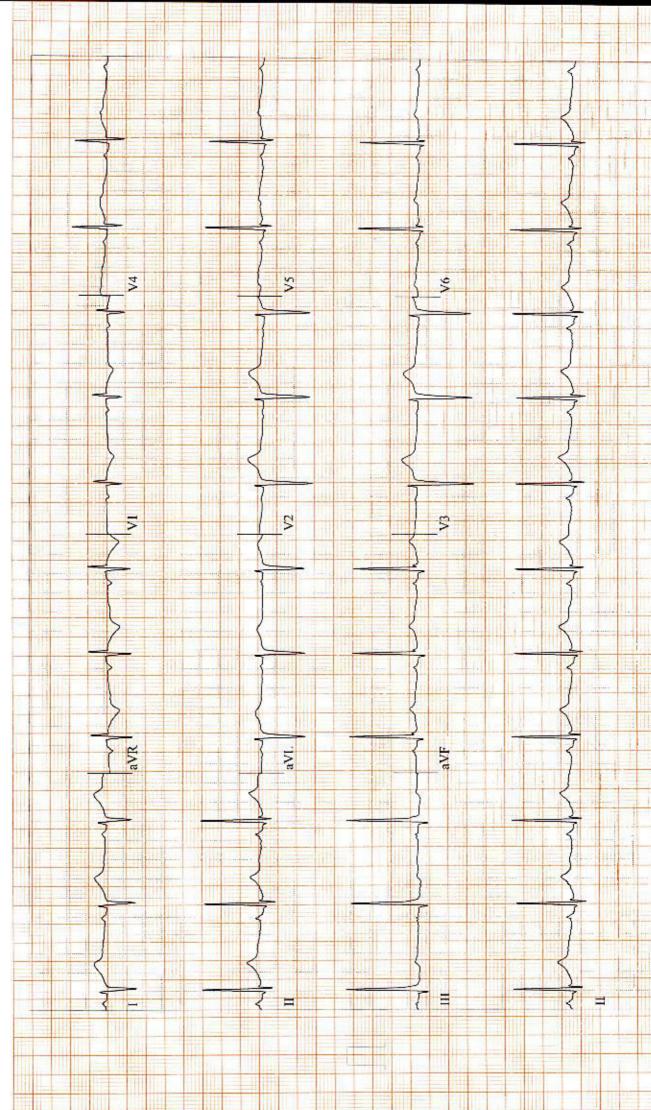
AJAY, KUMARPatient ID 12268039
01.02.2023
12:07:47pm

67 bpm 110/70 mmHg

PRETEST SUPINE 00:27

12-Lead Report

BRUCE 0.0 km/h 9.1 %



Page 2

Start of Test: 12:07:15pm

GE CardioSoft V6.73 (2) 25-mm/s 10 mm/mV 50Hz 0.01-40Hz S+ HR(V2,V5)

Раде 3 ST Level (mm) ST Slope (mV/s) Lead 94 Start of Test: 12:07:15pm V4-7-T SA 0.85 0.25 0.10 0.70 1.05 0.00 9/ BRUCE 2.7 km/h 10.0 % Ne_1 V3 1 1.05 0.75 0.45 0.25 0.53 1.10 -0.15 VS Comparative Medians Report 2 EXERCISE STAGE 1 02:50 GE CardioSoft V6.73 (2) 25 mm/s 10 mm/mV 50Hz 0.01 - 40Hz S+ HR(V2,V5) 84 bpm 110/70 mmHg Ser. CURRENT 60 ms post J 7 aVF aVR aVL -0.45 -1.49 -0.05 0.30 0.50 -0.35 0.65 Ħ BASELINE 60 ms post J くと avr." aVR. aVF. AJAY, KUMAR Patient ID 12268039 0.50 -0.55 0.20 0.40 -0.15 -0.42 0.60 E 01.02.2023 12:11:06pm

Page 4 Lead STLevel (mm) ST Slope (mV/s) 14 Start of Test: 12:07:15pm F 9/ NS Z V3 7 -0.39 -0.30 -0.30 0.10 0.30 1.00 0.80 0.15 74 BRUCE 4.0 km/h 12.0 % 1-- 1× 0.25 0.75 0.45 1.10 1.0 -0.15 0.53 9/ 75 5 EXERCISE STAGE 2 05:50 GE CardioSoft V6.73 (2) 25 mm/s 10 mm/mV 50Hz 0.01 - 40Hz S+ HR(II,V6) 94 bpm 120/80 mmHg SVR. 60 ms post J CURRENT aVF " aVE -0.15 -0.15 -1.38 -0.13 0.30 -1.02 0.00 -0.35 0.35 BASELINE 60 ms post J "VF AJAY, KUMAR Patient ID 12268039 01.02.2023 12:14:06pm 0.20 0.40 -0.01 0.15 -0.55 -1.13 avC 0.50 0.60

Comparative Medians Report

Page 5 ST Level (mm)
ST Slope (mV/s) Lead Start of Test: 12:07:15pm 1 - N 0.50 -0.65 0.80 0.57 0.80 0.05 9/ 0.35 -0.91 5.6 km/h 14.0 % BRUCE V3 -0.45 0.25 0.75 1.05 0.11 1.10 0.81 9/ -0.15 72 San Francis EXERCISE STAGE 3 08:50 MANA ANR JOSE GE CardioSoft V6.73 (2) 25 mm/s 10 mm/mV 50Hz 0.01 - 40Hz S+ HR(II.V6) 109 bpm 130/80 mmHg E 圣 60 ms post J ave. CURRENT aVE 0.35 -0.45 0.63 0.00 -0.55 0.20 0.30 1.12 Ξ = L tsod sm 09 BASELINE aVFaVR { aVE AJAY, KUMAR Patient ID 12268039 -0.55 0.40 0.20 -0.15 -0.42 0.50 0.36 0.60 12:17:06pm 01.02.2023

Comparative Medians Report

Page 6 ST Level (mm) ST Slope (mV/s) Lead Start of Test: 12:07:15pm V3-T -1.20 0.41 1.05 .28 -1.10 0.00 56.0 0.30 6.8 km/h 16.0 % 14-17 0.25 0.75 1.05 1.10 0.11 9/ STAGE 4 11:24 GE CardioSoft V6,73 (2) 25 mm/s 10 mm/mV 50Hz 0.01 - 40Hz S+ HR(V2,V6) 131 bpm 140/80 mmHg TO THE 60 ms post J CURRENT aVF -1.15 0.59 0.30 0,40 -1.95 0.80 -1.00 1.62 -1.30 60 ms post J BASELINE ave **₹** 0.40 0.20 aVL -0.55 -1.13 09.0 0.50 0.15 -0.42 12:19:40pm 01,02,2023

Comparative Medians Report (PEAK EXERCISE)

BRUCE

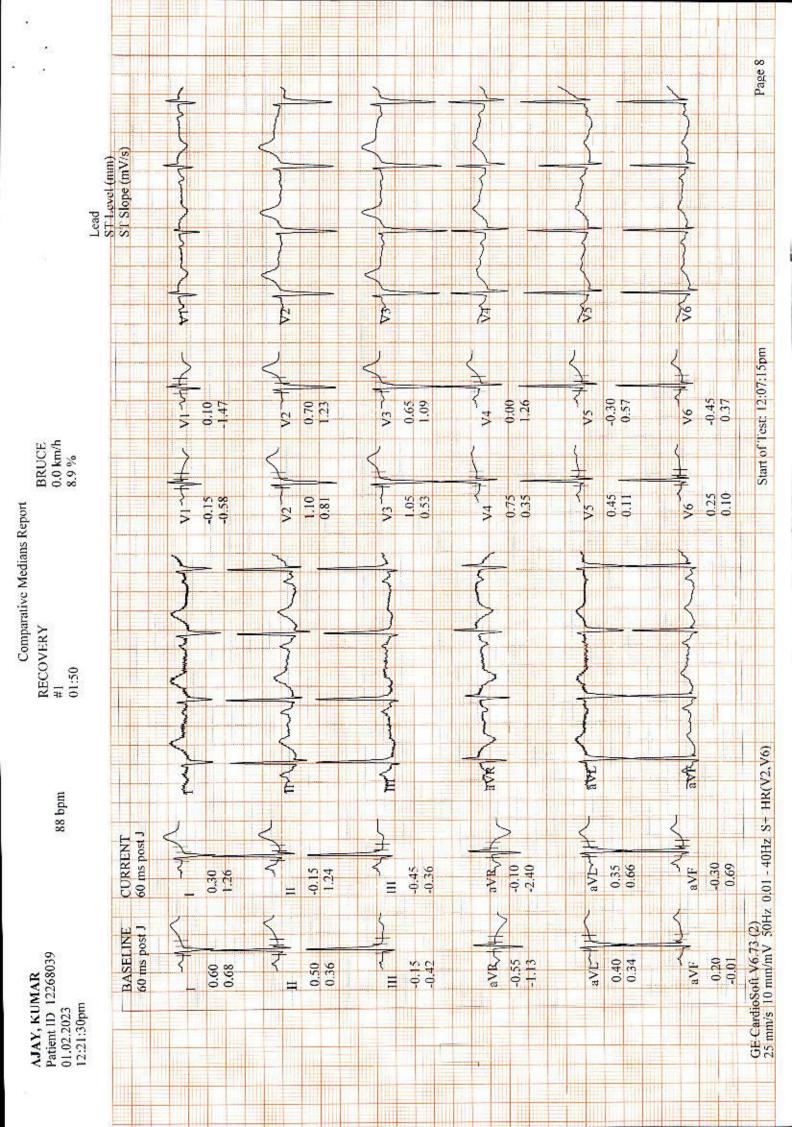
EXERCISE

Patient ID 12268039

AJAY, KUMAR

Page 7 ST Level (mm)
ST Slope (mV/s) Lead Start of Test: 12:07:15pm 0.50 20. 0.10 -0.20 1.54 1.05 -0.05 9/ 2.4 km/h 9.0 % BRUCE 李本 W674 V3---0.35 0.45 0.25 1.05 -0.15 0.81 0.11 RECOVERY 00:50 GE CardioSoft V6.73 (2) 25 mm/s 10 mm/mV 50Hz 0.01 - 40Hz S+ HR(V2.V6) 111 bpm 60 ms post J CURRENT aVITY -0.35 1.03 0.65 -0.30 0.70 2.25 0.60 BASELINE 60 ms post J 3 Patient ID 12268039 aVE 0.20 aVR_ -0.55 0.40 -0.15 -0.42 aVF 0.50 0.60 AJAY, KUMAR 01.02.2023 12:20:30pm

Comparative Medians Report







FORTIS MOHALI-CHC -SPLZD FORTIS HOSPITAL # MOHALI,

MOHALI 160062 7087030817 ACCESSION NO: **0006WB000563**PATIENT ID: FH.12268039

CLIENT PATIENT ID: UID:12268039

ABHA NO :

AGE/SEX :33 Years Male
DRAWN :01/02/2023 12:51:00
RECEIVED :01/02/2023 13:44:37
REPORTED :15/02/2023 14:06:31

CLINICAL INFORMATION:

UID:12268039 REQNO-1366087

CORP-OPD

BILLNO-10021230PCS001450 BILLNO-10021230PCS001450

Test Report Status	Final	Results	Biological Reference Interval	Unite
rest keport Status	<u>Finai</u>	Results	biological Reference Interval	Units

Н.	AEMATOLOGY - CBC		
CBC-5, EDTA WHOLE BLOOD			
BLOOD COUNTS, EDTA WHOLE BLOOD			
HEMOGLOBIN (HB) METHOD: SLS- HEMOGLOBIN DETECTION METHOD	15.4	13.0 - 17.0	g/dL
RED BLOOD CELL (RBC) COUNT METHOD: HYDRODYNAMIC FOCUSING	4.86	4.5 - 5.5	mil/μL
WHITE BLOOD CELL (WBC) COUNT METHOD: FLOWCYTOMETRY	6.90	4.0 - 10.0	thou/µL
PLATELET COUNT METHOD: HYDRO DYNAMIC FOCUSING METHOD / MICROSCOPY	130 Low	150 - 410	thou/µL
Comments			
MANY PLATELETS CLUMPS ARE SEEN. RBC AND PLATELET INDICES			
HEMATOCRIT (PCV) METHOD: HYDRODYNAMIC FOCUSING	45.3	40.0 - 50.0	%
MEAN CORPUSCULAR VOLUME (MCV) METHOD: CALCULATED PARAMETER	93.2	83.0 - 101.0	fL
MEAN CORPUSCULAR HEMOGLOBIN (MCH) METHOD: CALCULATED PARAMETER	31.7	27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION(MCHC) METHOD: CALCULATED PARAMETER	34.0	31.5 - 34.5	g/dL
RED CELL DISTRIBUTION WIDTH (RDW) METHOD: CALCULATED PARAMETER	13.6	11.6 - 14.0	%
MENTZER INDEX METHOD: CALCULATED PARAMETER	19.2		
MEAN PLATELET VOLUME (MPV) METHOD: CALCULATED PARAMETER	12.5 High	6.8 - 10.9	fL

WBC DIFFERENTIAL COUNT

Ritu Pantoy

Meenahahi Malhotra

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PUNJAB, INDIA

Tel: 0172-469-2222 Extn. 6726, 6727), 0172-469-2221 - CIN - L85110DL1996PLC076704







FORTIS MOHALI-CHC -SPLZD FORTIS HOSPITAL # MOHALI,

MOHALI 160062 7087030817 ACCESSION NO: **0006WB000563**PATIENT ID: FH.12268039

CLIENT PATIENT ID: UID:12268039

ABHA NO :

AGE/SEX :33 Years Male
DRAWN :01/02/2023 12:51:00
RECEIVED :01/02/2023 13:44:37
REPORTED :15/02/2023 14:06:31

CLINICAL INFORMATION:

UID:12268039 REQNO-1366087

CORP-OPD

BILLNO-10021230PCS001450 BILLNO-10021230PCS001450

DILLINO-1002123OFC3	3001430			
Test Report Status	<u>Final</u>	Results	Biological Reference	Interval Units
NEUTROPHILS		52	40.0 - 80.0	%
METHOD : FLOW CYTOMET	RY+LEISHMAIN STAIN+MICROSCOPY			
LYMPHOCYTES		37	20.0 - 40.0	%
METHOD : FLOW CYTOMET	RY+LEISHMAIN STAIN+MICROSCOPY			
MONOCYTES		7	2.0 - 10.0	%
METHOD : FLOW CYTOMET	RY+LEISHMAIN STAIN+MICROSCOPY			
EOSINOPHILS		4	1 - 6	%
METHOD : FLOW CYTOMET	RY+LEISHMAIN STAIN+MICROSCOPY			
BASOPHILS		00	0 - 2	%
METHOD : FLOW CYTOMET	RY+LEISHMAIN STAIN+MICROSCOPY			
ABSOLUTE NEUTRO	PHIL COUNT	3.73	2.0 - 7.0	thou/µL
METHOD : CALCULATED PA	RAMETER			
ABSOLUTE LYMPHO	CYTE COUNT	2.55	1.0 - 3.0	thou/µL
METHOD : CALCULATED PA	RAMETER			
ABSOLUTE MONOCY	TE COUNT	0.48	0.2 - 1.0	thou/µL
METHOD : CALCULATED PA	RAMETER			
ABSOLUTE EOSINO	PHIL COUNT	0.28	0.02 - 0.50	thou/µL
METHOD : CALCULATED PA	RAMETER			
NEUTROPHIL LYMPH	HOCYTE RATIO (NLR)	1.4		
METHOD : CALCULATED PA	` '			

Interpretation(s)

RBC AND PLATELET INDICES-Mentzer index (MCV/RBC) is an automated cell-counter based calculated screen tool to differentiate cases of Iron deficiency anaemia(>13) from Beta thalassaemia trait

(<13) in patients with microcytic anaemia. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HbA2 remains the gold standard for diagnosing a case of beta thalassaemia trait.

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

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



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MOHALI, 160062 PUNJAB, INDIA

Tel: 0172-469-2222 Extn. 6726, 6727), 0172-469-2221 - CIN - L85110DL1996PLC076704







REF. DOCTOR: SELF PATIENT NAME: AJAY KUMAR

FORTIS MOHALI-CHC -SPLZD FORTIS HOSPITAL # MOHALI,

MOHALI 160062 7087030817

ACCESSION NO: 0006WB000563 PATIENT ID : FH.12268039 CLIENT PATIENT ID: UID:12268039

ABHA NO

AGE/SEX :33 Years :01/02/2023 12:51:00 DRAWN RECEIVED: 01/02/2023 13:44:37 REPORTED :15/02/2023 14:06:31

CLINICAL INFORMATION:

UID:12268039 REQNO-1366087

CORP-OPD

BILLNO-10021230PCS001450 BILLNO-10021230PCS001450

Test Report Status Results **Biological Reference Interval** Units <u>Final</u>

HAEMATOLOGY

ERYTHROCYTE SEDIMENTATION RATE (ESR), WHOLE BLOOD

0 - 14mm at 1 hr E.S.R

METHOD: WESTERGREN METHOD

Interpretation(s)

ERYTHROCYTE SEDIMENTATION RATE (ESR), WHOLE BLOOD-TEST DESCRIPTION :-

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

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

Increase in: Infections, Vasculities, 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: Polycythermia vera, Sickle cell anemia

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)

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.



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FORTIS MOHALI-CHC -SPLZD FORTIS HOSPITAL # MOHALI,

MOHALI 160062 7087030817

ACCESSION NO: 0006WB000563 PATIENT ID : FH.12268039 CLIENT PATIENT ID: UID:12268039

ABHA NO

AGE/SEX :33 Years DRAWN :01/02/2023 12:51:00 RECEIVED: 01/02/2023 13:44:37 REPORTED :15/02/2023 14:06:31

CLINICAL INFORMATION:

UID:12268039 REQNO-1366087

CORP-OPD

BILLNO-10021230PCS001450 BILLNO-10021230PCS001450

Test Report Status	Final	Results	Biological Reference Interval	Units

	BIOCHEMISTRY		
LIVER FUNCTION PROFILE, SERUM			
BILIRUBIN, TOTAL METHOD: DIAZONIUM ION, BLANKED (ROCHE)	0.39	UPTO 1.2	mg/dL
BILIRUBIN, DIRECT METHOD: DIAZOTIZATION	0.13	0.00 - 0.30	mg/dL
BILIRUBIN, INDIRECT METHOD: CALCULATED PARAMETER	0.26	0.00 - 0.60	mg/dL
TOTAL PROTEIN METHOD: BIURET	7.6	6.6 - 8.7	g/dL
ALBUMIN METHOD: BROMOCRESOL GREEN	4.7	3.97 - 4.94	g/dL
GLOBULIN	2.9	2.0 - 4.0 Neonates - Pre Mature: 0.29 - 1.04	g/dL
METHOD: CALCULATED PARAMETER			
ALBUMIN/GLOBULIN RATIO METHOD: CALCULATED PARAMETER	1.6	1.0 - 2.0	RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	28	0 - 40	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT) METHOD: UV WITHOUT PYRIDOXAL-5 PHOSPHATE	35	0 - 41	U/L
ALKALINE PHOSPHATASE METHOD: PNPP - AMP BUFFER	110	40 - 129	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT) METHOD: GAMMA GLUTAMYLCARBOXY 4NITROANILIDE	64 High	8 - 61	U/L
LACTATE DEHYDROGENASE METHOD: LACTATE -PYRUVATE UV	150	135 - 225	U/L
GLUCOSE FASTING, FLUORIDE PLASMA			
FBS (FASTING BLOOD SUGAR) METHOD: HEXOKINASE	101	74 - 106	mg/dL

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Ms. Hardeep Kaur, M.Sc. **Biochemistry**

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Tel: 0172-469-2222 Extn. 6726, 6727), 0172-469-2221 - CIN - L85110DL1996PLC076704 Email: srl.mohali@fortishealthcare.com

Patient Ref. No. 6000002937607





FORTIS MOHALI-CHC -SPLZD FORTIS HOSPITAL # MOHALI,

MOHALI 160062 7087030817

ACCESSION NO: 0006WB000563 PATIENT ID : FH.12268039 CLIENT PATIENT ID: UID:12268039

ABHA NO

AGE/SEX :33 Years DRAWN :01/02/2023 12:51:00 RECEIVED: 01/02/2023 13:44:37 REPORTED :15/02/2023 14:06:31

CLINICAL INFORMATION:

UID:12268039 REQNO-1366087

CORP-OPD

BILLNO-10021230PCS001450 BILLNO-10021230PCS001450

Test Report Status <u>Final</u>	Results	Biological Reference Interv	al Units
BLOOD UREA NITROGEN (BUN), SERUM			
BLOOD UREA NITROGEN METHOD: UREASE - UV	12	6 - 20	mg/dL
URIC ACID, SERUM			
URIC ACID METHOD: URICASE, COLORIMETRIC	8.1 High	3.4 - 7.0	mg/dL
GLYCOSYLATED HEMOGLOBIN(HBA1C), EDT	A WHOLE BLOOD		
HBA1C	5.5	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 Therapeutic goals: < 7.0 Action suggested: > 8.0 (ADA Guideline 2021)	%
METHOD: HPLC		,	
ESTIMATED AVERAGE GLUCOSE(EAG) METHOD: CALCULATED PARAMETER CREATININE EGFR	111.2	< 116.0	mg/dL
CREATININE METHOD: ALKALINE PICRATE-KINETIC	0.80	0.70 - 1.20	mg/dL
AGE	33		years
GLOMERULAR FILTRATION RATE (MALE)	111	GFR of +90 normal or minimal kidney damage with normal GFR 89- 60 mild decrease 59-30 moderate decrease 29-15 severe decrease < 15 kidney failure (units: mL/min/1.73mSq.)	

GLUCOSE POST-PRANDIAL, PLASMA

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70 - 140



REF. DOCTOR: SELF PATIENT NAME: AJAY KUMAR

FORTIS MOHALI-CHC -SPLZD FORTIS HOSPITAL # MOHALI,

MOHALT 160062 7087030817

ACCESSION NO: 0006WB000563 PATIENT ID : FH.12268039 CLIENT PATIENT ID: UID:12268039

ABHA NO

AGE/SEX :33 Years Male :01/02/2023 12:51:00 DRAWN RECEIVED: 01/02/2023 13:44:37 REPORTED :15/02/2023 14:06:31

CLINICAL INFORMATION:

UID:12268039 REQNO-1366087

CORP-OPD

BILLNO-10021230PCS001450 BILLNO-10021230PCS001450

Test Report Status <u>Fir</u>	nal	Results	Biological Reference I	nterval Units
PPBS(POST PRANDIAL B	LOOD SUGAR)	106	Non-Diabetes	mg/dL

METHOD: HEXOKINASE

Interpretation(s)

LIVER FUNCTION PROFILE, SERUM-

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

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

hepatitis, obstruction of bile ducts, cirrhosis. **ALP** is a protein found in almost all body tissues. Tissues with higher amounts of ALP include the liver, bile ducts and bone. Elevated ALP levels are seen in Biliary obstruction, Osteoblastic bone tumors, osteomalacia, hepatitis, Hyperparathyroidism, Leukemia, Lymphoma, Pagets disease, Rickets, Sarcoidosis etc. Lower-than-normal ALP levels seen in Hypophosphatasia, Malnutrition, Protein deficiency, Wilsons disease.

GGT is an enzyme found in cell membranes of many tissues mainly in the liver, kidney and pancreas. It is also found in other tissues including intestine, spleen, heart, brain

and seminal vesicles. The highest concentration is in the kidney, but the liver is considered the source of normal enzyme activity. Serum GGT has been widely used as an index of liver dysfunction. Elevated serum GGT activity can be found in diseases of the liver, billiary system and pancreas. Conditions that increase serum GGT are obstructive liver disease, high alcohol consumption and use of enzyme-inducing drugs etc.

Total Protein also known as total protein, is a biochemical test for measuring the total amount of protein in serum. Protein in the plasma is made up of albumin and

globulin.Higher-than-normal levels may be due to:Chronic inflammation or infection,including HIV and hepatitis B or C,Multiple myeloma,Waldenstroms disease.Lower-than-normal levels may be due to: Agammaglobulinemia,Bleeding (hemorrhage),Burns,Glomerulonephritis,Liver disease, Malabsorption,Malnutrition,Nephrotic

syndrome, Protein-losing enteropathy etc. **Albumin** is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by:Liver disease like cirrhosis of the liver, nephrotic syndrome,protein-losing enteropathy,Burns,hemodilution,increased vascular permeability or decreased lymphatic clearance,malnutrition and wasting etc GLUCOSE FASTING,FLUORIDE PLASMA-**TEST DESCRIPTION**

Normally, the glucose concentration in extracellular fluid is closely regulated so that a source of energy is readily available to tissues and sothat no glucose is excreted in the

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 Hypoglycaemics & Insulin treatment,Renal Glyosuria,Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc.

BLOOD UREA NITROGEN (BUN), SERUM-Causes of Increased levels include Pre renal (High protein diet, Increased protein catabolism, GI haemorrhage, Cortisol,

Dehydration, CHF Renal), Renal Failure, Post Renal (Malignancy, Nephrolithiasis, Prostatism) Causes of decreased level include Liver disease, SIADH.

URIC ACID, SERUM-Causes of Increased levels:-Dietary(High Protein Intake,Prolonged Fasting,Rapid weight loss),Gout,Lesch nyhan syndrome,Type 2 DM,Metabolic

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View Report



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FORTIS MOHALI-CHC -SPLZD FORTIS HOSPITAL # MOHALI,

MOHALT 160062 7087030817

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AGE/SEX :33 Years :01/02/2023 12:51:00 DRAWN RECEIVED: 01/02/2023 13:44:37 REPORTED :15/02/2023 14:06:31

CLINICAL INFORMATION:

UID:12268039 REQNO-1366087

CORP-OPD

BILLNO-10021230PCS001450 BILLNO-10021230PCS001450

Test Report Status Results **Biological Reference Interval** Units <u>Final</u>

syndrome Causes of decreased levels-Low Zinc intake,OCP,Multiple Sclerosis GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD-Used For:

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

The ADA recommends measurement of HbA1c (typically 3-4 times per year for type 1 and poorly controlled type 2 diabetic patients, and 2 times per year for well-controlled type 2 diabetic patients) to determine whether a patients metabolic control has remained continuously within the target range.

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

HbA1c Estimation can get affected due to :

- 1. Shortened Erythrocyte survival: Any condition that shortens erythrocyte survival or decreases mean erythrocyte age (e.g. recovery from acute blood loss,hemolytic anemia) will falsely lower HbA1c test results.Fructosamine is recommended in these patients which indicates diabetes control over 15 days.
- 2.Vitamin C & E are reported to falsely lower test results.(possibly by inhibiting glycation of hemoglobin.
 3. Iron deficiency anemia is reported to increase test results. Hypertriglyceridemia, uremia, hyperbilirubinemia, chronic alcoholism, chronic ingestion of salicylates & opiates addiction are reported to interfere with some assay methods, falsely increasing results.

 4. Interference of hemoglobinopathies in HbA1c estimation is seen in
- a) Homozygous hemoglobinopathy. Fructosamine is recommended for testing of HbA1c. b) Heterozygous state detected (D10 is corrected for HbS & HbC trait.)
- c) HbF > 25% on alternate paltform (Boronate affinity chromatography) is recommended for testing of HbA1c.Abnormal Hemoglobin electrophoresis (HPLC method) is

recommended for detecting a hemoglobinopathy

CREATININE EGFR-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

This equation takes into account several factors that impact creatinine production, including age, gender, and race. In children, eGFR is calculated using original schwartz equation.

The equation has not been validated in children & will only be reported for patients > 16 years of age. The equation is normalized for an average adult body surface area of 1.73m², weight & height adjustment is not necessary.

The IDMS Traceable MDRD equation has not been validated in children & will only be reported for patients = 18 years of age. The equation is normalized for an average adult body surface area of 1.73m², weight & height adjustment is not necessary. Estimation of GFR in children and adolescence (0- < 18 years) is performed by bedside IDMS- Traceable Schwartz formula

GLUCOSE POST-PRANDIAL, PLASMA-Spectrophotometry Hexokinase

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MOHALI 160062 7087030817

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CLINICAL INFORMATION:

UID:12268039 REQNO-1366087

CORP-OPD

BILLNO-10021230PCS001450 BILLNO-10021230PCS001450

Test Report Status Results Biological Reference Interval Units <u>Final</u>

BIOCHEMISTRY - LIPID						
LIPID PROFILE, SERUM						
CHOLESTEROL, TOTAL	201 High	< 200 Desirable 200 - 239 Borderline High >/= 240 High	mg/dL			
METHOD: CHOLESTEROL OXIDASE, ESTERASE, PEROXIDASE						
TRIGLYCERIDES	107	< 150 Normal 150 - 199 Borderline High 200 - 499 High >/= 500 Very High	mg/dL			
METHOD: ENZYMATIC ASSAY		, -				
HDL CHOLESTEROL METHOD: DIRECT MEASURE - PEG	43	< 40 Low >/=60 High	mg/dL			
	144 High	. 100 Oakaral				
LDL CHOLESTEROL, DIRECT	144 High	< 100 Optimal 100 - 129 Near or above optimal 130 - 160 Borderline High 161 - 189 High >/= 190 Very High	mg/dL			
METHOD: CHOLESTEROL OXIDASE, ESTERASE, PEROXIDASE						
NON HDL CHOLESTEROL	158 High	Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL			
VERY LOW DENSITY LIPOPROTEIN	21.4	Desirable value : 10 - 35	mg/dL			
METHOD: CALCULATED PARAMETER						
CHOL/HDL RATIO	4.7 High	3.3-4.4 Low Risk 4.5-7.0 Average Risk 7.1-11.0 Moderate Risk > 11.0 High Risk				

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MOHALI 160062 7087030817 ACCESSION NO: **0006WB000563**PATIENT ID: FH.12268039

CLIENT PATIENT ID: UID:12268039

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CORP-OPD

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Test Report Status <u>Final</u> Results Biological Reference Interval Units

LDL/HDL RATIO

3.4 High

0.5 - 3.0 Desirable/Low Risk

3.1 - 6.0 Borderline/Moderate

Risk

>6.0 High Risk

METHOD: CALCULATED PARAMETER

Interpretation(s)

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Meenahahi Malhotra

Dr. Meenakshi Malhotra, MD Senior Consultant,48159





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CORP-OPD

BILLNO-10021230PCS001450 BILLNO-10021230PCS001450

Results **Test Report Status Biological Reference Interval** <u>Final</u>

CLINICAL PATH - URINALYSIS

URINALYSIS

PHYSICAL EXAMINATION, URINE

YELLOW COLOR

METHOD: MANUAL EXAMINATION

APPEARANCE CLEAR

METHOD: MANUAL EXAMINATION

CHEMICAL EXAMINATION, URINE

4.7 - 7.5 6.0

METHOD: DOUBLE INDICATOR PRINCIPLE

SPECIFIC GRAVITY 1.010 1.003 - 1.035

METHOD: REFLECTANCE PHOTOMETRY (IONIC CONCENTRATION)

NOT DETECTED NOT DETECTED **PROTFIN**

METHOD: REFLECTION PHOTOMETRY (PROTEIN ERROR INDICATOR)

NOT DETECTED NOT DETECTED GLUCOSE

METHOD: REFLECTANCE PHOTOMETRY (GLUCOSE OXIDASE METHOD)

NOT DETECTED KETONES NOT DETECTED

METHOD: REFLECTION PHOTOMETRY (NITROPRUSSIDE)

NOT DETECTED NOT DETECTED

METHOD: REFLECTANCE PHOTOMETRY (BENZIDINE REACTION)

BILIRUBIN NOT DETECTED NOT DETECTED

METHOD: REFLECTANCE SPECTROPHOTOMETRY (DIAZO REACTION)

NORMAL **NORMAL**

METHOD: REFLECTANCE PHOTOMETRY (EHRLICH'S REACTION)

NOT DETECTED NOT DETECTED

METHOD: REFLECTANCE SPECTROPHOTOMETRY (DIAZO REACTION)

MICROSCOPIC EXAMINATION, URINE

NOT DETECTED /HPF RED BLOOD CELLS NOT DETECTED

METHOD: MICROSCOPY

PUS CELL (WBC'S) /HPF NOT DETECTED 0-5

METHOD: REFLECTANCE PHOTOMETRY & MICROSCOPY

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CORP-OPD

BILLNO-10021230PCS001450 BILLNO-10021230PCS001450

BILLNO-1002123OPCS001450					
Test Report Status	<u>Final</u>	Results	Biological Reference Interval Units		
EPITHELIAL CELLS METHOD: MICROSCOPY		NOT DETECTED	0-5	/HPF	
CASTS METHOD: MICROSCOPY		NOT DETECTED			
CRYSTALS METHOD: MICROSCOPY		NOT DETECTED			
BACTERIA METHOD: MICROSCOPY		NOT DETECTED	NOT DETECTED		
YEAST		NOT DETECTED	NOT DETECTED		
Interpretation(s)					



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CLINICAL INFORMATION:

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CORP-OPD

BILLNO-10021230PCS001450 BILLNO-10021230PCS001450

SPECIALISED CHEMISTRY - HORMONE						
THYROID PANEL, SERUM						
T3 METHOD: SANDWICH (ECLIA)	144.0	80.00 - 200.00	ng/dL			
T4 METHOD: SANDWICH (ECLIA)	8.15	5.10 - 14.10	μg/dL			
TSH (ULTRASENSITIVE) METHOD: SANDWICH (ECLIA)	1.060	0.270 - 4.200	μIU/mL			
Interpretation(s)						

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

Meenahah: Malhotra

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