

: 2208211070

CID#

SID# : 177805051563

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Name : MR.JYOTI PRAKASH MALLIK Registered : 23-Mar-2022 / 09:27

Age / Gender : 38 Years/Male Collected : 23-Mar-2022 / 09:27

Consulting Dr. : - Reported : 25-Mar-2022 / 10:36

Reg.Location : Andheri West (Main Centre) Printed : 25-Mar-2022 / 16:11

PHYSICAL EXAMINATION REPORT

History and Complaints:

Asymptomatic

EXAMINATION FINDINGS:

Height (cms): 159 cms Weight (kg): 64 kgs
Temp (0c): Skin: Normal

Blood Pressure 120/80 mm of Hg Nails: Normal

(mm/hg):

Pulse: 90/min Lymph Node: Right submandibular nodes

palpable

Systems

Cardiovascular: S1S2 audible

Respiratory: AEBE Genitourinary: NAD

GI System: Liver & Spleen not palpable

CNS: NAD

IMPRESSION:

Kindly consult your family physician with all your reports for the treatment of Diabetes Mellitus and Dyslipidemia.

ADVICE:

FBS=213.3 mg/dl., PPBS=311.6 mg/dl., HbA1C=12.6 %, Diabetes Mellitus urine glucose=3+, Glycosuria, Total cholesterol=230.2 mg/dl., Triglycerides=232.4 mg/dl., LDL=161.2 mg/dl, Dyslipidemia, Grade II fatty liver

CHIEF COMPLAINTS:

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1) Hypertension: NO 2) IHD NO

3) Arrhythmia NO

4) Diabetes Mellitus NO 5) Tuberculosis

NO 6) Asthama NO

7) Pulmonary Disease

NO

8) Thyroid/ Endocrine disorders NO

9) Nervous disorders NO

10) GI system NO

11) Genital urinary disorder NO

12) Rheumatic joint diseases or symptoms NO

13) Blood disease or disorder NO 14) Cancer/lump growth/cyst

NO 15) Congenital disease NO

16) Surgeries NO

17) Musculoskeletal System NO

PERSONAL HISTORY:

1) Alcohol Occasionally

Smoking 2) NO 3) Diet Mixed 4) Medication NO

*** End Of Report ***

Sanguta Manwanie

Dr.Sangeeta Manwani M.B.B.S. Reg.No.71083

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MEDI ASSIST - BOB HC PACKAGE MALE

	CBC (Complete	e Blood Count), Blood	
PARAMETER	RESULTS	BIOLOGICAL REF RANGE	METHOD
RBC PARAMETERS			
Haemoglobin	14.9	13.0-17.0 g/dL	Spectrophotometric
RBC	6.71	4.5-5.5 mil/cmm	Elect. Impedance
PCV	45.5	40-50 %	Measured
MCV	67.8	80-100 fl	Calculated
MCH	22.2	27-32 pg	Calculated
MCHC	32.8	31.5-34.5 g/dL	Calculated
RDW	16.4	11.6-14.0 %	Calculated
WBC PARAMETERS			
WBC Total Count	7160	4000-10000 /cmm	Elect. Impedance
WBC DIFFERENTIAL AND A	ABSOLUTE COUNTS		
Lymphocytes	30.5	20-40 %	
Absolute Lymphocytes	2183.8	1000-3000 /cmm	Calculated
Monocytes	5.7	2-10 %	
Absolute Monocytes	408.1	200-1000 /cmm	Calculated
Neutrophils	57.9	40-80 %	Calculated
Absolute Neutrophils	4145.6	2000-7000 /cmm	Calculated
Eosinophils	5.7	1-6 %	Calculated
Absolute Eosinophils	408.1	20-500 /cmm	Calculated
Basophils	0.2	0.1-2 %	Calculated
Absolute Basophils	14.3	20-100 /cmm	Calculated
Immature Leukocytes			

WBC Differential Count by Absorbance & Impedance method/Microscopy.

PLATELET PARAMETERS

LAILLIA		150000-400000 /cmm	Elect. Impedance
Platelet Count	263000	150000-400000 / CITIII	Fig. 1. Company of the Company of th
MPV	9.6	6-11 fl	Calculated
MIFV		11-18 %	Calculated
PDW	19.8	11-10 %	Catediates

RBC MORPHOLOGY

Hypochromia +
Microcytosis +

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Macrocytosis

Anisocytosis

Poikilocytosis

Mild Mild

Polychromasia

Target Cells

Basophilic Stippling

Normoblasts

Others

Elliptocytes-occasional

WBC MORPHOLOGY

PLATELET MORPHOLOGY

COMMENT

Features suggest thalassemia trait

Advice: Hemoglobin studies by HPLC, reticulocyte count.

Specimen: EDTA Whole Blood

ESR, EDTA WB

2-15 mm at 1 hr.

Westergren

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Dr. AMAR DASGUPTA, MD, PhD Consultant Hematopathologist **Director - Medical Services**

Dr.TRUPTI SHETTY M. D. (PATH) **Pathologist**

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Hexokinase

Hexokinase

MEDI ASSIST - BOB HC PACKAGE MALE

PARAMETER	RESULTS	BIOLOGICAL REF RANGE	METHOD

GLUCOSE (SUGAR) FASTING,

Fluoride Plasma

213.3

Non-Diabetic: < 100 mg/dl

Impaired Fasting Glucose:

Reported

100-125 mg/dl

Diabetic: >/= 126 mg/dl

GLUCOSE (SUGAR) PP, Fluoride 311.6

Plasma PP/R

Non-Diabetic: < 140 mg/dl Impaired Glucose Tolerance:

140-199 mg/dl

Diabetic: >/= 200 mg/dl

BLOOD UREA, Serum 15.2 BUN, Serum 7.1

CREATININE, Serum 0.71 eGFR. Serum 132

URIC ACID, Serum 4.1 12.8-42.8 mg/dl

6-20 mg/dl

0.67-1.17 mg/dl >60 ml/min/1.73sqm

3.5-7.2 mg/dl

Enzymatic Calculated

Kinetic

Calculated

Enzymatic







Dr.ANUPA DIXIT M.D.(PATH) Consultant Pathologist & Lab

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MEDI ASSIST - BOB HC PACKAGE MALE GLYCOSYLATED HEMOGLOBIN (HbA1c)

PARAMETER RESULTS BIOLOGICAL REF RANGE METHOD

Glycosylated Hemoglobin 12.6 Non-Diabetic Level: < 5.7 % (HbA1c), EDTA WB - CC Prediabetic Level: 5.7-6.4 %

Diabetic Level: >/= 6.5 %

Collected

Estimated Average Glucose 314.9 mg/dl Calculated (eAG), EDTA WB - CC

Intended use:

In patients who are meeting treatment goals, HbA1c test should be performed at least 2 times a year

In patients whose therapy has changed or who are not meeting glycemic goals, it should be performed quarterly

• For microvascular disease prevention, the HbA1C goal for non pregnant adults in general is Less than 7%.

Clinical Significance:

HbA1c, Glycosylated hemoglobin or glycated hemoglobin, is hemoglobin with glucose molecule attached to it.

 The HbA1c test evaluates the average amount of glucose in the blood over the last 2 to 3 months by measuring the percentage of glycosylated hemoglobin in the blood.

Test Interpretation:

 The HbA1c test evaluates the average amount of glucose in the blood over the last 2 to 3 months by measuring the percentage of Glycosylated hemoglobin in the blood.

HbA1c test may be used to screen for and diagnose diabetes or risk of developing diabetes.

To monitor compliance and long term blood glucose level control in patients with diabetes.

Index of diabetic control, predicting development and progression of diabetic micro vascular complications.

Factors affecting HbA1c results:

Increased in: High fetal hemoglobin, Chronic renal failure, Iron deficiency anemia, Splenectomy, Increased serum triglycerides, Alcohol ingestion, Lead/opiate poisoning and Salicylate treatment.

Decreased in: Shortened RBC lifespan (Hemolytic anemia, blood loss), following transfusions, pregnancy, ingestion of large amount of Vitamin E or Vitamin C and Hemoglobinopathies

Reflex tests: Blood glucose levels, CGM (Continuous Glucose monitoring)

References: ADA recommendations, AACC, Wallach's interpretation of diagnostic tests 10th edition.

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Dr.MEGHA SHARMA M.D. (PATH), DNB (PATH) Pathologist

Page 4 of 10

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MEDI ASSIST - BOB HC PACKAGE MALE PROSTATE SPECIFIC ANTIGEN (PSA)

PARAMETER

RESULTS

BIOLOGICAL REF RANGE

Collected

METHOD

TOTAL PSA, Serum

0.407

0.03-2.5 ng/ml

ECLIA

Clinical Significance:

- · PSA is detected in the serum of males with normal, benign hyper-plastic, and malignant prostate tissue.
- Monitoring patients with a history of prostate cancer as an early indicator of recurrence and response to treatment.
- Prostate cancer screening 4. The percentage of Free PSA (FPSA) in serum is described as being significantly higher in patients with BPH
 than in patients with prostate cancer.
 5. Calculation of % free PSA (ie. FPSA/TPSA x 100), has been suggested as way of improving the
 differentiation of BPH and Prostate cancer.

Interpretation:

Increased In- Prostate diseases, Cancer, Prostatitis, Benign prostatic hyperplasia, Prostatic ischemia, Acute urinary retention, Manipulations like Prostatic massage, Cystoscopy, Needle biopsy, Transurethral resection, Digital rectal examination, Radiation therapy, Indwelling catheter, Vigorous bicycle exercise, Drugs (e.g., testosterone), Physiologic fluctuations. Also found in small amounts in other cancers (sweat and salivary glands, breast, colon, lung, ovary) and in Skene glands of female urethra and in term placenta, Acute renal failure, Acute myocardial infarction.

Decreased In- Ejaculation within 24-48 hours, Castration, Antiandrogen drugs (e.g., finasteride), Radiation therapy, Prostatectomy, PSA falls 17% in 3 days after lying in hospital, Artifactual (e.g., improper specimen collection; very high PSA levels). Finasteride (5-falpha; reductase inhibitor) reduces PSA by 50% after 6 months in men without cancer.

Reflex Tests: % FREE PSA, USG Prostate

Limitations:

tPSA 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. If there is a change in the tPSA assay procedure used while monitoring therapy, then the tPSA values obtained upon changing over to the new procedure must be confirmed by parallelmeasurements with both methods. Immediate PSA testing following digital rectal examination, ejaculation, prostatic massage, indwelling catheterization, ultrasonography and needle biopsy of prostate is not recommended as they falsely elevate levels.

Reference:

- Wallach's Interpretation of diagnostic tests, 10th Edition
- Pack insert

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Page 5 of 10



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MEDI ASSIST - BOB HC PACKAGE MALE URINE EXAMINATION REPORT

PARAMETER	RESULTS	BIOLOGICAL REF RANGE	METHOD	
PHYSICAL EXAMINATION				
Color	Yellow	Pale Yellow		
Reaction (pH)	5.0	4.5 - 8.0	Chemical Indicator	
Specific Gravity	1.025	1.001-1.030	Chemical Indicator	
Transparency	Slight hazy	Clear	•	
Volume (ml)	15			
CHEMICAL EXAMINATION				
Proteins	Trace	Absent	pH Indicator	
Glucose	3+	Absent	GOD-POD	
Ketones	Absent	Absent	Legals Test	
Blood	Trace	Absent	Peroxidase	
Bilirubin	Absent	Absent	Diazonium Salt	
Urobilinogen	Normal	Normal	Diazonium Salt	
Nitrite	Absent	Absent	Griess Test	
MICROSCOPIC EXAMINATION				
Leukocytes(Pus cells)/hpf	6-8	0-5/hpf		
Red Blood Cells / hpf	Occasional	0-2/hpf		
Epithelial Cells / hpf	1-2			
Casts	Absent	Absent		

Absent

Absent

Less than 20/hpf

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Absent

Absent

8-10



Crystals

Others

Amorphous debris

Bacteria / hpf



Hasti D Dr.SHASHIKANT DIGHADE M.D. (PATH) **Pathologist**

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MEDI ASSIST - BOB HC PACKAGE MALE BLOOD GROUPING & Rh TYPING

PARAMETER

RESULTS

ABO GROUP

В

Rh TYPING

POSITIVE

NOTE: Test performed by automated column agglutination technology (CAT) which is more sensitive than conventional methods.

Specimen: EDTA Whole Blood and/or serum

Clinical significance:

ABO system is most important of all blood group in transfusion medicine

Limitations:

- ABO blood group of new born is performed only by cell (forward) grouping because allo antibodies in cord blood are of maternal origin.
- Since A & B antigens are not fully developed at birth, both Anti-A & Anti-B antibodies appear after the first 4 to 6 months of life. As a
 result, weaker reactions may occur with red cells of newborns than of adults.
- Confirmation of newborn's blood group is indicated when A & B antigen expression and the isoagglutinins are fully developed at 2 to 4
 years of age & remains constant throughout life.
- Cord blood is contaminated with Wharton's jelly that causes red cell aggregation leading to false positive result
- The Hh blood group also known as Oh or Bombay blood group is rare blood group type. The term Bombay is used to refer the phenotype
 that lacks normal expression of ABH antigens because of inheritance of hh genotype.

Refernces:

- 1. Denise M Harmening, Modern Blood Banking and Transfusion Practices- 6th Edition 2012. F.A. Davis company. Philadelphia
- 2. AABB technical manual

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Page 7 of 10



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MEDI ASSIST - BOB HC PACKAGE MALE LIPID PROFILE

<u>PARAMETER</u>	RESULTS	BIOLOGICAL REF RANGE	METHOD
CHOLESTEROL, Serum	230.2	Desirable: <200 mg/dl Borderline High: 200-239mg/dl High: >/=240 mg/dl	Enzymatic
TRIGLYCERIDES, Serum	232.4	Normal: <150 mg/dl Borderline-high: 150 - 199 mg/dl High: 200 - 499 mg/dl Very high:>/=500 mg/dl	Enzymatic
HDL CHOLESTEROL, Serum	37.7	Desirable: >60 mg/dl Borderline: 40 - 60 mg/dl Low (High risk): <40 mg/dl	Enzymatic
NON HDL CHOLESTEROL, Serum	192.5	Desirable: <130 mg/dl Borderline-high:130 - 159 mg/dl High:160 - 189 mg/dl Very high: >/=190 mg/dl	Calculated
LDL CHOLESTEROL, Serum	161.2	Optimal: <100 mg/dl Near Optimal: 100 - 129 mg/dl Borderline High: 130 - 159 mg/dl High: 160 - 189 mg/dl Very High: >/= 190 mg/dl	Calculated
VLDL CHOLESTEROL, Serum	31.3	< /= 30 mg/dl	Calculated
CHOL / HDL CHOL RATIO, Serum	6.1	0-4.5 Ratio	Calculated
LDL CHOL / HDL CHOL RATIO, Serum	4.3	0-3.5 Ratio	Calculated

Note: LDL test is performed by direct measurement.







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Page 8 of 10

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MEDI ASSIST - BOB HC PACKAGE MALE

THYROID FUNCTION TESTS

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PARAMETER	<u>RESULTS</u>	BIOLOGICAL REF RANGE	METHOD
Free T3, Serum	4.7	3.5-6.5 pmol/L	ECLIA
Free T4, Serum	15.3	11.5-22.7 pmol/L	ECLIA
sensitiveTSH, Serum	4.03	0.35-5.5 microIU/ml	ECLIA

Interpretation:

A thyroid panel is used to evaluate thyroid function and/or help diagnose various thyroid disorders.

Clinical Significance:

1)TSH Values between 5.5 to 15 microIU/ml should be correlated clinically or repeat the test with new sample as physiological factors can give falsely high TSH.

2)TSH values may be trasiently altered becuase of non thyroidal illness like severe infections, liver disease, renal and heart severe burns, trauma and surgery etc.

TSH	FT4/T4	FT3/T3	Interpretation
High	Normal	Normal	Subclinical hypothyroidism, poor compliance with thyroxine, drugs like amiodarone, Recovery phase of non-thyroidal illness, TSH Resistance.
High	Low	Low	Hypothyroidism, Autoimmune thyroiditis, post radio iodine Rx, post thyroidectomy, Anti thyroid drugs, tyrosine kinase inhibitors & amiodarone, amyloid deposits in thyroid, thyroid tumors & congenital hypothyroidism.
Low	High	High	Hyperthyroidism, Graves disease, toxic multinodular goiter, toxic adenoma, excess iodine or thyroxine intake, pregnancy related (hyperemesis gravidarum, hydatiform mole)
Low	Normal	Normal	Subclinical Hyperthyroidism, recent Rx for Hyperthyroidism, drugs like steroids & dopamine), Non thyroidal illness.
Low	Low	Low	Central Hypothyroidism, Non Thyroidal Illness, Recent Rx for Hyperthyroidism.
High	High	High	Interfering anti TPO antibodies, Drug interference: Amiodarone, Heparin, Beta Blockers, steroids & anti-

Diurnal Variation:TSH follows a diurnal rhythm and is at maximum between 2 am and 4 am , and is at a minimum between 6 pm and 10 pm. The variation is on the order of 50 to 206%. Biological variation:19.7%(with in subject variation)

Reflex Tests:Anti thyroid Antibodies, USG Thyroid ,TSH receptor Antibody. Thyroglobulin, Calcitonin

Limitations: Samples should not be taken from patients receiving therapy with high biotin doses (i.e. >5 mg/day) until atleast 8 hours following the last biotin administration.

Reference:

- 1.O.koulouri et al. / Best Practice and Research clinical Endocrinology and Metabolism 27(2013)
- 2.Interpretation of the thyroid function tests, Dayan et al. THE LANCET . Vol 357
- 3. Tietz , Text Book of Clinical Chemistry and Molecular Biology -5th Edition
- 4.Biological Variation:From principles to Practice-Callum G Fraser (AACC Press)

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Page 9 of 10



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MEDI ASSIST - BOB HC PACKAGE MALE LIVER FUNCTION TESTS

P	PARAMETER	RESULTS	BIOLOGICAL REF RANGE	METHOD
В	BILIRUBIN (TOTAL), Serum	0.51	0.1-1.2 mg/dl	Colorimetric
В	BILIRUBIN (DIRECT), Serum	0.23	0-0.3 mg/dl	Diazo
В	BILIRUBIN (INDIRECT), Serum	0.28	0.1-1.0 mg/dl	Calculated
Т	OTAL PROTEINS, Serum	6.9	6.4-8.3 g/dL	Biuret
	LBUMIN, Serum	4.8	3.5-5.2 g/dL	BCG
G	SLOBULIN, Serum	2.1	2.3-3.5 g/dL	Calculated
A	VG RATIO, Serum	2.3	1 - 2	Calculated
S	GOT (AST), Serum	15.1	5-40 U/L	NADH (w/o P-5-P)
S	GPT (ALT), Serum	21.7	5-45 U/L	NADH (w/o P-5-P)
G	SAMMA GT, Serum	52.9	3-60 U/L	Enzymatic
	LKALINE PHOSPHATASE, Gerum	64.0	40-130 U/L	Colorimetric

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Page 10 of 10

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X-RAY CHEST PA VIEW

Both lung fields are clear.

Both costo-phrenic angles are clear.

No hilar abnormality is seen.

The cardiac size and shape are within normal limits.

The aorta shows normal radiological features.

The trachea is central.

The domes of diaphragm are normal in position and outlines.

The skeleton under review appears normal.

IMPRESSION:

NO SIGNIFICANT ABNORMALITY IS DETECTED.

*** End Of Report ***

Dr.R K BHANDARI M.D., D.M.R.E CONSULTANT RADIOLOGIST

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Patient's Name : JYOTI PRAKASH MALLIK

Age:38 YRS/MALE

Date: 23.03.2022

Requesting Doctor: --

CID. No

: 2208211070

2D-ECHO & COLOUR DOPPLER REPORT

Structurally Normal : MV / AV / TV / PV. No significant valvular stenosis.

Trivial Mitral Regurgitation , Trivial Aortic Regurgitation Trivial Pulmonary Regurgitation ,

Trivial Tricuspid regurgitation. No Pulmonary arterial hypertension. PASP by TRjet vel.method = 16 mm Hg.

LV / LA / RA / RV - Normal in dimension. IAS / IVS is Intact.

No Left Ventricular Diastolic Dysfunction [LVDD]. No doppler evidence of raised LVEDP

No regional wall abnormality. No thinning / scarring / dyskinesia of LV wall noted. Normal LV systolic function. LVEF = 60 % by visual estimation.

No e/o thrombus in LA /LV. No e/o Pericardial effusion.

IVC normal in dimension with good inspiratory collapse. Normal RV systolic function (by TAPSE)

Impression:

NORMAL LV SYSTOLIC FUNCTION, LVEF = 60 % , NO RWMA, NO PAH, NO LVDD, NO LV HYPERTROPHY.

ADDRESS: 2nd Floor, Aston, Sundervan Complex, Above Mercedes Showroom, Andheri West - 400053

HEALTHLINE - MUMBAI: 022-6170-0000 | OTHER CITIES: 1800-266-4343



Patient Name : MR JYOTI PRAKASH MALLIK

Age: 38 Years / MALE

Ref Dr.

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Date: 23.03.2022

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USG WHOLE ABDOMEN

<u>LIVER:</u> Liver is normal in size (14.5cm. cranio-caudal) and **shows bright echotexture.** There is no intra-hepatic biliary radical dilatation. No evidence of any focal lesion.

GALL BLADDER: Gall bladder is distended and appears normal. Wall thickness is within normal limits. There is no evidence of any calculus.

PORTAL VEIN: Portal vein is normal. CBD: CBD is normal.

<u>PANCREAS</u>: Pancreas appears normal in echotexture. There is no evidence of any focal lesion or calcification. Pancreatic duct is not dilated.

<u>KIDNEYS</u>: Right kidney measures 10.7 x 4.8cm. Left kidney measures 10.7 x 5.4cm. Both kidneys are normal in shape and echotexture. Corticomedullary differentiation is maintained. There is no evidence of any hydronephrosis, hydroureter or calculus.

SPLEEN: Spleen is normal in size (9.2cm.), shape and echotexture. No focal lesion is seen.

<u>URINARY BLADDER:</u> Urinary bladder is distended and normal. Wall thickness is within normal limits.

<u>PROSTATE:</u> Prostate measures 3.7 x 3.3 x 2.8cm. and prostatic weight is 18.7g. Median lobe does not show significant hypertrophy.

No free fluid or significant lymphadenopathy is seen.

MPRESSION: Grade II fatty liver.

*** End of Report ***

DR. NIKHIL DEV MD. RADIOLOGIST

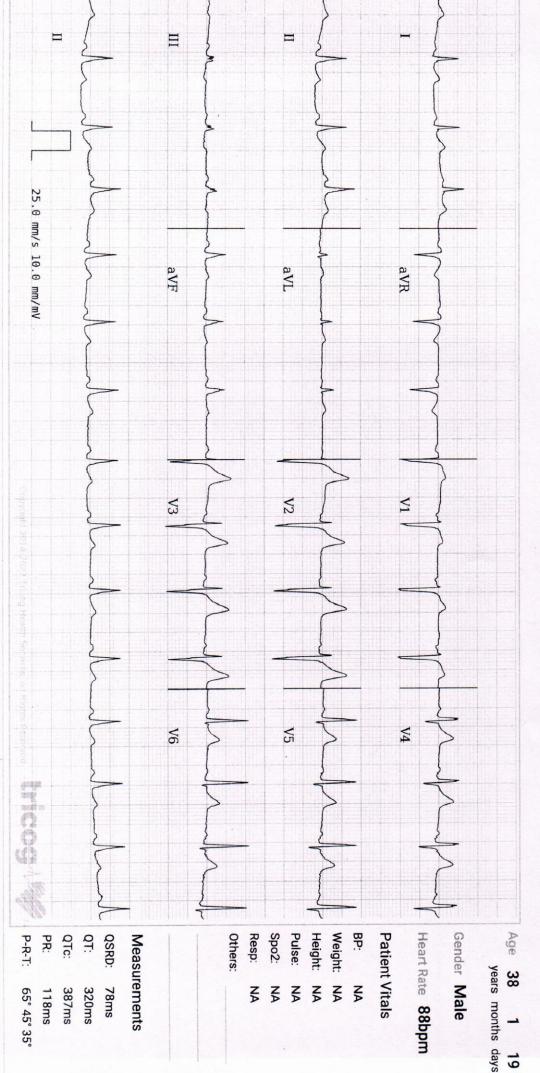


SUBURBAN DIAGNOSTICS - ANDHERI WEST

Patient ID: Patient Name: JYOTI PRAKASH MALLIK 2208211070

Date and Time: 23rd Mar 22 10:11 AM

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ECG Within Normal Limits: Sinus Rhythm, Normal Axis. Please correlate clinically.

REPORTED BY

78ms

118ms 387ms 320ms

65° 45° 35°

Cardiologist & Diabetologist MD, D.CARD, D. DIABETES DR RAVI CHAVAN 2004/06/2468

SUBURBAN DIAGNOSTICS

Patient Details Date: 23-Mar-22 Time: 12:25:57 PM

Name: JYOTI PRAKASH MALLIK ID: 2208211070

Age: 38 y Sex: M Height: 159 cms. Weight: 64 Kg.

Clinical History: NONE

Medications: NONE

Test Details

Protocol: Bruce Pr.MHR: 182 bpm THR: 154 (85 % of Pr.MHR) bpm

Total Exec. Time: 6 m 40 s Max. HR: 186 (102% of Pr. MHR)bpm Max. Mets: 10.20

Test Termination Criteria: Target HR attained

Protocol Details

Stage Name	Stage Time	Mets	Speed	Grade	Heart	Max. BP	Max. ST	Max. ST
	(min : sec)		(mph)	(%)	Rate (bpm)	(mm/Hg)	Level (mm)	Slope (mV/s)
Supine	0 : 15	1.0	0	0	95	120 / 80	-1.06 III	1.42 V2
Standing	0 : 15	1.0	0	0	101	120 / 80	-1.49 III	1.42 aVL
Hyperventilation	0:47	1.0	0	0	116	120 / 80	-5.94 V5	-4.25 V4
1	3:0	4.6	1.7	10	149	130 / 80	-2.55 III	2.83 V2
2	3:0	7.0	2.5	12	171	140 / 80	-2.97 III	3.89 V2
Peak Ex	0:40	10.2	3.4	14	186	180 / 80	-2.97 II	3.89 V2
Recovery(1)	1:0	1.8	1	0	151	160 / 80	-2.97 III	5.66 V2
Recovery(2)	1:0	1.0	0	0	136	160 / 80	-2.34 III	5.66 V2
Recovery(3)	1:0	1.0	0	0	122	140 / 80	-2.34 III	4.95 V2
Recovery(4)	1:0	1.0	0	0	115	140 / 80	-1.91 III	3.18 V2
Recovery(5)	0:29	1.0	0	0	118	120 / 80	-1. <mark>27 III</mark>	1.77 V2

Interpretation

GOOD EFFORT TOLERANCE ACCELERATED CHRONOTROPIC RESPONSE NORMAL INOTROPIC RESPONSE NO ANGINA/ ANGINA EQUIVALENTS

NO ARRHYTHMIAS

BASELINE ST-T CHANGES ABNORMALITIES IN INFEROLATERAL LEADS WHICH PERSISTED THROUGHOUT EXERCISE & REÇOVERY.

IMPRESSION:STRESS TEST IS EQUIVOCAL FOR STRESS INDUCIBLE ISCHAEMIA

Disclaimer: Negative stress test does not rule out Coronary Artery Disease.

Positive stress test is suggestive but not confirmatory of Coronary Artery Disease.

Hence clinical correlation is mandatory.

Ref. Doctor: MEDI ASSIST HEALTHCARE (Summary Report edited by user)

Doctor: DR. RAVI CHAVAN

(c) Schiller Healthcare India Pvt. Ltd. V 4.7

Dr. Ravi Chavan

sultani Zardiologist

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