

PHYSICAL EXAMINATION REPORT

Patient Name Girish G. Pedinkon Sex/Age M-49
Date 10-02-2024. Location thank

History and Complaints

C/o-thalasenuta Minor

Pignientation onfice NAD.

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EXAMINATION FINDINGS:

Height (cms):

7 O Temp (0c)

Weight (kg):

1.781.144

Blood Pressure

Nails:

Pulse

Lymph Node:

Systems:

Cardiovascular:

Respiratory:

Genitourinary:

GI System:

CNS:

Impression:

NAO.

I Hb (thalosemia trault | IDA)

1 AlGa Ratio, Lglobulein.

1 TOP'S, LHDL., 1 Bilisubin.

- Muld Hepatosperanguly, LVH



dvice: - Fron Supplement:
- Reg. Exercise, Low Fest Diest
- Physician's consultation: 0 Advice: R Hypertension: 1) IHD 2) Arrhythmia 3) **Diabetes Mellitus** 4) **Tuberculosis** 5) Asthama 6) **Pulmonary Disease** 7) Thyroid/ Endocrine disorders 8) Nervous disorders 9) GI system 10) Genital urinary disorder 11) Rheumatic joint diseases or symptoms 12) Blood disease or disorder 13) Cyston brehend Cancer/lump growth/cyst 14) Congenital disease 15) cyst Removal (Forehead) Surgeries 16) Musculoskeletal System 17) PERSONAL HISTORY: Alcohol 1) 2) Smoking Dr. Manasee Kulkarni Diet Medication 4) 2005/09/3439

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Date: 70/2/04

Name:- Cresh Pednelas Age: 19-49

EYE CHECK UP

Chief complaints: 1200

Systemic Diseases:

Past history:

Unaided Vision:

Aided Vision:

UM,

1326/6 HUMA 112

Refraction:

(Right Eye)

(Left Eye)

	Sph	Cyl	Axis	Vn	Sph	Cyl	Axis	Vn
Distance								
Vear								

Colour Vision: Normal / Abnormal

Remark: USC O QU

Spela



: 2404121825

Name

: MR. GIRISH PEDNEKAR

Age / Gender

: 49 Years / Male

Consulting Dr. Reg. Location

: -

: G B Road, Thane West (Main Centre)

Collected

Reported

: 10-Feb-2024 / 09:08 :10-Feb-2024 / 14:32

Authenticity Check

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Use a QR Code Scanner Application To Scan the Code

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

	CBC (Complet	te Blood Count), Blood	
PARAMETER	RESULTS	BIOLOGICAL REF RANGE	METHOD
RBC PARAMETERS			
Haemoglobin	10.2	13.0-17.0 g/dL	Spectrophotometric
RBC	4.94	4.5-5.5 mil/cmm	Elect. Impedance
PCV	31.7	40-50 %	Measured
MCV	64.2	80-100 fl	Calculated
MCH	20.6	27-32 pg	Calculated
MCHC	32.1	31.5-34.5 g/dL	Calculated
RDW	16.1	11.6-14.0 %	Calculated
WBC PARAMETERS			
WBC Total Count	6870	4000-10000 /cmm	Elect. Impedance
WBC DIFFERENTIAL AND	ABSOLUTE COUNTS		and an imposition
Lymphacytes	20.4		

VVDC Total Count	6870	4000-10000 /cmm	Elect. Impeda
WBC DIFFERENTIAL AND	ABSOLUTE COUNTS		
Lymphocytes	30.1	20-40 %	
Absolute Lymphocytes	2067.9	1000-3000 /cmm	Calculated
Monocytes	6.4	2-10 %	
Absolute Monocytes	439.7	200-1000 /cmm	Calculated
Neutrophils	60.1	40-80 %	
Absolute Neutrophils	4128.9	2000-7000 /cmm	Calculated
Eosinophils	2.7	1-6 %	
Absolute Eosinophils	185.5	20-500 /cmm	Calculated
Basophils	0.7	0.1-2 %	
Absolute Basophils	48.1	20-100 /cmm	Calculated

WBC Differential Count by Absorbance & Impedance method/Microscopy.

PLATELET PARAMETERS Platelet Count

Immature Leukocytes

Hypochromia Microcytosis

260000 MPV 8.6 PDW 11.8 **RBC MORPHOLOGY**

6-11 fl 11-18 %

150000-400000 /cmm

Elect. Impedance Calculated

Calculated

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Macrocytosis

Anisocytosis

Mild

Poikilocytosis

Mild

Polychromasia

Target Cells

Basophilic Stippling

Normoblasts

Others

Elliptocytes-occasional

WBC MORPHOLOGY

PLATELET MORPHOLOGY

COMMENT

Features suggest thalassemia trait and/or iron deficiency anemia

Advice: Iron studies, Serum ferritin, Hb Electrophoresis & Reticulocyte count estimation recommended.

Specimen: EDTA Whole Blood

ESR, EDTA WB-ESR

5

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:

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

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*** End Of Report ***

Dr.IMRAN MUJAWAR M.D (Path) Pathologist

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Collected Reported : 10-Feb-2024 / 12:51 : 10-Feb-2024 / 17:31

Hexokinase

Hexokinase

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

PARAMETER	RESULTS	BIOLOGICAL REF RA	ANGE METHOD

GLUCOSE (SUGAR) FASTING, Fluoride Plasma

IG.

93.7

Non-Diabetic: < 100 mg/dl Impaired Fasting Glucose:

100-125 mg/dl

Diabetic: >/= 126 mg/dl

GLUCOSE (SUGAR) PP, Fluoride 117.3

Plasma PP/R

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

140-199 mg/dl

Absent Absent

Diabetic: >/= 200 mg/dl

Urine Sugar (Fasting)

Absent

Urine Ketones (Fasting)

Absent

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

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J. Mujawar Dr. IMRAN MUJAWAR

M.D (Path) Pathologist

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

PARAMETER	RESULTS	BIOLOGICAL REF RANGE	METHOD
BLOOD UREA, Serum	18.0	12.8-42.8 mg/dl	Urease & GLDH
BUN, Serum	8.4	6-20 mg/dl	Calculated
CREATININE, Serum	0.92	0.67-1.17 mg/dl	Enzymatic
eGFR, Serum	102	(ml/min/1.73sqm) Normal or High: Above 90 Mild decrease: 60-89 Mild to moderate decrease: 45-59 Moderate to severe decrease: 3-44 Severe decrease: 15-29 Kidney failure: <15	

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

TOTAL PROTEINS, Serum	6.7	6.4-8.3 g/dL	Biuret
ALBUMIN, Serum	4.8	3.5-5.2 g/dL	BCG
GLOBULIN, Serum	1.9	2.3-3.5 g/dL	Calculated
A/G RATIO, Serum	2.5	1 - 2	Calculated
URIC ACID, Serum	6.9	3.5-7.2 mg/dl	Uricase
PHOSPHORUS, Serum	3.3	2.7-4.5 mg/dl	Ammonium molybdate
CALCIUM, Serum	8.8	8.6-10.0 mg/dl	N-BAPTA
SODIUM, Serum	137	135-148 mmol/l	ISE
POTASSIUM, Serum	4.8	3.5-5.3 mmol/l	ISE
CHLORIDE, Serum	104	98-107 mmol/l	ISE

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

Dr.IMRAN MUJAWAR M.D (Path)

M.D (Path)

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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 4.9

Non-Diabetic Level: < 5.7 % Prediabetic Level: 5.7-6.4 %

Diabetic Level: >/= 6.5 %

HPLC

(HbA1c), EDTA WB - CC Estimated Average Glucose

(eAG), EDTA WB - CC

93.9

mg/dl

Calculated

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 G B Road Lab, Thane West End Of Report *

> Dr.IMRAN MUJAWAR M.D (Path)

Mujawar

Pathologist

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Reg. Location

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

PARAMETER

RESULTS

BIOLOGICAL REF RANGE METHOD

TOTAL PSA, Serum

0.62

<4.0 ng/ml

CLIA

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



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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.

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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: 2404121825

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Age / Gender

: 49 Years / Male

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: 10-Feb-2024 / 09:08 :10-Feb-2024 / 12:14

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

PARAMETER	RESULTS	BIOLOGICAL REF RANGE	METHOD
PHYSICAL EXAMINATION			
Color	Pale yellow	Pale Yellow	
Reaction (pH)	Acidic (6.0)	4.5 - 8.0	Chemical Indicator
Specific Gravity	1.030	1.010-1.030	Chemical Indicator
Transparency	Clear	Clear	
Volume (ml)	40	Tambu. (803)	
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 EXAMINATIO	N		
Leukocytes(Pus cells)/hpf	2-3	0-5/hpf	
Red Blood Cells / hpf	Absent	0-2/hpf	
Epithelial Cells / hpf	2-3		
Casts	Absent	Absent	
Crystals	Absent	Absent	
Amorphous debris	Absent	Absent	
Bacteria / hpf	3-4	Less than 20/hpf	
Others	THE STATE OF THE PARTY		
Internation Theory	1 (6)		

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 G B Road Lab, Thane West *** End Of Report **

> Dr.IMRAN MUJAWAR M.D (Path)

Mujawar

Pathologist

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

PARAMETER

RESULTS

ABO GROUP

В

Rh TYPING

Positive

NOTE: Test performed by Semi- automated column agglutination technology (CAT)

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 G B Road Lab, Thane West
*** End Of Report ***

Dr.IMRAN MUJAWAR M.D (Path) Pathologist

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

PARAMETER	RESULTS	BIOLOGICAL REF RANGE	METHOD
CHOLESTEROL, Serum	125.3	Desirable: <200 mg/dl Borderline High: 200-239mg/dl High: >/=240 mg/dl	CHOD-POD
TRIGLYCERIDES, Serum	205.2	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	26.9	Desirable: >60 mg/dl Borderline: 40 - 60 mg/dl Low (High risk): <40 mg/dl	Homogeneous enzymatic colorimetric assay
NON HDL CHOLESTEROL, Serum	98.4	Desirable: <130 mg/dl Borderline-high:130 - 159 mg/dl High:160 - 189 mg/dl Very high: >/=190 mg/dl	Calculated
LDL CHOLESTEROL, Serum	76.5	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	21.9	< /= 30 mg/dl	Calculated
CHOL / HDL CHOL RATIO, Serum	4.7	0-4.5 Ratio	Calculated
LDL CHOL / HDL CHOL RATIO, Serum	2.8	0-3.5 Ratio	Calculated

Kindly correlate clinically.

Note: LDL test is performed by direct measurement.

*Sample processed at SUBURBAN DIAGNOSTICS (INDIA) PVT. LTD G B Road Lab, Thane West
** End Of Report ***

Dr.VANDANA KULKARNI

M.D (Path)
Pathologist

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

PARAMETER	RESULTS	BIOLOGICAL REF RANGE	METHOD
Free T3, Serum	5.0	3.5-6.5 pmol/L	ECLIA
Free T4, Serum	15.5	11.5-22.7 pmol/L	ECLIA
sensitiveTSH, Serum	1.4	0.35-5.5 microIU/ml mIU/ml	ECLIA



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:10-Feb-2024 / 13:16 Reported

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 upto 15 microIU/ml should be correlated clinically or repeat the test with new sample as physiological

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

- 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.

- 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 G B Road Lab, Thane West *** End Of Report ***

> Mujawar Dr.IMRAN MUJAWAR M.D (Path) Pathologist

> > Page 13 of 14



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

PARAMETER	RESULTS	BIOLOGICAL REF RANGE	METHOD
BILIRUBIN (TOTAL), Serum	1.41	0.1-1.2 mg/dl	Diazo
BILIRUBIN (DIRECT), Serum	0.52	0-0.3 mg/dl	Diazo
BILIRUBIN (INDIRECT), Serum	0.89	0.1-1.0 mg/dl	Calculated
TOTAL PROTEINS, Serum	6.7	6.4-8.3 g/dL	Biuret
ALBUMIN, Serum	4.8	3.5-5.2 g/dL	BCG
GLOBULIN, Serum	1.9	2.3-3.5 g/dL	Calculated
A/G RATIO, Serum	2.5	1 - 2	Calculated
SGOT (AST), Serum	30.5	5-40 U/L	IFCC without pyridoxal phosphate activation
SGPT (ALT), Serum	20.0	5-45 U/L	IFCC without pyridoxal phosphate activation
GAMMA GT, Serum	21.1	3-60 U/L	IFCC
ALKALINE PHOSPHATASE, Serum	69.5	40-130 U/L	PNPP

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

Dr.IMRAN MUJAWAR M.D (Path) Pathologist

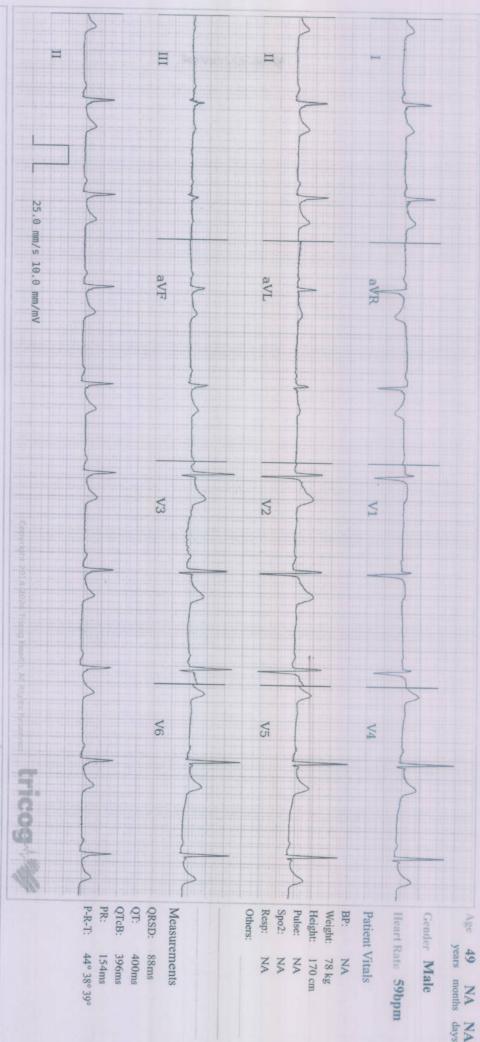
Page 14 of 14

SUBURBAN DIAGNOSTICS - G B ROAD, THANE WEST



Patient Name: GIRISH PEDNEKAR Patient ID: 2404121825

Date and Time: 10th Feb 24 9:47 AM



ECG Within Normal Limits: Early repolarization with an ascending ST segment, Sinus Bradycardia. Please correlate clinically.

REPORTED BY

DR SHAILAJA PILLAI MBBS, MD Physican MD Physican 49972



Authenticity Check <<ORCode>>

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CID

: 2404121825

Name

: Mr GIRISH PEDNEKAR

Age / Sex

Reg. Location

: 49 Years/Male

Ref. Dr

: G B Road, Thane West Main Centre

Reg. Date

Reported

Use a QR Code Seanner Application To Scan the Code

: 10-Feb-2024

: 10-Feb-2024 / 13:19

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.

-End of Report---

Dr Gauri Varma Consultant Radiologist MBBS / DMRE

MMC- 2007/12/4113

Charles

Click here to view images << lmageLink>>



Reg. No. : 2404121825	Sex : MALE	
Name : MR. GIRISH PEDNEKAR	Age: 49 YRS	
Ref. By :	Date: 10.02.2024	

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USG ABDOMEN AND PELVIS

<u>LIVER:</u> Liver appears mildly enlarged in size(16.4 cm) and shows homogenous echoreflectivity. 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 $11.0 \times 4.6 \text{ cm}$. Left kidney measures $10.3 \times 4.5 \text{ cm}$. Both kidneys are normal in shape and echotexture. Corticomedullary differentiation is maintained. There is no evidence of any hydronephrosis, hydroureter or calculus.

<u>SPLEEN:</u> Spleen is mildly enlarged in size (15.7 cm) and normal 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 is normal in size, 20 cc in volume and shows normal echotexture. No evidence of any focal lesion. Median lobe does not show significant hypertrophy.

No free fluid or significant lymphadenopathy is seen.



Reg. No. : 2404121825	Sex : MALE
Name : MR. GIRISH PEDNEKAR	Age: 49 YRS
Ref. By :	Date: 10.02.2024

IMPRESSION:

- MILD HEPATOMEGALY.
- MILD SPLENOMEGALY.

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.

Advice: Clinical co-relation and further evaluation.

DR. SHIVANGINI V. INGOLE M.B.B.S., DMRE (CONSULTANT RADIOLOGIST) REG NO. 2018/12/6130



REG NO: 2404121825	SEX : MALE
NAME : MR. GIRISH PEDNEKAR	AGE: 49 YRS
REF BY DR:	DATE: 10.02.2024

2D ECHOCARDIOGRAPHY

M - MODE FINDINGS:

LVIDD	54	mm
LVIDS	27	mm
LVEF	60	%
IVS	13	mm
PW	7	mm
AO	20	mm
LA	36	mm

2D ECHO:

- · All cardiac chambers are normal in size
- Left ventricular contractility: Normal
- · Regional wall motion abnormality: Absent.
- Systolic thickening: Normal. LVEF = 60%
- Mitral, tricuspid, aortic, pulmonary valves are: Normal.
- · Great arteries: Aorta and pulmonary artery are: Normal.
- Inter artrial and inter ventricular septum are intact.
- · Pulmonary veins, IVC, hepatic veins are normal.
- No pericardial effusion. No intracardiac clots or vegetation.



PATIENT NAME: MR.GIRISH PEDNEKAR

COLOR DOPPLER:

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- Mitral valve doppler E- 0.9 m/s, A- 0.6 m/s.
- Mild TR.
- No aortic / mitral regurgition. Aortic velocity 1.5 m/s, PG 9.6 mmHg
- No significant gradient across aortic valve.
- No diastolic dysfunction.

IMPRESSION:

- MILD CONCENTRIC HYPERTROPHY OF LV
- NO REGIONAL WALL MOTION ABNORMALITY AT REST.
- NORMAL LV SYSTOLIC FUNCTION.

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

DR. VOGESH KHARCHE
DNB (MEDICINE) DNB (CARDIOLOGY)
CONSULTANAT INTERVENTIONAL CARDIOLOGIST.