

Name : MR.MURUGESH NAINAR

Age / Gender : 43 Years / Male

Consulting Dr. : -

Reg. Location

: Bhayander East (Main Centre)

Authenticity Check

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Reported

:29-Mar-2024 / 09:20 :29-Mar-2024 / 12:23

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

| CBC (Complete Blood Count), Blood | | | |
|-----------------------------------|----------------|-----------------------------|--------------------|
| <u>PARAMETER</u> | <u>RESULTS</u> | BIOLOGICAL REF RANGE | <u>METHOD</u> |
| RBC PARAMETERS | | | |
| Haemoglobin | 13.5 | 13.0-17.0 g/dL | Spectrophotometric |
| RBC | 4.74 | 4.5-5.5 mil/cmm | Elect. Impedance |
| PCV | 39.5 | 40-50 % | Measured |
| MCV | 83 | 80-100 fl | Calculated |
| MCH | 28.5 | 27-32 pg | Calculated |
| MCHC | 34.2 | 31.5-34.5 g/dL | Calculated |
| RDW | 13.9 | 11.6-14.0 % | Calculated |
| WBC PARAMETERS | | | |
| WBC Total Count | 6990 | 4000-10000 /cmm | Elect. Impedance |
| WBC DIFFERENTIAL AND ABSO | LUTE COUNTS | | |
| Lymphocytes | 31.4 | 20-40 % | |
| Absolute Lymphocytes | 2194.9 | 1000-3000 /cmm | Calculated |
| Monocytes | 10.2 | 2-10 % | |
| Absolute Monocytes | 713.0 | 200-1000 /cmm | Calculated |
| Neutrophils | 40.9 | 40-80 % | |
| Absolute Neutrophils | 2858.9 | 2000-7000 /cmm | Calculated |
| Eosinophils | 16.5 | 1-6 % | |
| Absolute Eosinophils | 1153.3 | 20-500 /cmm | Calculated |
| Basophils | 1.0 | 0.1-2 % | |
| Absolute Basophils | 69.9 | 20-100 /cmm | Calculated |

WBC Differential Count by Absorbance & Impedance method/Microscopy.

PLATELET PARAMETERS

| Platelet Count | 304000 | 150000-400000 /cmm | Elect. Impedance |
|----------------|--------|--------------------|------------------|
| MPV | 6.9 | 6-11 fl | Calculated |
| PDW | 9.8 | 11-18 % | Calculated |

RBC MORPHOLOGY

Immature Leukocytes

Hypochromia -Microcytosis -

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:29-Mar-2024 / 12:50

Macrocytosis

Anisocytosis

Poikilocytosis

Polychromasia

Target Cells

Basophilic Stippling

Normoblasts

Others Normocytic, Normochromic

WBC MORPHOLOGY PLATELET MORPHOLOGY

COMMENT Eosinophilia

Advice:1)Stool examination for parasites

2) Allergy testing

Specimen: EDTA Whole Blood

ESR, EDTA WB-ESR 2-15 mm at 1 hr. Sedimentation 6



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:29-Mar-2024 / 12:23

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:

- Pack Insert
- 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
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: 29-Mar-2024 / 13:29

:29-Mar-2024 / 17:43

Hexokinase

Hexokinase

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

<u>PARAMETER</u> <u>RESULTS</u> <u>BIOLOGICAL REF RANGE</u> <u>METHOD</u>

GLUCOSE (SUGAR) FASTING, 119.0 Non-Diabetic: < 100 mg/dl Fluoride Plasma Impaired Fasting Glucose:

100-125 mg/dl

Diabetic: >/= 126 mg/dl

Collected

Reported

GLUCOSE (SUGAR) PP, Fluoride 175.0 Non-Diabetic: < 140 mg/dl

Plasma PP/R Impaired Glucose Tolerance:

140-199 mg/dl

Diabetic: >/= 200 mg/dl

Urine Sugar (Fasting)AbsentAbsentUrine Ketones (Fasting)AbsentAbsent

Urine Sugar (PP)AbsentAbsentUrine Ketones (PP)AbsentAbsent

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Kidney failure: <15

: 29-Mar-2024 / 09:20 : 29-Mar-2024 / 16:04

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 | 26.7 | 12.8-42.8 mg/dl | Kinetic |
| BUN, Serum | 12.5 | 6-20 mg/dl | Calculated |
| CREATININE, Serum | 1.19 | 0.67-1.17 mg/dl | Enzymatic |
| eGFR, Serum | 78 | (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 |

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

| TOTAL PROTEINS, Serum | 7.2 | 6.4-8.3 g/dL | Biuret |
|-----------------------|-----|----------------|--------------|
| ALBUMIN, Serum | 4.3 | 3.5-5.2 g/dL | BCG |
| GLOBULIN, Serum | 2.9 | 2.3-3.5 g/dL | Calculated |
| A/G RATIO, Serum | 1.5 | 1 - 2 | Calculated |
| URIC ACID, Serum | 5.3 | 3.5-7.2 mg/dl | Enzymatic |
| PHOSPHORUS, Serum | 3.5 | 2.7-4.5 mg/dl | Molybdate UV |
| CALCIUM, Serum | 9.8 | 8.6-10.0 mg/dl | N-BAPTA |
| SODIUM, Serum | 138 | 135-148 mmol/l | ISE |
| POTASSIUM, Serum | 5.1 | 3.5-5.3 mmol/l | ISE |
| CHLORIDE, Serum | 101 | 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)

PARAMETER RESULTS BIOLOGICAL REF RANGE METHOD

Glycosylated Hemoglobin (HbA1c), EDTA WB - CC

6.9 Non-Diabetic Level: < 5.7 %

Prediabetic Level: 5.7-6.4 % Diabetic Level: >/= 6.5 %

Collected

Reported

Estimated Average Glucose (eAG), EDTA WB - CC

151.3

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.

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





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

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Name : MR.MURUGESH NAINAR

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TOTAL PSA, Serum

: -

Reg. Location: Bhayander East (Main Centre)



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

<4.0 ng/ml

PARAMETER RESULTS BIOLOGICAL REF RANGE METHOD

1.361

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



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Consulting Dr. : - Collected : 29-Mar-2024 / 09:20

Reg. Location : Bhayander East (Main Centre) Reported :29-Mar-2024 / 15:45

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-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.NAMRATA RAUL M.D (Biochem) Biochemist

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Collected

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

PARAMETER RESULTS

ABO GROUP AB

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

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





Dr.VRUSHALI SHROFF M.D.(PATH) Pathologist

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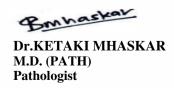
MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO LIPID PROFILE

| <u>PARAMETER</u> | <u>RESULTS</u> | BIOLOGICAL REF RANGE | <u>METHOD</u> |
|-------------------------------------|----------------|--|--|
| CHOLESTEROL, Serum | 189.0 | Desirable: <200 mg/dl Borderline High: 200-239mg/dl High: >/=240 mg/dl | CHOD-POD |
| TRIGLYCERIDES, Serum | 105.0 | Normal: <150 mg/dl Borderline-high: 150 - 199 mg/dl High: 200 - 499 mg/dl Very high:>/=500 mg/dl | GPO-POD |
| HDL CHOLESTEROL, Serum | 45.5 | Desirable: >60 mg/dl Borderline: 40 - 60 mg/dl Low (High risk): <40 mg/dl | Homogeneous enzymatic colorimetric assay |
| NON HDL CHOLESTEROL, Serum | 143.5 | Desirable: <130 mg/dl Borderline-high:130 - 159 mg/dl High:160 - 189 mg/dl Very high: >/=190 mg/dl | Calculated |
| LDL CHOLESTEROL, Serum | 123.0 | 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 | 20.5 | < /= 30 mg/dl | Calculated |
| CHOL / HDL CHOL RATIO, Serum | 4.2 | 0-4.5 Ratio | Calculated |
| LDL CHOL / HDL CHOL RATIO, Serum | 2.7 | 0-3.5 Ratio | Calculated |

^{*}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 **THYROID FUNCTION TESTS**

| <u>PARAMETER</u> | <u>RESULTS</u> | BIOLOGICAL REF RANGE | <u>METHOD</u> |
|---------------------|----------------|----------------------|---------------|
| Free T3, Serum | 5.7 | 3.5-6.5 pmol/L | ECLIA |
| Free T4, Serum | 18.6 | 11.5-22.7 pmol/L | ECLIA |
| sensitiveTSH, Serum | 4.31 | 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 | | | 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. |
| | | 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 illness. | | 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)

*Sample processed at SUBURBAN DIAGNOSTICS (INDIA) PVT. LTD Borivali Lab, Borivali West
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Dr.KETAKI MHASKAR M.D. (PATH) Pathologist



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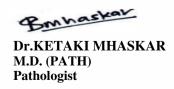
MEDIWHEEL FULL BODY HEALTH CHECKUP MALE ABOVE 40/2D ECHO LIVER FUNCTION TESTS

| <u>PARAMETER</u> | <u>RESULTS</u> | BIOLOGICAL REF RANGE | <u>METHOD</u> |
|--------------------------------|----------------|----------------------|------------------|
| BILIRUBIN (TOTAL), Serum | 0.52 | 0.1-1.2 mg/dl | Colorimetric |
| BILIRUBIN (DIRECT), Serum | 0.29 | 0-0.3 mg/dl | Diazo |
| BILIRUBIN (INDIRECT), Serum | 0.23 | 0.1-1.0 mg/dl | Calculated |
| TOTAL PROTEINS, Serum | 7.2 | 6.4-8.3 g/dL | Biuret |
| ALBUMIN, Serum | 4.3 | 3.5-5.2 g/dL | BCG |
| GLOBULIN, Serum | 2.9 | 2.3-3.5 g/dL | Calculated |
| A/G RATIO, Serum | 1.5 | 1 - 2 | Calculated |
| SGOT (AST), Serum | 28.5 | 5-40 U/L | NADH (w/o P-5-P) |
| SGPT (ALT), Serum | 18.7 | 5-45 U/L | NADH (w/o P-5-P) |
| GAMMA GT, Serum | 30.9 | 3-60 U/L | Enzymatic |
| ALKALINE PHOSPHATASE, Serum | 48.1 | 40-130 U/L | Colorimetric |

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







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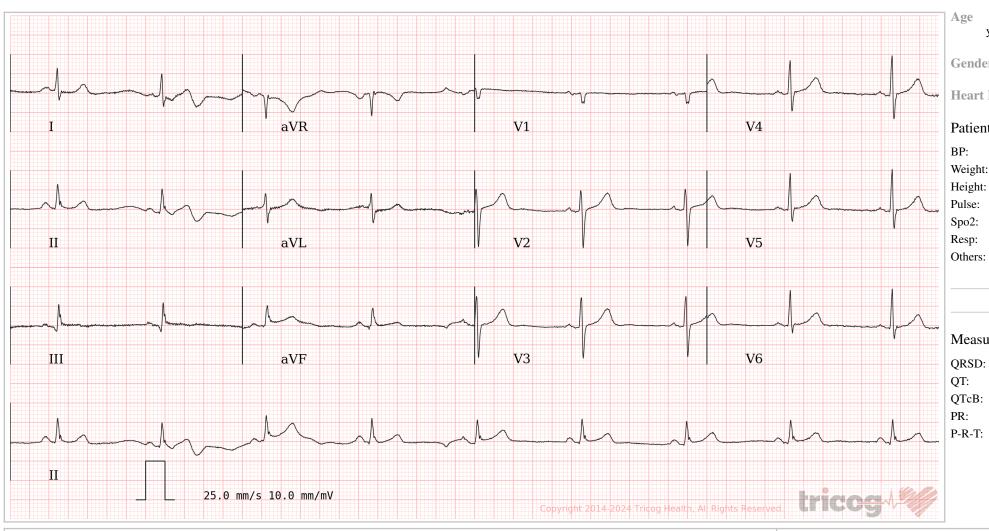
SUBURBAN DIAGNOSTICS - BHAYANDER EAST



Patient Name: MURUGESH NAINAR

Date and Time: 29th Mar 24 10:58 AM

Patient ID: 2408913101



months days

Gender Male

Heart Rate 56bpm

Patient Vitals

110/80 mmHg

74 kg Weight:

161 cm

NA NA

NA

Others:

Measurements

QRSD: 76ms 390ms QTcB: 376ms 134ms

44° 69° 39°

ECG Within Normal Limits: Sinus Bradycardia, Normal axis. No significant ST-T changes. Please correlate clinically.

REPORTED BY

Dr. Smita Valani MBBS, D. Cardiology 2011/03/0587

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.

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Name G N O S MR MURUGESH NAINAR

Age / Gender : 43 Years/Male

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: 29-Mar-2024 / 09:08

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: 29-Mar-2024 / 15:17

PHYSICAL EXAMINATION REPORT

History and Complaints:

No Complaint

EXAMINATION FINDINGS:

Height (cms):

161

Weight (kg):

74

Temp (0c):

Afebrile

Skin:

NAD

Blood Pressure (mm/hg): 110/80

Nails:

NAD

Pulse:

60 74/min

Lymph Node:

AB+me/

Not Palpable

Systems

Cardiovascular: S1S2-Normal

Respiratory:

Chest-Clear

Genitourinary: GI System:

NAD NAD

CNS:

NAD

IMPRESSION: HO AIC -

RESSION: HIS AIC - 6.9%.

US a in Sto. Sp. II factory Wine

ICE: CBC, Bio chemitry are NNL

Ly Emper- commetanta.

ADVICE:

CHIEF COMPLAINTS:

Hypertension:

No

IHD

No

Arrhythmia

No

4) Diabetes Mellitus

No No

Tuberculosis 6) Asthama

No

7) Pulmonary Disease

No

Name S NOS MR. MURUGESH NAINAR

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: 29-Mar-2024 / 15:17

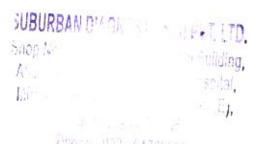
| | 8) | Thyroid/ Endocrine disorders | No |
|----|-----|--------------------------------------|----|
| | 9) | Nervous disorders | No |
| | 10) | GI system | No |
| | 11) | Genital urinary disorder | |
| | 12) | Rheumatic joint diseases or symptoms | No |
| | 13) | Blood disease or disorder | No |
| | 14) | Cancer/lump growth/cyst | No |
| | 15) | Congenital disease | No |
| 83 | 16) | Surgeries | No |
| 9 | 17) | Musculoskeletal System | No |
| | | | |

PERSONAL HISTORY:

| 1) | Alcohol | No |
|----|------------|-------|
| 2) | Smoking | No |
| 3) | Diet | Mixed |
| 4) | Medication | No |

*** End Of Report ***

CONSULTANT FRYSICIAN
Reg. No. 2017/12/3553



REGD. OFFICE: Dr. Lal PathLabs Ltd., Block E, Sector-18, Rohini, New Delhi - 110085. | CIN No.: L74899DL1995PLC065388



| PATIENT NAME :MR.MURUGESH NAINAR | SEX : MALE |
|----------------------------------|--------------------|
| REFERRED BY : DR | AGE : 43 YEARS |
| • CID NO : 2408913101 | • DATE: 29/03/2024 |

2D-Echocardigram & Doppler Report

Cardiac Evalution:

DIMENSIONS:

| | | 4 | |
|-------|------|-------|----|
| IVSd | | 11.0 | mm |
| IVSs | E 12 | 12.7 | mm |
| LVIDd | | 34.7 | mm |
| LVIDs | | 18.0 | mm |
| LVPWd | | 13.1 | mm |
| LVPWS | | 11.0 | mm |
| LVEF | | 55-60 | % |
| AO | | 29.8 | mm |
| LA | | 34.7 | mm |
| AVC | | 14.9 | mm |
| | | | |

MORPHOLOGICAL DATA

| Normal |
|--------|
| Normal |
| Normal |
| Normal |
| Normal |
| Intact |
| Normal |
| |



DOPPLER DATA:

| Mitral E velocity | 0.80 | cm/s | |
|-------------------|------|------|-------------|
| Mitral A velocity | 0.65 | cm/s | |
| Mitral E/A | 1.23 | | |
| AV max | 1.00 | cm/s | PG 4.0 mmhg |
| PV max | 0.89 | cm/s | PG 3.2 mmhg |
| TR max | 1.77 | cm/s | PG 25 mmhg |

IMPRESSION:

- Normal dimensions of all cardiac chambers.
- Good LV systolic Function. LVEF = 55-60 %.
- No RWMA.
- No clot/vegetation/effusion.
- No PH. (PASP by TR jet 25 mm Hg).

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

DR. SMITA VALANI M.B.B.S., D. Cardiology Reg. No. 2011/03/0587 CONSULTANT CARDIOLOGIST



DR. ANITA CHOUDWARY

CONSUL 2017/12/5553 Reg. No. 2017/12/5553

myrugesh Namar sex/Age: 43/m

EYE CHECK UP

Chief complaints:

Systemic Diseases:

Past history:

Unaided Vision:

Aided Vision:

Refraction:

RF (F 616 616 N/6 N/6

(Right Eye)

(Left Eye)

| | | | | (Leit Ey | e) | | | |
|----------|-----|-----|-------|----------|-----|-----|------|---|
| - 0 | Sph | Cyl | *Axis | Vn | Sph | Cyl | Axis | Vn |
| Distance | | | | | | | | • |
| Near | | | | | | | | |

Colour Vision: Normal / Abnormal

Remark:

SUBURBANTO CALLETTER (1) FVY. LTD.
Shop No. 10 Mira - Bhayar - La grayander (E),

Dist. Thane - 401 105.

Phone:: 022 - 61700000



Name : Mr MURUGESH NAINAR

Age / Sex : 43 Years/Male

Ref. Dr Reg. Date : 29-Mar-2024

: 29-Mar-2024/21:47 Reg. Location : Bhayander East Main Centre Reported



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USG WHOLE ABDOMEN

LIVER:

The liver is normal in size (13.2 cm), normal in shape and shows smooth margins. It shows raised parenchymal echotexture. No obvious cystic or solid lesion made out in the parenchyma. The intra hepatic biliary and portal radicals appear normal. The main portal vein appears normal. (PV-6.3 mm)

GALL BLADDER:

The gall bladder is folded and physiologically distended. Neck region is not well visualised. Gall bladder wall appears normal. No evidence of any calculus, mass lesion or sludge seen in the visualised lumen.

COMMON BILE DUCT:

The visualized common bile duct is normal in caliber. Terminal common bile duct is obscured due to bowel gas artefacts.

PANCREAS:

The pancreas appears normal. No evidence of solid or cystic mass lesion seen.

KIDNEYS:

Right kidney measures 9.8 x 4.7 cm. Left kidney measures 10.1 x 5.7 cm. Both the kidneys are normal in size, shape, position and echotexture. Corticomedullary differentiation is well maintained. Pelvicalyceal system is normal. A 3.9 mm calculus seen in the mid pole of left kidney. No evidence of any calculus seen in the right kidney. No evidence of any hydronephrosis or mass lesion seen on both sides.

SPLEEN:

The spleen is normal in size (10.0 cm) and echotexture. No evidence of focal lesion is noted.

URINARY BLADDER:

The urinary bladder is well distended and reveals no intraluminal abnormality. Bladder wall appears normal. No obvious calculus or mass lesion made out in the lumen.

PROSTATE:

The prostate is normal in size 3.4 x 2.6 x 2.5 cm and weighs 12.0 gms. It shows normal parenchymal echotexture. No obvious calcification or mass lesion made out.

There is no evidence of any lymphadenopathy or ascites.

Click here to view images http://3.111.232.119/iRISViewer/NeoradViewer?AccessionNo=2024032909091512



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IMPRESSION:

- Grade II fatty infiltration of liver.
- Non obstructive left renal calculus.

Kindly correlate clinically.

Investigations have their limitation. Solitary pathological/Radiological & other investigations never confirm the final diagnosis. They only help in diagnosing the disease in correlation to clinical symptoms & other related tests. Please interpret accordingly.

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

Dr. Aisha Lakhani Mbbs, Md (Radio Diagnosis)

Bhayander center



Name : Mr MURUGESH NAINAR

Age / Sex : 43 Years/Male

Ref. Dr

Reg. Location: Bhayander East Main Centre



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Name : Mr MURUGESH NAINAR

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X-RAY CHEST PA VIEW

Both the lung fields are clear with no active parenchymal lesion.

The cardiothoracic ratio is maintained and the cardiac outline is normal.

The domes of the diaphragm and hila are normal.

The cardio and costophrenic angles are clear.

Bony thorax is normal.

IMPRESSION:

• No obvious active parenchymal lesion made out.

Kindly correlate clinically.

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

Dr. Aisha Lakhani Mbbs, Md (Radio Diagnosis)

Bhayander center



: Mr MURUGESH NAINAR Name

Age / Sex : 43 Years/Male

Ref. Dr

Reg. Location : Bhayander East Main Centre

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