

Name : MR.SHRIKANT SATAM

Age / Gender : 43 Years / Male

Consulting Dr. : -

Reg. Location : Borivali West (Main Centre)



R

E

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METHOD

MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO

CBC (Complete	Blood Count), Blood
<u>RESULTS</u>	BIOLOGICAL REF RANGE

· · · · · · · · · · · · · · · · · · ·	· · · · · · · · · · · · · · · · · · ·	•	
RBC PARAMETERS			
Haemoglobin	14.0	13.0-17.0 g/dL	Spectrophotometric
RBC	5.04	4.5-5.5 mil/cmm	Elect. Impedance
PCV	40.9	40-50 %	Measured
MCV	81	80-100 fl	Calculated
MCH	27.8	27-32 pg	Calculated
MCHC	34.2	31.5-34.5 g/dL	Calculated
RDW	14.3	11.6-14.0 %	Calculated

WBC PARAMETERS

PARAMETER

WBC Total Count 6330 4000-10000 / cmm Elect. Impedance

WBC DIFFERENTIAL AND ABSOLUTE COUNTS

WEC DILLERING AND I	ADSOLUTE COUNTS		
Lymphocytes	38.8	20-40 %	
Absolute Lymphocytes	2456.0	1000-3000 /cmm	Calculated
Monocytes	8.0	2-10 %	
Absolute Monocytes	506.4	200-1000 /cmm	Calculated
Neutrophils	50.3	40-80 %	
Absolute Neutrophils	3184.0	2000-7000 /cmm	Calculated
Eosinophils	2.2	1-6 %	
Absolute Eosinophils	139.3	20-500 /cmm	Calculated
Basophils	0.7	0.1-2 %	
Absolute Basophils	44.3	20-100 /cmm	Calculated

Immature Leukocytes -

WBC Differential Count by Absorbance & Impedance method/Microscopy.

PLATELET PARAMETERS

Platelet Count	294000	150000-400000 /cmm	Elect. Impedance
MPV	7.6	6-11 fl	Calculated
PDW	12.3	11-18 %	Calculated

RBC MORPHOLOGY

Hypochromia -Microcytosis -

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Name : MR.SHRIKANT SATAM

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Macrocytosis

Anisocytosis

Poikilocytosis

Polychromasia

Target Cells

Basophilic Stippling

Normoblasts

Others Normocytic, Normochromic

WBC MORPHOLOGY

PLATELET MORPHOLOGY

COMMENT

Specimen: EDTA Whole Blood

ESR, EDTA WB-ESR 8 2-15 mm at 1 hr. Sedimentation

Clinical Significance: The erythrocyte sedimentation rate (ESR), also called a sedimentation rate is the rate red blood cells sediment in a period of time.

Interpretation:

Factors that increase ESR: Old age, Pregnancy, Anemia

Factors that decrease ESR: Extreme leukocytosis, Polycythemia, Red cell abnormalities- Sickle cell disease

Limitations:

- It is a non-specific measure of inflammation.
- The use of the ESR as a screening test in asymptomatic persons is limited by its low sensitivity and specificity.

Reflex Test: C-Reactive Protein (CRP) is the recommended test in acute inflammatory conditions.

Reference:

- Brigden ML. Clinical utility of the erythrocyte sedimentation rate. American family physician. 1999 Oct 1;60(5):1443-50.

*Sample processed at SUBURBAN DIAGNOSTICS (INDIA) PVT. LTD Borivali Lab, Borivali West *** End Of Report ***





BMhaskar Dr.KETAKI MHASKAR M.D. (PATH) **Pathologist**

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:01-Apr-2024 / 16:12

Hexokinase

MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO

PARAMETER RESULTS BIOLOGICAL REF RANGE METHOD

GLUCOSE (SUGAR) FASTING, 113.5 Non-Diabetic: < 100 mg/dl Impaired Fasting Glucose:

100-125 mg/dl

Diabetic: >/= 126 mg/dl

Collected

Reported

GLUCOSE (SUGAR) PP, Fluoride 130.0 Non-Diabetic: < 140 mg/dl Hexokinase

Plasma PP/R Impaired Glucose Tolerance:

140-199 mg/dl

Diabetic: >/= 200 mg/dl

Urine Sugar (Fasting)AbsentAbsentUrine Ketones (Fasting)AbsentAbsent

Urine Sugar (PP) Absent Absent
Urine Ketones (PP) Absent Absent

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Kidney failure: <15

:01-Apr-2024 / 08:27 :01-Apr-2024 / 11:34

MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO KIDNEY FUNCTION TESTS

<u>PARAMETER</u>	<u>RESULTS</u>	BIOLOGICAL REF RANGE	<u>METHOD</u>
BLOOD UREA, Serum	15.4	12.8-42.8 mg/dl	Kinetic
BUN, Serum	7.2	6-20 mg/dl	Calculated
CREATININE, Serum	1.05	0.67-1.17 mg/dl	Enzymatic
eGFR, Serum	90	(ml/min/1.73sqm) Normal or High: Above 90 Mild decrease: 60-89 Mild to moderate decrease: 45- 59 Moderate to severe decrease:30 -44 Severe decrease: 15-29	Calculated
		Severe decrease: 15-29	

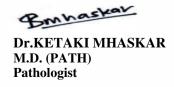
Note: eGFR estimation is calculated using 2021 CKD-EPI GFR equation w.e.f 16-08-2023

TOTAL PROTEINS, Serum	7.4	6.4-8.3 g/dL	Biuret
ALBUMIN, Serum	4.2	3.5-5.2 g/dL	BCG
GLOBULIN, Serum	3.2	2.3-3.5 g/dL	Calculated
A/G RATIO, Serum	1.3	1 - 2	Calculated
URIC ACID, Serum	6.2	3.5-7.2 mg/dl	Enzymatic
PHOSPHORUS, Serum	3.1	2.7-4.5 mg/dl	Molybdate UV
CALCIUM, Serum	9.4	8.6-10.0 mg/dl	N-BAPTA
SODIUM, Serum	139	135-148 mmol/l	ISE
POTASSIUM, Serum	4.8	3.5-5.3 mmol/l	ISE
CHLORIDE, Serum	105	98-107 mmol/l	ISE

^{*}Sample processed at SUBURBAN DIAGNOSTICS (INDIA) PVT. LTD Borivali Lab, Borivali West
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MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO **GLYCOSYLATED HEMOGLOBIN (HbA1c)**

BIOLOGICAL REF RANGE PARAMETER RESULTS METHOD

Glycosylated Hemoglobin 6.2 (HbA1c), EDTA WB - CC

Prediabetic Level: 5.7-6.4 %

Non-Diabetic Level: < 5.7 % Diabetic Level: >/= 6.5 %

Collected

Estimated Average Glucose (eAG), EDTA WB - CC

131.2

mg/dl

Calculated

HPLC

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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BMhaskar Dr.KETAKI MHASKAR M.D. (PATH) **Pathologist**

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CLIA

MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO **PROSTATE SPECIFIC ANTIGEN (PSA)**

PARAMETER RESULTS

Kindly note change in platform w.e.f. 24-01-2024

BIOLOGICAL REF RANGE METHOD

TOTAL PSA, Serum 0.731 <4.0 ng/ml

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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.

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

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-α 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.
- Patients who have been regularly exposed to animals or have received immunotherapy or diagnostic procedures utilizing immunoglobulins or immunoglobulin fragments may produce antibodies, e.g. HAMA, that interferes with immunoassays.
- PSA results should be interpreted in light of the total clinical presentation of the patient, including: symptoms, clinical history, data from additional tests, and other appropriate information.
- Serum PSA concentrations should not be interpreted as absolute evidence for the presence or absence of prostate cancer.

Note: The concentration of PSA in a given specimen, determined with assay from different manufacturers, may not be comparable due to differences in assay methods and reagent specificity.

Reference:

- Wallach's Interpretation of diagnostic tests
- Total PSA Pack insert

*Sample processed at SUBURBAN DIAGNOSTICS (INDIA) PVT. LTD SDRL, Vidyavihar Lab ** End Of Report *





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

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MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO URINE EXAMINATION REPORT

<u>PARAMETER</u>	<u>RESULTS</u>	BIOLOGICAL REF RANGE	<u>METHOD</u>
PHYSICAL EXAMINATION			
Color	Pale yellow	Pale Yellow	-
Reaction (pH)	5.0	4.5 - 8.0	Chemical Indicator
Specific Gravity	1.005	1.001-1.030	Chemical Indicator
Transparency	Clear	Clear	-
Volume (ml)	40	-	-
CHEMICAL EXAMINATION			
Proteins	Absent	Absent	pH Indicator
Glucose	Absent	Absent	GOD-POD
Ketones	Absent	Absent	Legals Test
Blood	Absent	Absent	Peroxidase
Bilirubin	Absent	Absent	Diazonium Salt
Urobilinogen	Normal	Normal	Diazonium Salt
Nitrite	Absent	Absent	Griess Test
MICROSCOPIC EXAMINATION	<u>on</u>		
Leukocytes(Pus cells)/hpf	1-2	0-5/hpf	
Red Blood Cells / hpf	Absent	0-2/hpf	
Epithelial Cells / hpf	0-1		
Casts	Absent	Absent	
Crystals	Absent	Absent	
Amorphous debris	Absent	Absent	
Bacteria / hpf	2-3	Less than 20/hpf	
Others	-		

Interpretation: The concentration values of Chemical analytes corresponding to the grading given in the report are as follows:

- Protein (1+ = 25 mg/dl , 2+ = 75 mg/dl , 3+ = 150 mg/dl , 4+ = 500 mg/dl)
- Glucose(1+ = 50 mg/dl, 2+ =100 mg/dl, 3+ =300 mg/dl, 4+ =1000 mg/dl)
- Ketone (1+ = 5 mg/dl, 2+ = 15 mg/dl, 3+ = 50 mg/dl, 4+ = 150 mg/dl)

Reference: Pack inert

*Sample processed at SUBURBAN DIAGNOSTICS (INDIA) PVT. LTD Borivali Lab, Borivali West *** End Of Report **





BMhaskar Dr.KETAKI MHASKAR M.D. (PATH) **Pathologist**

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MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO **BLOOD GROUPING & Rh TYPING**

RESULTS PARAMETER

ABO GROUP В

Rh TYPING Positive

NOTE: Test performed by automated Erythrocytes magnetized technology (EMT) 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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Dr.VRUSHALI SHROFF M.D.(PATH) **Pathologist**

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MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO LIPID PROFILE

CHOLESTEROL, Serum 225.0 Desirable: <200 mg/dl Borderline High: 200-239mg/dl High: >/=240 mg/dl Normal: <150 mg/dl Borderline-high: 150 - 199 mg/dl High: 200 - 499 mg/dl Homogeneous enzymatic colorimetric assay NON HDL CHOLESTEROL, Serum NON HDL CHOLESTEROL, Serum Perinable: <130 mg/dl Homogeneous enzymatic colorimetric assay Calculated NON HDL CHOLESTEROL, Serum Perinable: <130 mg/dl Borderline-high: 130 - 159 mg/dl High: 160 - 189 mg/dl Very high: >/=190 mg/dl NON HDL CHOLESTEROL, Serum Perinable: <200 mg/dl Homogeneous enzymatic colorimetric assay Calculated Calculated Calculated Very high: >/=190 mg/dl Very high: 130 - 159 mg/dl Borderline High: 130 - 159 mg/dl High: 160 - 189 mg/dl Very high: >/= 190 mg/dl Very hig	<u>PARAMETER</u>	<u>RESULTS</u>	BIOLOGICAL REF RANGE	<u>METHOD</u>
Borderline-high: 150 - 199 mg/dl High: 200 - 499 mg/dl Very high:>/=500 mg/dl HDL CHOLESTEROL, Serum 32.9 Desirable: >60 mg/dl Low (High risk): <40 mg/dl Enzymatic Colorimetric assay NON HDL CHOLESTEROL, Serum 192.1 Desirable: <130 mg/dl Borderline-high: 130 - 159 mg/dl High: 160 - 189 mg/dl Very high: >/=190 mg/dl Very high: >/=190 mg/dl Desirable: <130 mg/dl Calculated	CHOLESTEROL, Serum	225.0	Borderline High: 200-239mg/dl	CHOD-POD
Borderline: 40 - 60 mg/dl colorimetric assay NON HDL CHOLESTEROL, 192.1 Serum Desirable: <130 mg/dl Borderline-high:130 - 159 mg/dl High:160 - 189 mg/dl Very high: >/=190 mg/dl Very high: >/=190 mg/dl Borderline High: 130 - 129 mg/dl Borderline High: 130 - 129 mg/dl Borderline High: 130 - 159 mg/dl Borderline High: 130 - 159 mg/dl Borderline High: 130 - 159 mg/dl High: 160 - 189 mg/dl High: 160 - 189 mg/dl Very High: >/= 190 mg/dl Calculated VLDL CHOLESTEROL, Serum 68.9 VLDL CHOLESTEROL, Serum 68.9 CHOL / HDL CHOL RATIO, 6.8 O-4.5 Ratio Calculated	TRIGLYCERIDES, Serum	471.0	Borderline-high: 150 - 199 mg/dl High: 200 - 499 mg/dl	GPO-POD
Serum Borderline-high:130 - 159 mg/dl High:160 - 189 mg/dl Very high: >/=190 mg/dl Very high: >/=190 mg/dl Calculated Near Optimal: 100 - 129 mg/dl Borderline High: 130 - 159 mg/dl High: 160 - 189 mg/dl Very High: >/= 190 mg/dl Very High: >/= 190 mg/dl Very High: >/= 190 mg/dl VLDL CHOLESTEROL, Serum CHOL / HDL CHOL RATIO, 6.8 Serum LDL CHOL / HDL CHOL RATIO, 3.7 O-3.5 Ratio Calculated	HDL CHOLESTEROL, Serum	32.9	Borderline: 40 - 60 mg/dl	enzymatic
Near Optimal: 100 - 129 mg/dl Borderline High: 130 - 159 mg/dl High: 160 - 189 mg/dl Very High: >/= 190 mg/dl Very High: >/= 190 mg/dl Calculated CHOL / HDL CHOL RATIO, Serum LDL CHOL / HDL CHOL RATIO, 3.7 O-3.5 Ratio Calculated	•	192.1	Borderline-high:130 - 159 mg/dl High:160 - 189 mg/dl	
CHOL / HDL CHOL RATIO, 6.8 0-4.5 Ratio Calculated Serum LDL CHOL / HDL CHOL RATIO, 3.7 0-3.5 Ratio Calculated	LDL CHOLESTEROL, Serum	123.2	Near Optimal: 100 - 129 mg/dl Borderline High: 130 - 159 mg/dl High: 160 - 189 mg/dl	Calculated
Serum LDL CHOL / HDL CHOL RATIO, 3.7 0-3.5 Ratio Calculated	VLDL CHOLESTEROL, Serum	68.9	< /= 30 mg/dl	Calculated
		6.8	0-4.5 Ratio	Calculated
		3.7	0-3.5 Ratio	Calculated

Note: LDL test is performed by direct measurement.

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MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO THYROID FUNCTION TESTS

<u>PARAMETER</u>	<u>RESULTS</u>	BIOLOGICAL REF RANGE	<u>METHOD</u>
Free T3, Serum	5.5	3.5-6.5 pmol/L	ECLIA
Free T4, Serum	10.7	11.5-22.7 pmol/L	ECLIA
sensitiveTSH, Serum	20.8	0.35-5.5 microIU/ml	ECLIA



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Interpretation:

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

Clinical Significance:

- 1)TSH Values between high abnormal upto15 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 epileptics.

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:

- 1. 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.
- 2. Patient samples may contain heterophilic antibodies that could react in immunoassays to give falsely elevated or depressed results. this assay is designed to minimize interference from heterophilic antibodies.

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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Dr.JYOT THAKKER
M.D. (PATH), DPB
Pathologist & AVP(Medical Services)

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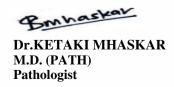
MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO LIVER FUNCTION TESTS

<u>PARAMETER</u>	<u>RESULTS</u>	BIOLOGICAL REF RANGE	<u>METHOD</u>
BILIRUBIN (TOTAL), Serum	0.56	0.1-1.2 mg/dl	Colorimetric
BILIRUBIN (DIRECT), Serum	0.28	0-0.3 mg/dl	Diazo
BILIRUBIN (INDIRECT), Serum	0.28	0.1-1.0 mg/dl	Calculated
TOTAL PROTEINS, Serum	7.4	6.4-8.3 g/dL	Biuret
ALBUMIN, Serum	4.2	3.5-5.2 g/dL	BCG
GLOBULIN, Serum	3.2	2.3-3.5 g/dL	Calculated
A/G RATIO, Serum	1.3	1 - 2	Calculated
SGOT (AST), Serum	20.8	5-40 U/L	NADH (w/o P-5-P)
SGPT (ALT), Serum	25.0	5-45 U/L	NADH (w/o P-5-P)
GAMMA GT, Serum	21.5	3-60 U/L	Enzymatic
ALKALINE PHOSPHATASE, Serum	84.9	40-130 U/L	Colorimetric

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*** End Of Report ***







SUBURBAN DIAGNOSTICS - BORIVALI WEST

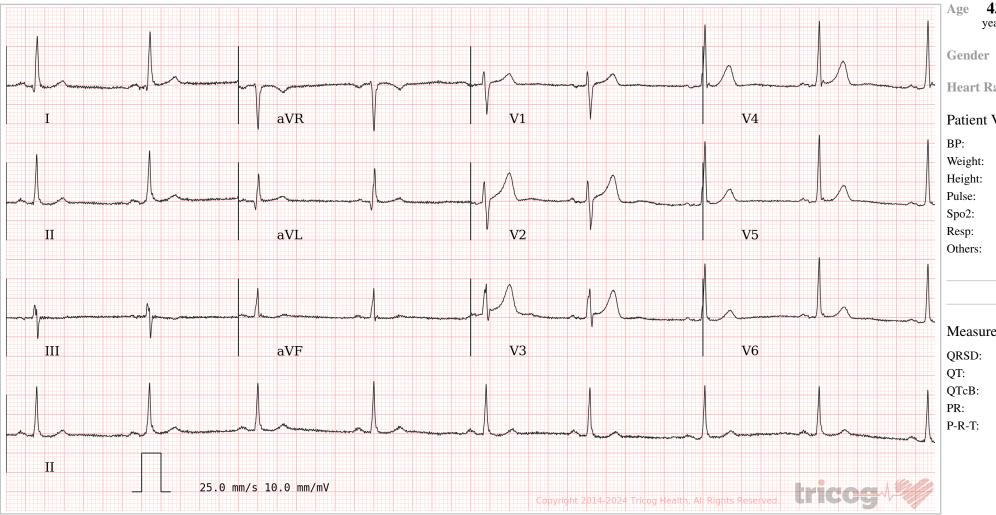


Patient Name: SHRIKANT SATAM

Patient ID:

2409200295

Date and Time: 1st Apr 24 9:08 AM



years months days

Gender Male

Heart Rate 52bpm

Patient Vitals

NA NA NA NA NA NA

Measurements

82ms 378ms 351ms 176ms 36° 24° 3°

Sinus Bradycardia. Please correlate clinically.

REPORTED BY

Dr Nitin Sonavane M.B.B.S.AFLH, D.DIAB, D.CARD Consultant Cardiologist 87714

Disclaimer: 1) Analysis in this report is based on ECG alone and should be used as an adjunct to clinical history, symptoms, and results of other invasive and non-invasive tests and must be interpreted by a qualified physician. 2) Patient vitals are as entered by the clinician and not derived from the ECG.



PCID# TESTING . HEA 2409200295

Name

: MR.SHRIKANT SATAM

Age / Gender : 43 Years/Male

Consulting Dr. :

Reg.Location : Borivali West (Main Centre)

Collected

: 01-Apr-2024 / 08:23

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Reported

: 01-Apr-2024 / 16:46

PHYSICAL EXAMINATION REPORT

History and Complaints:

Nil

EXAMINATION FINDINGS:

Height (cms):

166

Weight (kg):

79

Temp (0c):

Afebrile

Skin:

NAD

Blood Pressure (mm/hg): 140/80

Nails:

NAD

Pulse:

76/min

Lymph Node:

Not Palpable

Systems

Cardiovascular: S1S2-Normal Chest-Clear

Respiratory: **Genitourinary:**

NAD

GI System:

NAD

CNS:

NAD

IMPRESSION: 🔈

ADVICE:

Lipid prehie TSH physician Refu

CHIEF COMPLAINTS:

1) Hypertension:

No

2) IHD

No

3) Arrhythmia

No

4) Diabetes Mellitus

No

5) Tuberculosis

No

6) Asthama

No

7) Pulmonary Disease

No



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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	No
2)	Smoking	No
3)	Diet	Veg/Mix
4)	Medication	No

*** End Of Report ***

Dr.NITIN SONAVANE PHYSICIAN

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> DR. NITEM SONAVA NE M.B.B.S. AFLH, O DIAB, G 47-80. CONSULTANT-CARDIOLOGIST RESD. NO.: 87714



R E 0 R T

Date:-

Name:-

Shrikant Satemsex/Age: 43 m

EYE CHECK UP

Chief complaints:

Systemic Diseases:

Past history:

Unaided Vision:

Aided Vision:

Refraction:

NO

MILO

MILO

(Left Eye)

(, - ,		
T .	T	

	Sph	Cyl	Axis	Vn	Sph	Cyl	Axis	Vn
Distance					- , Hills			
Near			n hebr					

Colour Vision: Normal / Abnormal

(Right Eve)

Remark:

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CID NO: 2409200295	
PATIENT'S NAME: MR.SHRIKANT SATAM	AGE/SEX: 43 Y/M
REF BY:	DATE: 01/04/2024

2-D ECHOCARDIOGRAPHY

- 1. RA, LA RV is Normal Size.
- 2. No LV Hypertrophy.
- 3. Normal LV systolic function. LVEF 60 % by bi-plane
- 4. No RWMA at rest.
- 5. Aortic, Pulmonary, Tricuspid valves normal, Trivial MR.
- 6. Great arteries: Aorta: Normal
 - a. No mitral valve prolaps.
- 7. Inter-ventricular septum is intact and normal.
- 8. Intra Atrial Septum intact.
- 9. Pulmonary vein, IVC, hepatic are normal.
- 10.No LV clot.
- 11. No Pericardial Effusion
- 12. No Diastolic disfunction. No Doppler evidence of raised LVEDP.



PATIENT'S NAME: MR.SHRIKANT SATAM

REF BY: ----
DATE: 01/04/2024

1.	AO root diameter	2.7 cm
2.	IVSd	1.1 cm
3.	LVIDd	4.3 cm
4.	LVIDs	2.3 cm
5.	LVPWd	1.1 cm
6.	LA dimension	3.7 cm
7.	RA dimension	3.7 cm
8.	RV dimension	3.0 cm
9.	Pulmonary flow vel:	0.8 m/s
10.	Pulmonary Gradient	3.2 m/s
11.	. Tricuspid flow vel	1.4 m/s
12	. Tricuspid Gradient	8 m/s
13	. PASP by TR Jet	18 mm Hg
14	. TAPSE	3.0 cm
15	. Aortic flow vel	1.1 m/s
16	. Aortic Gradient	5 m/s
17	. MV:E	0.8 m/s
18	. A vel	0.6 m/s
19	. IVC	17 mm
20). E/E'	10

Impression:

Normal 2d echo study.

Disclaimer

Echo may have inter/Intra observer variations in measurements as the study is observer dependent and changes with Pt's hemodynamics. Please co-relate findings with patients clinical status.

End of Report

DR. S. NITIN Consultant Cardiologist Reg. No. 87714 R

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Name : Mr SHRIKANT SATAM

Age / Sex : 43 Years/Male

Ref. Dr :

Reg. Location: Borivali West



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USG WHOLE ABDOMEN

<u>LIVER:</u>Liver is normal in size 14.9 cm, with mild generalized increase in parenchymal echotexture. There is no intra-hepatic biliary radical dilatation. No evidence of any focal lesion.

GALL BLADDER: Gall bladder is distended and appears normal. No obvious wall thickening is noted. There is no evidence of any calculus.

(Tiny polyps/calculi may be missed due to technical limitations, sub-optimal distension of GB, adjacent gases and inter-machine variability in resolution settings)

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

PANCREAS: Pancreas appears normal in echotexture. There is no evidence of any focal lesion or calcification.

KIDNEYS: Right kidney measures 9.7 x 3.8 cm. Left kidney measures 10.5 x 5.2 cm.

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, shape and echotexture. No focal lesion is seen.

URINARY BLADDER: Urinary bladder is distended and normal. Wall thickness is within normal limits.

PROSTATE: Prostate is normal in size and echotexture. Prostate measures 3.4 x 2.7 x 3.4 cm and prostatic weight is 18 gm. No evidence of any obvious focal lesion.

No free fluid or size significant lymphadenopathy is seen.



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Opinion:

Grade I fatty infiltration of liver.

For clinical correlation and follow up.

Note: Investigations have their limitations. Solitary radiological investigations never confirm the final diagnosis. They only help in diagnosing the disease in correlation to clinical symptoms and other related tests. USG is known to have inter-observer variations. Further / Follow-up imaging may be needed in some cases for confirmation / exclusion of diagnosis. Patient was explained in detail verbally about the USG findings, USG measurements and its limitations. In case of any typographical error in the report, patient is requested to immediately contact the center for rectification within 7 days post which the center will not be responsible for any rectification. Please interpret accordingly.

-----End of Report-----

DR.SUDHANSHU SAXENA Consultant Radiologist M.B.B.S DMRE (RadioDiagnosis) RegNo .MMC 2016061376.



Name : Mr SHRIKANT SATAM

Age / Sex : 43 Years/Male

Ref. Dr

Reg. Location: Borivali West



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X-RAY CHEST PA VIEW

Both lung fields are clear.

Both costo-phrenic angles are clear.

The cardiac size and shape are within normal limits.

The domes of diaphragm are normal in position and outlines.

The skeleton under review appears normal.

IMPRESSION:

NO SIGNIFICANT ABNORMALITY IS DETECTED.

Kindly correlate clinically.

Note: Investigations have their limitations. Solitary radiological investigations never confirm the final diagnosis. They only help in diagnosing the disease in correlation to clinical symptoms and other related tests. X ray is known to have inter-observer variations. Further / follow-up imaging may be needed in some cases for confirmation / exclusion of diagnosis. Please interpret accordingly. In case of any typographical error / spelling error in the report, patient is requested to immediately contact the centre within 7 days post which the center will not be responsible for any rectification.

-----End of Report-----

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