TOTTIS MEDCENTRE NAM BABO SINICH4 . 23-3-24. Name CHANDIGARH Date : ____ UHID : (A unit of Fortis Hospital Mohali) SCO 11, Sector 11-D, Chandigarh - 160011 ma Gender : __ Age

Nursing Assessment

P	rofile
Height (cm): 65CM	Waist Circumference (cm): 36 thehee
Weight (Kg.): 65-2KC1 '	Body Mass Index :
Occupation: Reiniate Job	Marital Status Single Married
Vita	al Signs
Pulse Rate (min): 883/mint Stord-98	Respiratory Rate (/min): 20.5/min-1
Blood Pressure (mmHg): 160/Sommy	Temperature (if febrile) : A Joboulle
Past	History
Hypertension :	Diabetes :
Heart disease :	Dyslipidemia :
Asthma :	Tuberculosis :
Allergies :	
/ For V	Nomen
LMP:	Last Pap smear done in
Menopause 🗌 Yes 🖉 No	Last Mammography done in
Consent for X-ray & Mammography	
Current M	Aedications
	*

Signature, Name and Emp. ID of the Nurse : 1009

St Fortis MEDCENTRE CHANDIGARH

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

MR. SHMAM BABU Sihicity. 13049279 Date: 23-3-24. 54 Year Gender: Malc. Name UHID Age

Internal Medicine Consultation

Relevant History:

Diagnosis:

Examination Findings:

Advice / Treatment Plan:

Investigations:

Signature and stamp of the Consultant

m, Singh 13049279	Maie	23 23 23 23 23 23 23 23 23 23 23 23 23 2	23.03.2024 10:42:35 Fortis Med Centre sector 11 Chandgarh	Location: Order Number: Vist: Indication: Medication:	93 bpm / mmHg
	Technician: Ordering Ph: Referring Ph: Attending Ph:			Medication 2: Medication 3:	
	QT / QTBaz : PR P P P P P P P P P P P P P P P P P P	68 ms 348 / 432 ms 168 ms 110 ms 648 / 645 ms 50 / -12 / 72 degrees	Normal sinus rhythm Minimal voltage criteria for LVH, may be normal variant Nonspecific T wave abnormality Abnormal ECG		
1	Junt	Junt			
- ^{MR}					22
- Ne					1
*					
1		-			Internet
R	GE MAC2000 1.1	l 125L ^m v241	25 mm/s 10 mm/mV	ADS 0.56-40 Hz 50 Hz 20	Unconfirmed 2x5x6_25_R1 1/1

CHANDIGARH	Name Min Shoam Baby Singh.
(A unit of Fortis Hospital Mohali)	UHID: 130492 Age Date: 23-3-24
SCO 11, Sector 11-D, Chandigarh - 160011	Age: 54, Gender: Male.
	Age :

Ophthalmology Consultation

_R 619

666

Visual acuity with glasses

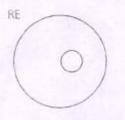
History: NIL

Examination findings: Visual acuity

Slit Lamp Examinat RE Clear

LE clear

Fundus Examination

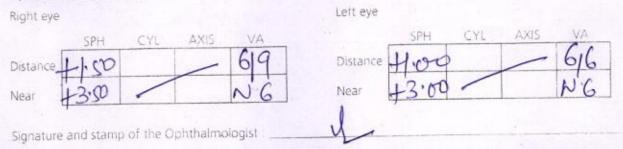


Colour Vision

Diagnosis: Pusbyolia BE

Treatment"

Spectacle prescription:





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

NAME: MR. SHYAM BABU SINGH AGE AND SEX: 54Y/M UHID NO: 13049279 DATE:23/03/2024 ROI: WHOLE ABDOMEN

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

Gall bladder is normally distended. Wall thickness is normal. Multiple echogenic foci of size 6-7 mm are seen along both the walls as well as in the lumen? Cholesterol stones with small polyps. 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 enlarged in size (16.7 cm), normal in outline and echotexture. No focal lesion seen. splenic vein is dilated at the hilum.

Right kidney is marginally small in size (8.8 cm), normal in outline and echogenicity. Corticomedullary 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 echo pattern. No focal lesion seen.

No free fluid is seen.

Opinion: ECHOGENIC FOCI IN GALL BLADDER? CHOLESTEROL STONES WITH GB POLYPS SPLENOMEGALY WITH DILATED SPLENIC VEIN COARSE ECHOTEXTURE OF LIVER MINIMALLY SMALL RIGHT KIDNEY

Suggested clinical / CECT ABDOMEN correlation.

Dr. ADITI PANWAR PMC - 41230 Consultant Radiologist

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

SHYAM BABU SINGH 54 M Study Date: 23/03/2024 Patient ID: 13049279 Accession #: Alt ID: DOB: Age: Gender: M Ht: Wt: BSA: Institution: Fortis MEDCENTRE, Chandigarh Referring Physician: Performed By: Physician of Record: Performed By: Performed By:

Images



Signature

Signature: Name(Print):

Date:



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

DEPARTMENT OF FMC-RADIOLOGY LAB

Date: 23/Mar/2024

Name: Mr. Shyam Babu Singh Age | Sex: 54 YEAR(S) | Male Order Station : FRONTOFFICE-FMC Bed Name : UHID | Episode No : 13049279 | 4220/24/10021 Order No | Order Date: 10021/PN/OP/2403/10779 | 23-Mar-2024 Admitted On | Reporting Date : 23-Mar-2024 11:03:38 Order Doctor Name : Dr.SELF .

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.

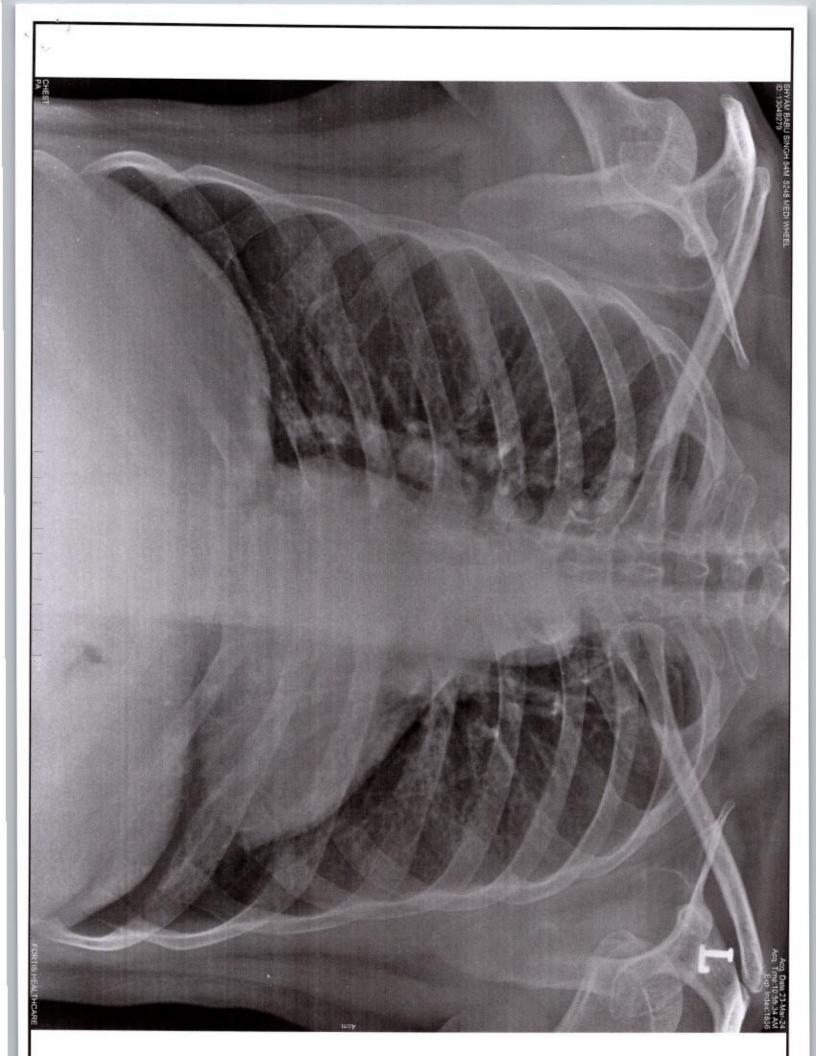
Please correlate clinically and with other relevant investigations.

Dr. ADITI PANWAR

PMC - 41230

Consultant Radiologist

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



CARDIOPRIN

SCO 11, Sector 11 D Chandigarh Station Telephone:

EXERCISE STRESS TEST REPORT

Age: 54yrs Gender: Male Race: Indian

DOB: 05.09.1969

Referring Physician: --

Attending Physician: DR MANJEET/DR VIJAY HARJAI

Patient Nan	ne: Babu, Shayam
Patient ID:	13049279
Height: 165	cm
Weight: 65	kg

Study Date: 23.03.2024 Test Type: --Protocol: BRUCE

Medications:

Medical History:

--

-

Reason for Exercise Test:

Exercise Test Summary

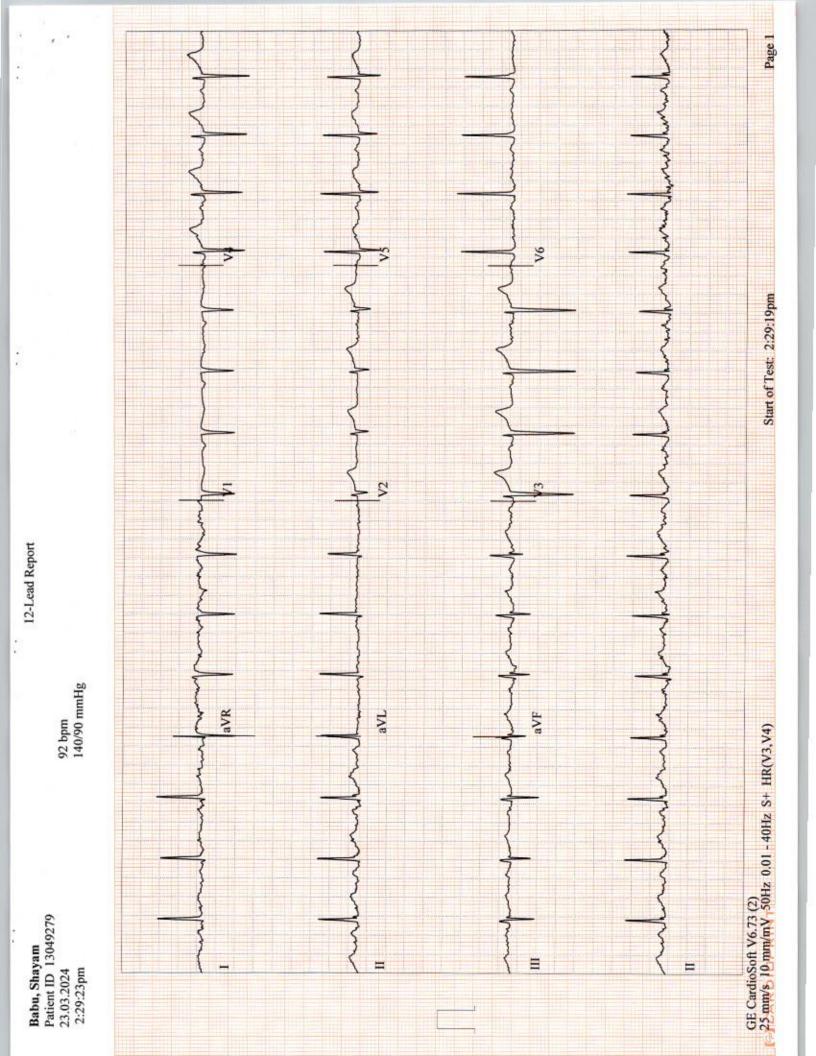
Phase Name	Stage Name	Time in Stage	Speed (km/h)	Grade (%)	HR (bpm)	BP (mmHg)	Comment
PRETEST	SUPINE	00:04	0.00	0.00	93	140/90	
	STANDING	00:14	0.00	0.00	95	110/20	
EXERCISE	STAGE 1	03:00	2.70	10.00	131	140/90	
	STAGE 2	03:00	4.00	12.00	146	150/100	
	STAGE 3	03:00	5.50	14.00	162	160/100	
	STAGE 4	00:01	5.60	14.00	162	100/100	
RECOVERY		02:10	0.00	7.70	130	170/110	

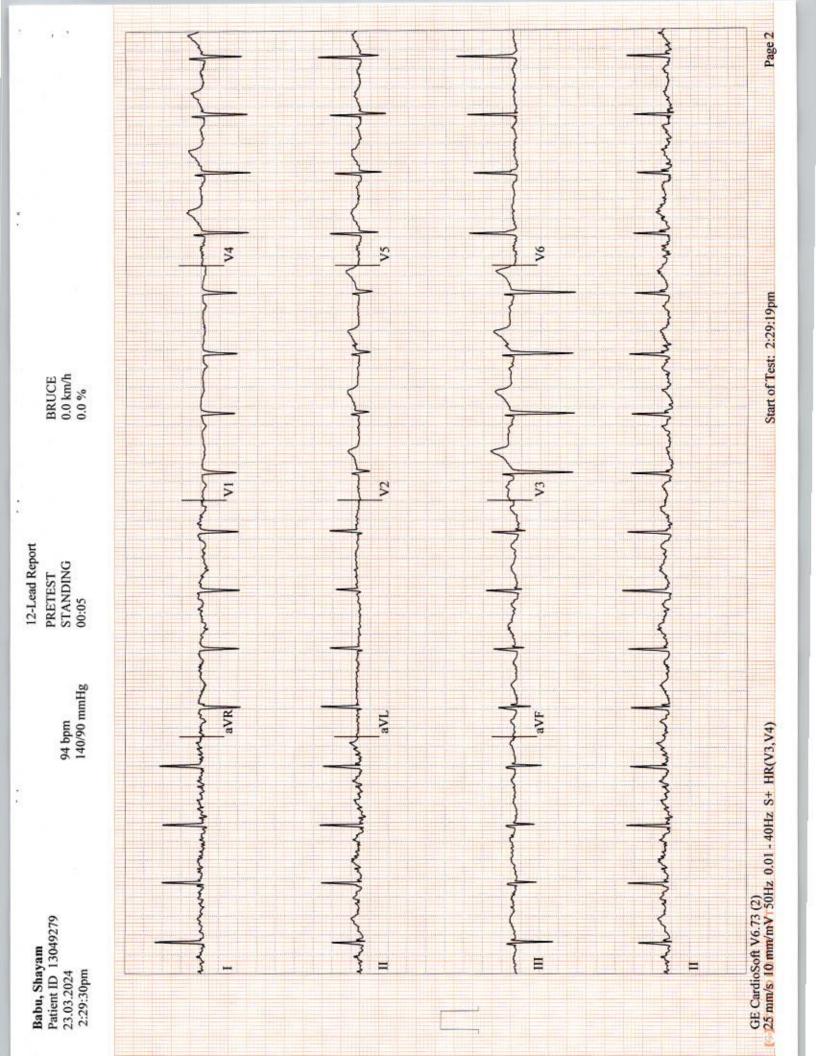
The patient exercised according to the BRUCE for 9:01 min:s, achieving a work level of Max. METS: 10.30. The resting heart rate of 92 bpm rose to a maximal heart rate of 162 bpm. This value represents 97 % of the maximal, age-predicted heart rate. The resting blood pressure of 140/90 mmHg, rose to a maximum blood pressure of 170/110 mmHg. The exercise test was stopped due to Exaggerated BP increase, Target heart rate achieved.

Interpretation

Summary: Resting ECG: normal. Functional Capacity: normal. HR Response to Exercise: appropriate. BP Response to Exercise: resting hypertension - exaggerated response. Chest Pain: none. Arrhythmias: none. ST Changes: none.

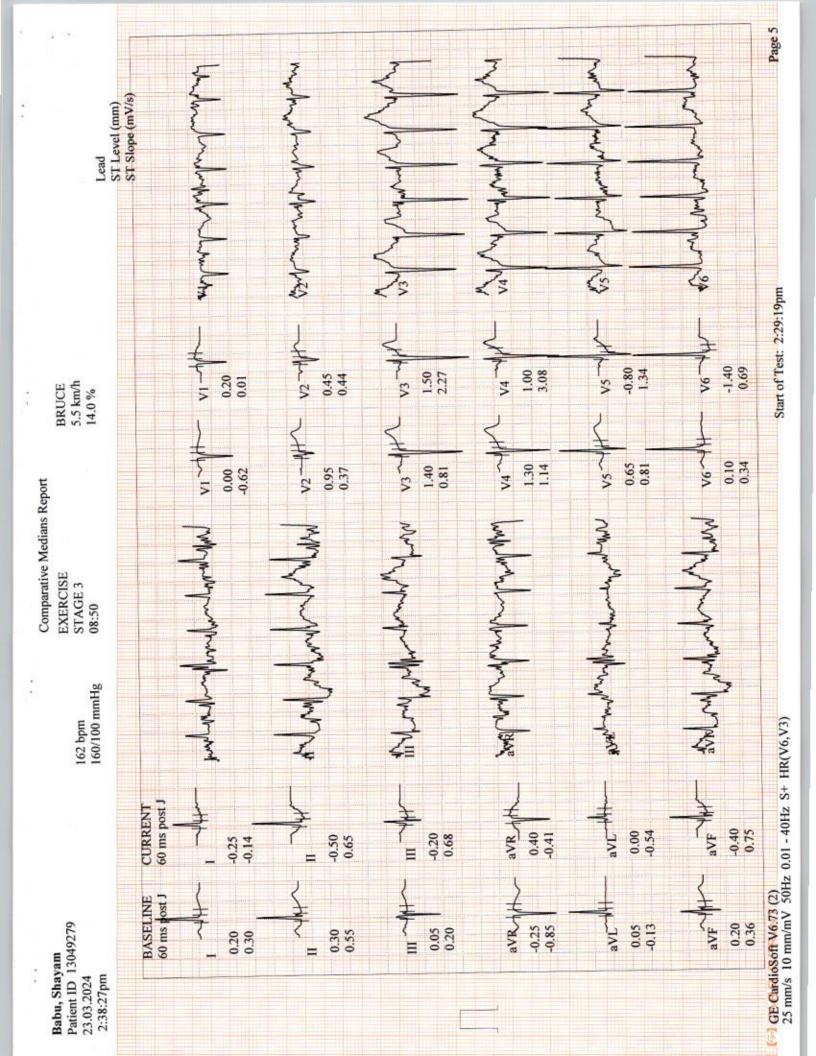
Conclusions

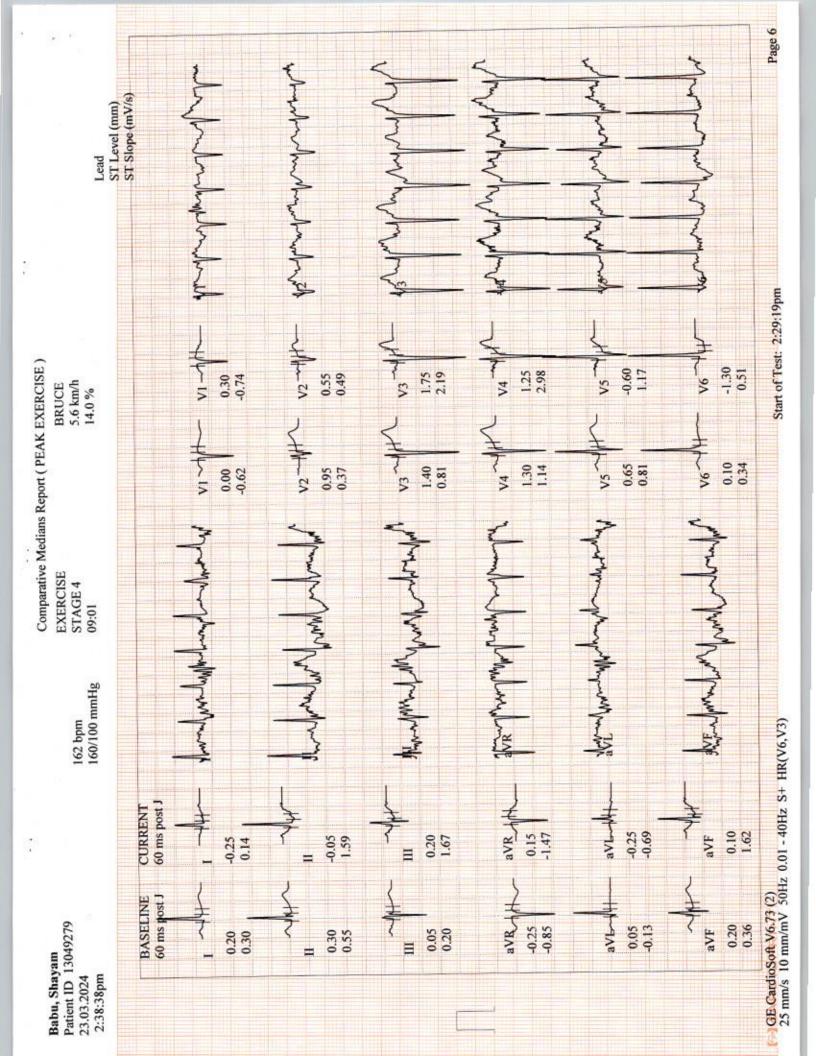


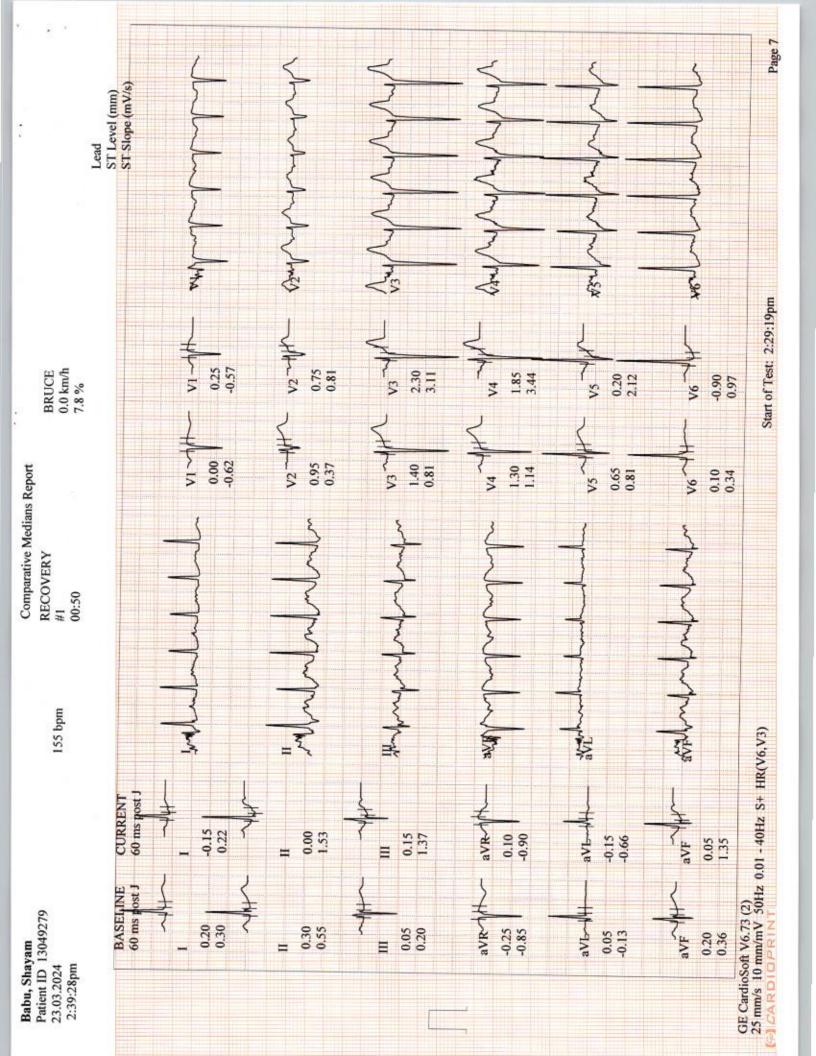


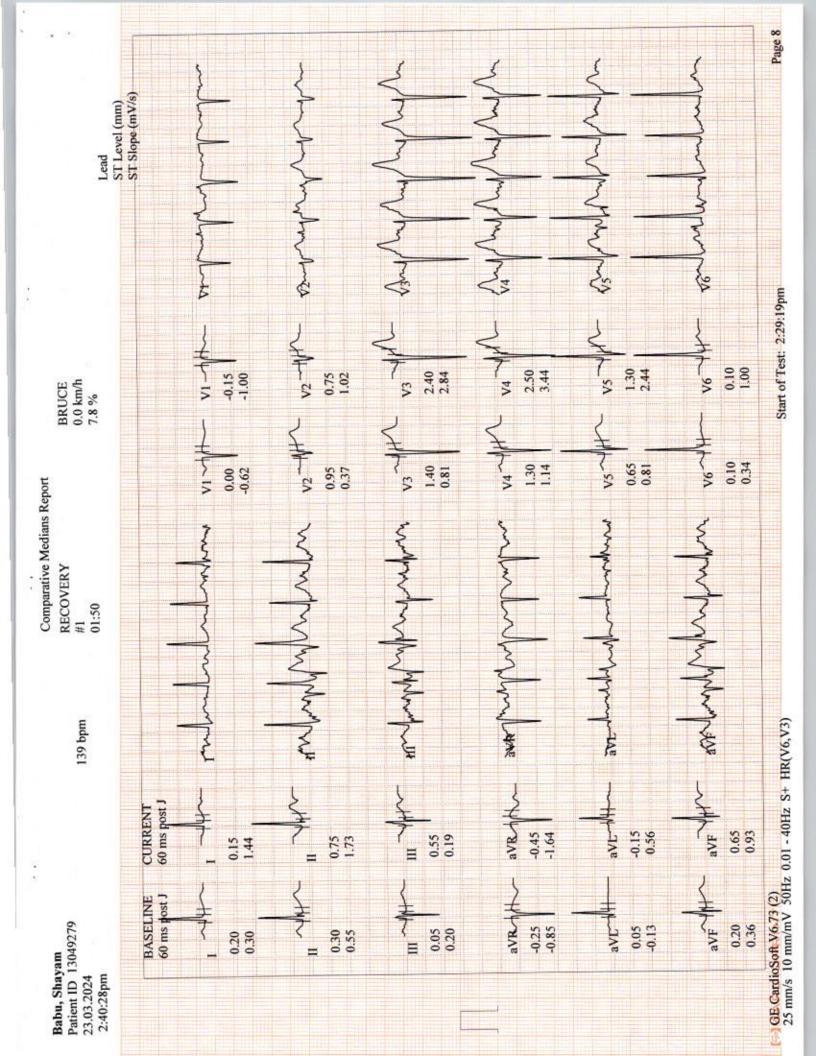
Lead ST Level (mm)	ST Slope (mV/s)	mohandomohander	markon Am Jun Jun Jung	- In many hard	may my my my my	man man man has	value and when he walked	Darre 3
eport BRUCE 2.7 km/h 10.0 %		VI	v2-W+- v2-W+- 0.95 0.35 0.37 0.36	V3-V4- V3-V4- V3-V4- V3-V4- V3-V4- V3-V4- V3-V4- V3-V4- V3-V4-V4-V4-V4-V4-V4-V4-V4-V4-V4-V4-V4-V4-	V4 V4 V4 V4 V1 1.30 0.80 0.80 0.80 1.142	V5 44 V5 46 0.65 0.05 0.81 0.93	v6 + + + v6 + + + - v6 - 0.10 - 0.60 0.34 0.28	Start of Test: 2:20-10mm
Comparative Medians Report 133 bpm EXERCISE 140/90 mmHg 02:50		for the multimentant of the south of the second	My Manufar Marine		stry from how have been a bridge	everth much many with my Munder	sould when he was however	(V6,V3)
	60 ms post J	- 1 ~444	- II ++++	- III	ave.	avr.44	- avr +++	0.01 - 40Hz S+ HR
Babu, Shayam Patient ID 13049279 23.03.2024 2:32:27pm	BASELINE 60 ms post J		11 744	ш 0.05 0.20	aVR-1	avr. 111	avr 11	GE CardioSoft V6.73 (2) 25 mm/s 10 mm/mV 50Hz 0.01 - 40Hz S+ HR(V6.V3)

Lead	S.I. Level (mm) ST Slope (mV/s)	handrahan por	Martin Martin	In In have been and the	In I when have	Involution -	serlindundundundund	9pm Page 4
BRUCE 4.0 km/h 12.0 %		- VI - 11	- v2-111-	0.85 1.14	V4-11	V5 -4 4	V6-140	Start of Test: 2:29:19pm
		V1 ~1 +++	26:0 75:0	V3-440	V4 - 44	V5 44	V6 + + +	St
Comparative Medians Report EXERCISE STAGE 2 05:50		Lupular Manuel	1 marshar	Andread	Indrading	al marken and	-	
146 bpm 150/100 mmHg		endrandium.	Mr Man	month	averyman	at Bull man marker h	Munhappan	(V6,V3)
	CURRENT 60 ms post J	- 1	11	0.68	aVR-1	aVL-Wh	aVF~/H/	25 mm/s 10 mm/mV 50Hz 0.01 - 40Hz S+ HR(V6,V3)
Babu, Shayam Patient ID 13049279 23.03.2024 2:35:27pm	BASELINE 60 ms post J	0.20	II 744	111	aVR/1//	aVL-14	ave 14	0 mm/mV S0Hz (
Babu, Shayam Patient ID 1304 23.03.2024 2:35:27pm							GE Candio	25 mm/s 1 [5-] CARDI













PATIENT NAME : SHYAM BABU SINGH REF. DOCTOR : SELF ACCESSION NO : 0006XC024723 AGE/SEX :54 Years Male FORTIS MOHALI-CHC -SPLZD PATIENT ID DRAWN :23/03/2024 09:21:00 : FH.13049279 FORTIS HOSPITAL # MOHALI, CLIENT PATIENT ID: UID:13049279 RECEIVED : 23/03/2024 19:38:36 MOHALI 160062 ABHA NO REPORTED :24/03/2024 01:29:58 : 7087030817

CLINICAL INFORMATION :

UID:13049279 REQNO-1681647 CORP-OPD BILLNO-10021240PCS005248 BILLNO-10021240PCS005248

Test Report Status	<u>Preliminary</u>	Results	Biological Reference Interval Units		
	HA	NEMATOLOGY - CBC			
CBC-5, EDTA WHOLE	BLOOD				
BLOOD COUNTS, EDT	TA WHOLE BLOOD				
HEMOGLOBIN (HB) METHOD : SLS- HEMOGLOB	IN DETECTION METHOD	11.9 Low	13.0 - 17.0	g/dL	
RED BLOOD CELL (R METHOD : HYDRODYNAMIC	,	3.67 Low	4.5 - 5.5	mil/µL	
WHITE BLOOD CELL METHOD : FLOWCYTOMETRY	· · ·	2.92 Low	4.0 - 10.0	thou/µL	
PLATELET COUNT	FOCUSING METHOD / MICROSCOPY	47 C.Low	150 - 410	thou/µL	
Comments					
PLATELETS ARE DONE MA					
HEMATOCRIT (PCV) METHOD : HYDRODYNAMIC	FOCUSING	41.6	40.0 - 50.0	%	
MEAN CORPUSCULA METHOD : CALCULATED PAR	R VOLUME (MCV)	113.4 High	83.0 - 101.0	fL	
	R HEMOGLOBIN (MCH)	32.4 High	27.0 - 32.0	pg	
MEAN CORPUSCULA CONCENTRATION(MO METHOD : CALCULATED PAR	R HEMOGLOBIN CHC)	28.6 Low	31.5 - 34.5	g/dL	
RED CELL DISTRIBU METHOD : CALCULATED PAR	TION WIDTH (RDW)	14.3 High	11.6 - 14.0	%	
MENTZER INDEX METHOD : CALCULATED PAR		30.9			
METHOD : CALCULATED PAR MEAN PLATELET VOI METHOD : CALCULATED PAR	LUME (MPV)	12.9 High	6.8 - 10.9	fL	

Subhijit Koul

Shafua

Dr. Subhijit kaur (MD, Pathology) Senior Resident, 49300

Dr. Shafira Garg (MD, Pathology) Attending Consultant, 47150



Dr. Irneet Mundi (MD,DNB Pathology) Associate Consultant, 34080 Page 1 Of 15



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PATIENT NAME : SH	IYAM BABU SINGH	REF. DOCTOR : SELF				
FORTIS MOHALI-CHC - FORTIS HOSPITAL # N MOHALI 160062 7087030817	-	ACCESSION NO: 0006 PATIENT ID : FH.1: CLIENT PATIENT ID:UII ABHA NO :	8049279 DF 0:13049279 RE	RAWN :	54 Years 23/03/2024 23/03/2024 24/03/2024	19:38:36
CLINICAL INFORMATIO	DN :		ł			
UID:13049279 REQNO CORP-OPD BILLNO-10021240PCS BILLNO-10021240PCS	005248					
Test Report Status	Preliminary	Results	Biological Re	ference	Interval l	Inits

WBC DIFFERENTIAL COUNT

NEUTROPHILS	61	40.0 - 80.0	%
METHOD : FLOW CYTOMETRY+LEISHMAIN STAIN+MICROSCOPY LYMPHOCYTES	29	20.0 - 40.0	%
METHOD : FLOW CYTOMETRY+LEISHMAIN STAIN+MICROSCOPY			0/
MONOCYTES METHOD : FLOW CYTOMETRY+LEISHMAIN STAIN+MICROSCOPY	8	2.0 - 10.0	%
EOSINOPHILS	2	1 - 6	%
METHOD : FLOW CYTOMETRY+LEISHMAIN STAIN+MICROSCOPY BASOPHILS	00	0 - 2	%
METHOD : FLOW CYTOMETRY+LEISHMAIN STAIN+MICROSCOPY	00	0 - 2	70
ABSOLUTE NEUTROPHIL COUNT	1.78 Low	2.0 - 7.0	thou/µL
METHOD : CALCULATED PARAMETER ABSOLUTE LYMPHOCYTE COUNT	0.85 Low	1.0 - 3.0	thou/µL
METHOD : CALCULATED PARAMETER			
ABSOLUTE MONOCYTE COUNT METHOD : CALCULATED PARAMETER	0.23	0.2 - 1.0	thou/µL
ABSOLUTE EOSINOPHIL COUNT	0.06	0.02 - 0.50	thou/µL
METHOD : CALCULATED PARAMETER NEUTROPHIL LYMPHOCYTE RATIO (NLR)	2.1		
METHOD : CALCULATED PARAMETER	2.1		

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.

Subhijit Koul

Shafia

Dr. Subhijit kaur (MD, Pathology) Senior Resident, 49300

Dr. Shafira Garg (MD, Pathology) Attending Consultant, 47150

Dr. Irneet Mundi (MD,DNB Pathology) Associate Consultant, 34080

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CLINICAL INFORMATION :

UID:13049279 REQNO-1681647 CORP-OPD BILLNO-10021240PCS005248 BILLNO-10021240PCS005248

Test Report Status	Preliminary	Results	val Units	
		HAEMATOLOGY		
ERYTHROCYTE SEDI	MENTATION RATE (ESP	R <u>),EDTA BLOOD</u>		
E.S.R METHOD : WESTERGREN ME	THOD	66 High	0 - 14	mm at 1 hr
GLYCOSYLATED HEM	IOGLOBIN(HBA1C), ED	TA WHOLE BLOOD		
HBA1C		6.0 High	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 AVERAG METHOD : CALCULATED PAR		125.5 High	< 116.0	mg/dL

Interpretation(s)

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

Erythrocyte sedimentation rate (ESR) is a test that indirectly measures the degree of inflammation present in the body. The test actually measures the rate of fall (sedimentation) of erythrocytes in a sample of blood that has been placed into a tall, thin, vertical tube. Results are reported as the 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, 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

LIMITATIONS

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)

Subhijit Koul

Shafua



Dr. Subhijit kaur (MD, Pathology) Senior Resident, 49300 Dr. Shafira Garg (MD, Pathology) Attending Consultant,47150 Dr. Irneet Mundi (MD,DNB Pathology) Associate Consultant, 34080





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REF. DOCTOR : SELF PATIENT NAME : SHYAM BABU SINGH ACCESSION NO : 0006XC024723 AGE/SEX :54 Years Male FORTIS MOHALI-CHC -SPLZD :23/03/2024 09:21:00 PATIENT ID : FH.13049279 DRAWN FORTIS HOSPITAL # MOHALI, CLIENT PATIENT ID: UID:13049279 RECEIVED : 23/03/2024 19:38:36 MOHALI 160062 REPORTED :24/03/2024 01:29:58 ABHA NO : 7087030817 **CLINICAL INFORMATION :** UID:13049279 REQNO-1681647 CORP-OPD BILLNO-10021240PCS005248 BILLNO-10021240PCS005248 **Test Report Status** Results **Biological Reference Interval** Units <u>Preliminary</u>

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. 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. 1. 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.
 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

Subhijit Koul

Dr. Subhijit kaur (MD, Pathology) Senior Resident, 49300

Chafua

Dr. Shafira Garg (MD, Pathology) Attending Consultant, 47150



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PATIENT NAME : SHYAM BABU SINGH REF. DOCTOR : SELF ACCESSION NO : 0006XC024723 AGE/SEX :54 Years Male FORTIS MOHALI-CHC -SPLZD PATIENT ID DRAWN :23/03/2024 09:21:00 : FH.13049279 FORTIS HOSPITAL # MOHALI, CLIENT PATIENT ID: UID:13049279 RECEIVED : 23/03/2024 19:38:36 MOHALI 160062 ABHA NO REPORTED :24/03/2024 01:29:58 : 7087030817

CLINICAL INFORMATION :

UID:13049279 REQNO-1681647 CORP-OPD BILLNO-10021240PCS005248 BILLNO-10021240PCS005248

est Report Status <u>Preliminary</u>	Results Biological Reference Interval		
	BIOCHEMISTRY		
VER FUNCTION PROFILE, SERUM			
ILIRUBIN, TOTAL METHOD : DIAZONIUM ION, BLANKED (ROCHE)	1.40 High	UPTO 1.2	mg/dL
	0.59 High	0.00 - 0.30	mg/dL
ILIRUBIN, INDIRECT METHOD : CALCULATED PARAMETER	0.81 High	0.00 - 0.60	mg/dL
OTAL PROTEIN METHOD : BIURET	8.3	6.6 - 8.7	g/dL
LBUMIN METHOD : BROMOCRESOL GREEN	4.2	3.97 - 4.94	g/dL
IOBULIN	4.1 High	2.0 - 4.0 Neonates - Pre Mature: 0.29 - 1.04	g/dL
METHOD : CALCULATED PARAMETER _BUMIN/GLOBULIN RATIO METHOD : CALCULATED PARAMETER	1.0	1.0 - 2.0	RATIO
SPARTATE AMINOTRANSFERASE(AST/SGOT)	38	0 - 40	U/L
LANINE AMINOTRANSFERASE (ALT/SGPT)	27	0 - 41	U/L
LKALINE PHOSPHATASE	130 High	40 - 129	U/L
AMMA GLUTAMYL TRANSFERASE (GGT) METHOD : GAMMA GLUTAMYLCARBOXY 4NITROANILIDE	446 High	8 - 61	U/L
ACTATE DEHYDROGENASE METHOD : LACTATE -PYRUVATE UV	146	135 - 225	U/L

GLUCOSE FASTING, FLUORIDE PLASMA

FBS (FASTING BLOOD SUGAR) METHOD : HEXOKINASE

Ritu Pankoy

Dr. Ritu Pankaj (MD,Pathology), PDCC Additional Director, 30897

Ms. Hardeep Kaur, M.Sc. Biochemistry

140 High

74 - 106

mg/dL

Meenahsh Malhotra

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PATIENT NAME : SHYAM BABU SINGH	REF. DOCTOR : SELF			
FORTIS MOHALI-CHC -SPLZD			AGE/SEX : 54 Years Male DRAWN : 23/03/2024 09:21:00	
FORTIS HOSPITAL # MOHALI,				
MOHALI 160062	CLIENT PATIENT ID: U	i	RECEIVED : 23/03/2024 19:38:36	
7087030817	ABHA NO :	l F	REPORTED :24/03/2024 01:29:58	
CLINICAL INFORMATION :				
UID:13049279 REQNO-1681647				
CORP-OPD				
BILLNO-10021240PCS005248				
BILLNO-10021240PCS005248				
Test Report Status <u>Preliminary</u>	Results	Biological R	eference Interval Units	
<u>BLOOD UREA NITROGEN (BUN), SERUM</u>				
BLOOD UREA NITROGEN METHOD : UREASE - UV	11	6 - 20	mg/dL	
URIC ACID, SERUM URIC ACID METHOD : URICASE, COLORIMETRIC	4.9	3.4 - 7.0	mg/dL	
CREATININE EGFR				
CREATININE METHOD : ALKALINE PICRATE-KINETIC	1.00	0.70 - 1.20	mg/dL	
AGE	54		years	
GLOMERULAR FILTRATION RATE (MALE)	89	damage wit 89- 60 mild decrea 59-30 moderate d 29-15 severe decr < 15 kidney	ninimal kidney h normal GFR se ecrease ease	

Interpretation(s)

Ritu Pankay

Dr. Ritu Pankaj (MD,Pathology), PDCC Additional Director, 30897

Ms. Hardeep Kaur, M.Sc. Biochemistry

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REF. DOCTOR : SELF PATIENT NAME : SHYAM BABU SINGH ACCESSION NO : 0006XC024723 AGE/SEX :54 Years Male FORTIS MOHALI-CHC -SPLZD :23/03/2024 09:21:00 PATIENT ID : FH.13049279 DRAWN FORTIS HOSPITAL # MOHALI, CLIENT PATIENT ID: UID:13049279 RECEIVED : 23/03/2024 19:38:36 MOHALT 160062 REPORTED :24/03/2024 01:29:58 ABHA NO 7087030817 **CLINICAL INFORMATION :** UID:13049279 REQNO-1681647 CORP-OPD BILLNO-10021240PCS005248 BILLNO-10021240PCS005248 **Test Report Status** Results Biological Reference Interval Units **Preliminary GLUCOSE POST-PRANDIAL, PLASMA** 141 High PPBS(POST PRANDIAL BLOOD SUGAR) Non-Diabetes ma/dL 70 - 140 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 extertion (eg, bernolysis and ineffective erythropoiesis), decreased bilirubin metabolism (eg, hereditary and neonatal jaundice). Conjugated (direct) bilirubin in Viral hepatitis, Drug reactions, Alcoholic liver disease Conjugated (direct) bilirubin is also elevated more than unconjugated (indirect) bilirubin in Viral hepatitis, Drug reactions, Alcoholic liver disease Conjugated (direct) bilirubin is also elevated more than unconjugated (indirect) bilirubin when there is some kind of blockage of the bile ducts like in Gallstones getting into the bile ducts, tumors &Scarring of the bile ducts. Increased unconjugated (indirect) bilirubin may be a result of Hemolytic or pernicious anemia, Transfusion reaction & a common metabolic condition termed Gilbert syndrome, due to low levels of the enzyme that attaches sugar molecules to bilirubin. AST is an enzyme found in various parts of the body. AST is found in the liver, heart, skeletal muscle, kidneys, brain, and red blood cells, and it is commonly measured clinically as a marker for liver health. AST levels increase during chronic viral hepatitis, blockage of the bile duct, cirrhosis of the liver, liver cancer, kidney failure, hemolytic anemia, pancreatitis, hemochromatosis. AST levels may also increase after a heart attack or strenuous activity. ALT test measures the amount of this enzyme in the blood. ALT is found mainly in the liver, but also in smaller amounts in the kidneys, heart, muscles, and pancreas. It is commonly measured as a part of a diagnostic evaluation of hepatocellular injury, to determine liver health.AST levels increase during acute hepatitis, sometimes due to a viral infection, ischemia to the liver, chronic hepatitis, obstruction of bile ducts, cirrhosis. ALP is a protein found in almost all body tissues. Tissues with higher amounts of ALP include the liver, bile ducts and bone. Elevated ALP levels are seen in Biliary obstruction, Osteoblastic bone tumors, osteomalacia, hepatitis, Hyperparathyroidism, Leukemia, Lymphoma, Pagets disease, Rickets, Sarcoidosis etc. Lower-than-normal ALP levels seen in Hypophosphatasia, Malnutrition, Protein deficiency, Wilsons disease. GGT is an enzyme found in cell membranes of many tissues mainly in the liver, kidney and pancreas. It is also found in other tissues including intestine, spleen, heart, brain and seminal vesicles. The highest concentration is in the kidney, but the liver is considered the source of normal enzyme activity. Serum GGT has been widely used as an index of liver dysfunction. Elevated serum GGT activity can be found in diseases of the liver, biliary system and pancreas. Conditions that increase serum GGT are obstructive liver disease, high alcohol consumption and use of enzyme-inducing drugs etc. Total Protein also known as total protein is a biochemical test for measuring the total amount of protein in serum. Protein in the plasma is made up of albumin and globulin. Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenstroms disease.Lower-than-normal levels may be due to: Agammaglobulinemia,Bleeding (hemorrhage),Burns,Glomerulonephritis,Liver disease, Malabsorption,Malnutrition,Nephrotic syndrome, Protein-losing enteropathy etc. Albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by:Liver disease like cirrhosis of the liver, nephrotic syndrome,protein-losing enteropathy,Burns,hemodilution,increased vascular permeability or decreased lymphatic clearance,malnutrition and wasting etc GLUCOSE FASTING,FLUORIDE PLASMA-**TEST DESCRIPTION** Normally, the glucose concentration in extracellular fluid is closely regulated so that a source of energy is readily available to tissues and 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 Ritu Pankay Meenahah Malhotra Page 7 Of 15

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Biochemistry

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PATIENT NAME : SHYAM BABU SINGH	REF. DOCTOR	: SELF
FORTIS MOHALI-CHC -SPLZD FORTIS HOSPITAL # MOHALI, MOHALI 160062 7087030817	ACCESSION NO: 0006XC024723 PATIENT ID : FH.13049279 CLIENT PATIENT ID:UID:13049279 ABHA NO :	AGE/SEX :54 Years Male DRAWN :23/03/2024 09:21:00 RECEIVED :23/03/2024 19:38:36 REPORTED :24/03/2024 01:29:58
CLINICAL INFORMATION : UID:13049279 REQNO-1681647 CORP-OPD BILLNO-10021240PCS005248 BILLNO-10021240PCS005248		
Test Report Status <u>Preliminary</u>	Results Biologi	cal Reference Interval Units

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 syndrome Causes of decreased levels-Low Zinc intake,OCP,Multiple Sclerosis GLUCOSE POST-PRANDIAL, PLASMA-Spectrophotometry Hexokinase

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CLINICAL INFORMATION :

UID:13049279 REQNO-1681647 CORP-OPD BILLNO-10021240PCS005248 BILLNO-10021240PCS005248

Test Report Status	<u>Preliminary</u>	Results	Biological Reference Interva	l Units
		BIOCHEMISTRY - LIPI	D	
LIPID PROFILE, SER	<u>UM</u>			
CHOLESTEROL, TOT		162	< 200 Desirable 200 - 239 Borderline High >/= 240 High	mg/dL
	XIDASE, ESTERASE, PEROXIDASE			
TRIGLYCERIDES		104	< 150 Normal 150 - 199 Borderline High 200 - 499 High >/= 500 Very High	mg/dL
METHOD : ENZYMATIC ASSA	ΑY			
HDL CHOLESTEROL		61 High	< 40 Low >/=60 High	mg/dL
METHOD : DIRECT MEASURE	E - PEG		_	
LDL CHOLESTEROL,	DIRECT	83	< 100 Optimal 100 - 129 Near or above optimal 130 - 160 Borderline High 161 - 189 High >/= 190 Very High	mg/dL
METHOD : CHOLESTEROL O	XIDASE, ESTERASE, PEROXIDASE			
NON HDL CHOLESTE	EROL	101	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	20.8	Desirable value : 10 - 35	mg/dL
METHOD : CALCULATED PAR	AMETER			
CHOL/HDL RATIO		2.7 Low	3.3-4.4 Low Risk 4.5-7.0 Average Risk 7.1-11.0 Moderate Risk > 11.0 High Risk	

Meenahshi Malhotra

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PATIENT NAME : SHYAM BABU SINGH	REF. DOCTOR : SELF		
	ACCESSION NO : 0006XC024723	AGE/SEX : 54 Years Male	
FORTIS MOHALI-CHC -SPLZD	PATIENT ID : FH.13049279	DRAWN :23/03/2024 09:21:00	
FORTIS HOSPITAL # MOHALI,	CLIENT PATIENT ID: UID:13049279	RECEIVED : 23/03/2024 19:38:36	
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CLINICAL INFORMATION :	· ·	•	
UID:13049279 REONO-1681647			

UID:13049279 REQNO-1681647 CORP-OPD BILLNO-1002124OPCS005248 BILLNO-1002124OPCS005248

Test Report Status	<u>Preliminary</u>	Results	Biological Reference Interval Units
LDL/HDL RATIO		1.4	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk
METHOD : CALCULATED PARA	AMETER		

Interpretation(s)

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PATIENT NAME : SHYAM BABU SINGH	REF. DOCTOR : SELF		
	ACCESSION NO : 0006XC024723	AGE/SEX : 54 Years Male	
FORTIS MOHALI-CHC -SPLZD	PATIENT ID : FH.13049279	DRAWN :23/03/2024 09:21:00	
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Test Report Status	<u>Preliminary</u>	Results	Biological Reference Interval Units
	CL	INICAL PATH - URINAL	YSIS
URINALYSIS			
PHYSICAL EXAMINA	TION, URINE		
COLOR METHOD : MANUAL EXAMIN	ΑΠΟΝ	YELLOW	
APPEARANCE METHOD : MANUAL EXAMINA	ΑΤΙΟΝ	CLEAR	

CHEMICAL EXAMINATION, URINE

PH	6.0	4.7 - 7.5
METHOD : DOUBLE INDICATOR PRINCIPLE		
SPECIFIC GRAVITY	1.020	1.003 - 1.035
METHOD : REFLECTANCE PHOTOMETRY (IONIC CONCENTRATION)		
PROTEIN	NOT DETECTED	NOT DETECTED
METHOD : REFLECTION PHOTOMETRY (PROTEIN ERROR INDICATOR)		
GLUCOSE	NOT DETECTED	NOT DETECTED
METHOD : REFLECTANCE PHOTOMETRY (GLUCOSE OXIDASE METHO	D)	
KETONES	NOT DETECTED	NOT DETECTED
METHOD : REFLECTION PHOTOMETRY (NITROPRUSSIDE)		
BLOOD	NOT DETECTED	NOT DETECTED
METHOD : REFLECTANCE PHOTOMETRY (BENZIDINE REACTION)		
BILIRUBIN	NOT DETECTED	NOT DETECTED
METHOD : REFLECTANCE SPECTROPHOTOMETRY (DIAZO REACTION)		
UROBILINOGEN	NORMAL	NORMAL
METHOD : REFLECTANCE PHOTOMETRY (EHRLICH'S REACTION)		
NITRITE	NOT DETECTED	NOT DETECTED
METHOD : REFLECTANCE SPECTROPHOTOMETRY (DIAZO REACTION)		

MICROSCOPIC EXAMINATION, URINE

Shafua

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reet

Dr. Irneet Mundi (MD,DNB Pathology) Associate Consultant, 34080

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Test Report Status	Preliminary	Results	Biological Reference Ir	nterval Units
			NOT DETECTED	
RED BLOOD CELLS		NOT DETECTED	NOT DETECTED	/HPF
PUS CELL (WBC'S)		NOT DETECTED	0-5	/HPF
EPITHELIAL CELLS		NOT DETECTED	0-5	/HPF
CASTS		NOT DETECTED		
CRYSTALS		NOT DETECTED		
BACTERIA METHOD : REFLECTANCE SPECTF	ROPHOTOMETRY	NOT DETECTED	NOT DETECTED	
YEAST		NOT DETECTED	NOT DETECTED	

Interpretation(s)

Shafia

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PATIENT NAME : SHYAM BABU SINGH	REF. DOCTOR :	SELF
	ACCESSION NO : 0006XC024723	AGE/SEX : 54 Years Male
FORTIS MOHALI-CHC -SPLZD	PATIENT ID : FH.13049279	DRAWN :23/03/2024 09:21:00
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Test Report Status	Preliminary	Results	Biological Reference Interval	Units
		CLINICAL PATH - STOOL ANALYSIS		
STOOL: OVA & PARA	SITE	RESULT PENDING		
PHYSICAL EXAMINA	TION,STOOL	RESULT PENDING		
CHEMICAL EXAMINA	TION, STOOL	RESULT PENDING		
MICROSCOPIC EXAM	INATION, STOOL	RESULT PENDING		

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PATIENT NAME : SHYAM BABU SINGH	REF. DOCTOR : SELF		
	ACCESSION NO : 0006XC024723	AGE/SEX : 54 Years Male	
FORTIS MOHALI-CHC -SPLZD	PATIENT ID : FH.13049279	DRAWN :23/03/2024 09:21:00	
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Test Report Status	<u>Preliminary</u>	Results	Biological Reference I	nterval Units				
SPECIALISED CHEMISTRY - HORMONE								
THYROID PANEL, SE	<u>RUM</u>	101.8	80.00 - 200.00	ng/dL				
T4		4.61 Low	5.10 - 14.10	µg/dL				
TSH (ULTRASENSITIVE)		4.040	0.270 - 4.200	µIU/mL				

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0.0 - 3.1



ng/mL

REF. DOCTOR : SELF PATIENT NAME : SHYAM BABU SINGH ACCESSION NO : 0006XC024723 AGE/SEX :54 Years Male FORTIS MOHALI-CHC -SPLZD :23/03/2024 09:21:00 PATIENT ID : FH.13049279 DRAWN FORTIS HOSPITAL # MOHALI, CLIENT PATIENT ID: UID:13049279 RECEIVED : 23/03/2024 19:38:36 MOHALI 160062 REPORTED :24/03/2024 01:29:58 ABHA NO : 7087030817

CLINICAL INFORMATION :

UID:13049279 REQNO-1681647 CORP-OPD BILLNO-10021240PCS005248 BILLNO-10021240PCS005248

PROSTATE SPECIFIC ANTIGEN

Test Report Status	Preliminary	Results	Biological Reference Interval	Units				
SPECIALISED CHEMISTRY - TUMOR MARKER								
PROSTATE SPECIFIC ANTIGEN, SERUM								

Interpretation(s)

PROSTATE SPECIFIC ANTIGEN, SERUM-- PSA is detected in the male patients with normal, benign hyperplastic and malignant prostate tissue and in patients with prostatitis. - PSA is not detected (or detected at very low levels) in the patients without prostate tissue (because of radical prostatectomy or cystoprostatectomy) and also in the female patients.

- It a suitable marker for monitoring of patients with Prostate Cancer and it is better to be used in conjunction with other diagnostic procedures.

0.921

- Serial PSA levels can help determine the success of prostatectomy and the need for further treatment, such as radiation, endocrine or chemotherapy and useful in detecting residual disease and early recurrence of tumor.

 Elevated levels of PSA can be also observed in the patients with non-malignant diseases like Prostatitis and Benign Prostatic Hyperplasia.
 Specimens for total PSA assay should be obtained before biopsy, prostatectomy or prostatic massage, since manipulation of the prostate gland may lead to elevated PSA (false positive) levels persisting up to 3 weeks.

- As per American urological guidelines, PSA screening is recommended for early detection of Prostate cancer above the age of 40 years. Following Age specific reference range can be used as a guide lines.

- Measurement of total PSA alone may not clearly distinguish between benign prostatic hyperplasia (BPH) from cancer, this is especially true for the total PSA values between 4-10 ng/mL.

- Total PSA values determined on patient samples by different testing procedures cannot be directly compared with one another and could be the cause of erroneous medical interpretations. Recommended follow up on same platform as patient result can vary due to differences in assay method and reagent specificity.

References

1. Burtis CA, Ashwood ER, Bruns DE. Teitz textbook of clinical chemistry and Molecular Diagnostics. 4th edition.

2. Williamson MA, Snyder LM. Wallach's interpretation of diagnostic tests. 9th edition.

End Of Report

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

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