

PATIENT NAME: CHINTAN TRIVEDI (PKG10000289) EC 85778 REF. DOCTOR: DR. BANK OF BARODA

CODE/NAME & ADDRESS: C000138355 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, LADO SARAI, MEHRAULISOUTH WEST

DELHI

NEW DELHI 110030 8800465156

ACCESSION NO: 0290WL001393

PATIENT ID : CHINM310882290

CLIENT PATIENT ID: ABHA NO

AGE/SEX :41 Years

DRAWN

RECEIVED: 09/12/2023 13:17:27 REPORTED :11/12/2023 16:24:52

Test Report Status Results **Biological Reference Interval** Units <u>Final</u>

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

XRAY-CHEST

BOTH THE LUNG FIELDS ARE CLEAR

BOTH THE COSTOPHRENIC AND CARIOPHRENIC ANGELS ARE CLEAR

BOTH THE HILA ARE NORMAL

CARDIAC AND AORTIC SHADOWS APPEAR NORMAL **»**» BOTH THE DOMES OF THE DIAPHRAM ARE NORMAL >> >>

VISUALIZED BONY THORAX IS NORMAL **»**»

NO ABNORMALITY DETECTED **IMPRESSION**

> Dr G.S. Saluja, (MBBS, DMRD) (Consultant Radiologist)

ECG

ECG SINUS RHYTHM.

OTHERWISE NORMAL ECG.

MEDICAL HISTORY

RELEVANT PRESENT HISTORY NOT SIGNIFICANT RELEVANT PAST HISTORY NOT SIGNIFICANT RELEVANT PERSONAL HISTORY NOT SIGNIFICANT **NOT SIGNIFICANT** RELEVANT FAMILY HISTORY OCCUPATIONAL HISTORY **NOT SIGNIFICANT** HISTORY OF MEDICATIONS **NOT SIGNIFICANT**

ANTHROPOMETRIC DATA & BMI

HEIGHT IN METERS 1.69 mts WEIGHT IN KGS. Kgs 55

RMI 19 BMI & Weight Status as follows/sqmts

Below 18.5: Underweight 18.5 - 24.9: Normal 25.0 - 29.9: Overweight 30.0 and Above: Obese

GENERAL EXAMINATION

MENTAL / EMOTIONAL STATE NORMAL

Dr. Arpita Pasari, MD **Consultant Pathologist**





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PHYSICAL ATTITUDE NORMAL
GENERAL APPEARANCE / NUTRITIONAL HEALTHY

STATUS

BUILT / SKELETAL FRAMEWORK

FACIAL APPEARANCE

SKIN

UPPER LIMB

LOWER LIMB

NORMAL

NORMAL

NORMAL

NORMAL

NECK LYMPHATICS / SALIVARY GLANDS NOT ENLARGED OR TENDER

THYROID GLAND NOT ENLARGED

CAROTID PULSATION NORMAL TEMPERATURE AFEBRILE

PULSE 65/MIN, REGULAR, ALL PERIPHERAL PULSES WELL FELT, NO CAROTID

BRUIT

RESPIRATORY RATE NORMAL

CARDIOVASCULAR SYSTEM

BP 110/70 MM HG mm/Hg

(SUPINE)

PERICARDIUM NORMAL
APEX BEAT NORMAL
HEART SOUNDS NORMAL
MURMURS ABSENT

RESPIRATORY SYSTEM

SIZE AND SHAPE OF CHEST NORMAL
MOVEMENTS OF CHEST SYMMETRICAL
BREATH SOUNDS INTENSITY NORMAL

BREATH SOUNDS QUALITY VESICULAR (NORMAL)

ADDED SOUNDS ABSENT

PER ABDOMEN

APPEARANCE NORMAL
VENOUS PROMINENCE ABSENT
LIVER NOT PALPABLE
SPLEEN NOT PALPABLE

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HERNIA ABSENT

CENTRAL NERVOUS SYSTEM

HIGHER FUNCTIONS

CRANIAL NERVES

CEREBELLAR FUNCTIONS

SENSORY SYSTEM

MOTOR SYSTEM

REFLEXES

NORMAL

NORMAL

MUSCULOSKELETAL SYSTEM

SPINE NORMAL JOINTS NORMAL

BASIC EYE EXAMINATION

CONJUNCTIVA NORMAL
EYELIDS NORMAL
EYE MOVEMENTS NORMAL
CORNEA NORMAL

DISTANT VISION RIGHT EYE WITHOUT

GLASSES

DISTANT VISION LEFT EYE WITHOUT

GLASSES

NEAR VISION RIGHT EYE WITHOUT GLASSES N6, WITHIN NORMAL LIMIT NEAR VISION LEFT EYE WITHOUT GLASSES N6, WITHIN NORMAL LIMIT

COLOUR VISION

NORMAL

6/12, VISUAL ACUITY FOR CORRECTION

6/12, VISUAL ACUITY FOR CORRECTION

BASIC ENT EXAMINATION

EXTERNAL EAR CANAL NORMAL TYMPANIC MEMBRANE NORMAL

NOSE NO ABNORMALITY DETECTED

SINUSES NORMAL THROAT NORMAL

TONSILS NOT ENLARGED

BASIC DENTAL EXAMINATION

TEETH NORMAL GUMS HEALTHY

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SUMMARY

RELEVANT HISTORY

RELEVANT GP EXAMINATION FINDINGS

NOT SIGNIFICANT

REMARKS / RECOMMENDATIONS

NONE

FITNESS STATUS

FITNESS STATUS FIT (WITH MEDICAL ADVICE) (AS PER REQUESTED PANEL OF TESTS)

Comments

CLINICAL FINDINGS:-

DYSLIPIDEMIA.

RAISED HbA1C AND ESTIMATED AVERAG GLUCOSE (EAG)

FITNESS STATUS :-

FITNESS STATUS: FIT (WITH MEDICAL ADVICE) (AS PER REQUESTED PANEL OF TESTS)

ADVICE :- LOW FAT WITH HIGH FIBER DIET AND REGULAR PHYSICAL EXERCISE FOR DYSLIPIDEMIA.

NEED PHYSICIAN CONSULTATION FOR LIFE STYLE MODIFICATION.

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ULTRASOUND ABDOMEN

ULTRASOUND ABDOMEN

Liver is normal in size, shape and echotexture. Intra & Extra hepatic biliary radicals are normal. Portal vein and C.B.D are normal in caliber

Gall Bladder is normal, thin walled & its lumen is echo free.

Spleen is normal in size, shape & echotexture.

Pancreas is normal in size, shape & echotexture.

Both Kidneys are normal in size, shape and echotexture. Central pelvicalyceal system is normal. Corticomedullary differentiation is maintained.

IVC and AO is normal in caliber.

Urinary Bladder is normal thin walled, there is no calculus.

Prostate is normal in size & echotexture.

IMPRESSION- No Significant abnormality seen in USG of Whole Abdomen.

Dr G S Saluja (MBBS.DMRD) REG.NO 4005 (Consultant Radiologist) TMT OR ECHO

CLINICAL PROFILE

Parasternal long axis, Parasternal short axis at multiple levels, apical 4-C & apical & 5-C views taken.

All cardiac valves are normal in structure & move normally.

All cardiac chambers and great vessels are normal in size.

The left ventricular wall is normal in thickness & contractility.

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There is no evidence of any regional wall motion abnormality.

There is no evidence of any vegetation or clot or pericardial effusion.

The calculated LVEF 70%.

IMPRESSION:

- Normal 2D Echo Study
- LVEF 70%

M-MODE ECHOCARDIOGRAPHY

(1) MITRAL VALVE DIMENSIONS Normal Value

> **EPSS** 2-7 mm: mm

(2) AORTIC VALVE DIMENSIONS

Aortic Root 27 20-37 mm : mm Left atrium 31 19-40 mm : mm Cusp Opening 2115-26 mm : mm

(3) LEFT VENTRICULAR DIMENSIONS

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DIMENSION	OBSERVED	NORMAL VALUES
LVID (Diastolic)	35 : mm	37-56 mm
LVID (Systolic)	24 : mm	24-42 mm
RVID (Diastolic)	12 : mm	7-23 mm
IVST (Diastolic)	9 : mm	6-11 mm
LVPWT (Diastolic)	9 : mm	6-11 mm

LEFT VENTRICULAR FUNCTION

LVEDV : ml LVESV : ml EF 70 : %

COLOR DOPPLER FUNCTION

PEAK VELOCITY M/SEC	MAX.	GRADIENT	MMHG	REGURGITATION	
PV8					
MV8/.3					
AV- 1					
TV 1					

Interpretation(s)

THIS REPORT CARRIES THE SIGNATURE OF OUR LABORATORY DIRECTOR. THIS IS AN INVIOLABLE FEATURE OF OUR LAB MANAGEMENT SOFTWARE. HOWEVER, ALL EXAMINATIONS AND INVESTIGATIONS HAVE BEEN CONDUCTED BY OUR PANEL OF DOCTORS.



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:41 Years

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FITNESS STATUS-Conclusion on an individual's Fitness, which is commented upon mainly for Pre employment cases, is based on multi factorial findings and does not depend on any one single parameter. The final Fitness assigned to a candidate will depend on the Physician's findings and overall judgement on a case to case basis, details of the candidate's past and personal history as well as the comprehensiveness of the diagnostic panel which has been requested for .These are then further correlated with details of the job under consideration to eventually fit the right man to the right job.

Basis the above, Agilus diagnostic classifies a candidate's Fitness Status into one of the following categories:

- Fit (As per requested panel of tests) AGILUS Limited gives the individual a clean chit to join the organization, on the basis of the General Physical Examination and the specific test panel requested for.
- Fit (with medical advice) (As per requested panel of tests) This indicates that although the candidate can be declared as FIT to join the job, minimal problems have been detected during the Pre- employment examination. Examples of conditions which could fall in this category could be cases of mild reversible medical abnormalities such as height weight disproportions, borderline raised Blood Pressure readings, mildly raised Blood sugar and Blood Lipid levels, Hematuria, etc. Most of these relate to sedentary lifestyles and come under the broad category of life style disorders. The idea is to caution an individual to bring about certain lifestyle changes as well as seek a Physician """"s consultation and counseling in order to bring back to normal the mildly deranged parameters. For all purposes the individual is FIT to join the job.
 Fitness on Hold (Temporary Unfit) (As per requested panel of tests) Candidate's reports are kept on hold when either the diagnostic tests or the physical findings reveal
- Fitness on Hold (Temporary Unfit) (As per requested panel of tests) Candidate's reports are kept on hold when either the diagnostic tests or the physical findings reveal the presence of a medical condition which warrants further tests, counseling and/or specialist opinion, on the basis of which a candidate can either be placed into Fit, Fit (With Medical Advice), or Unfit category. Conditions which may fall into this category could be high blood pressure, abnormal ECG, heart murmurs, abnormal vision, grossly elevated blood sugars, etc.
- Unfit (As per requested panel of tests) An unfit report by Agilus diagnostic Limited clearly indicates that the individual is not suitable for the respective job profile e.g. total color blindness in color related jobs.

Berita

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H.	AEMATOLOGY - CBC		
MEDI WHEEL FULL BODY HEALTH CHECK UP AI	BOVE 40 MALE		
BLOOD COUNTS,EDTA WHOLE BLOOD			
HEMOGLOBIN (HB)	16.4	13.0 - 17.0	g/dL
RED BLOOD CELL (RBC) COUNT	5.40	4.5 - 5.5	mil/μL
WHITE BLOOD CELL (WBC) COUNT	6.37	4.0 - 10.0	thou/µL
PLATELET COUNT	246	150 - 410	thou/µL
RBC AND PLATELET INDICES			
HEMATOCRIT (PCV)	46.7	40 - 50	%
MEAN CORPUSCULAR VOLUME (MCV)	86.5	83 - 101	fL
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	30.4	27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC)	35.1 High	31.5 - 34.5	g/dL
RED CELL DISTRIBUTION WIDTH (RDW)	10.4 Low	11.6 - 14.0	%
MENTZER INDEX	16.0		
MEAN PLATELET VOLUME (MPV)	8.6	6.8 - 10.9	fL
WBC DIFFERENTIAL COUNT			
NEUTROPHILS	50	40 - 80	%
LYMPHOCYTES	38	20 - 40	%
MONOCYTES	06	2 - 10	%
EOSINOPHILS	06	1 - 6	%
BASOPHILS	00	0 - 2	%
ABSOLUTE NEUTROPHIL COUNT	3.18	2.0 - 7.0	thou/µL
ABSOLUTE LYMPHOCYTE COUNT	2.42	1 - 3	thou/µL
ABSOLUTE MONOCYTE COUNT	0.38	0.20 - 1.00	thou/µL
ABSOLUTE EOSINOPHIL COUNT	0.38	0.02 - 0.50	thou/µL

Interpretation(s)
BLOOD COUNTS,EDTA WHOLE BLOOD-The cell morphology is well preserved for 24hrs. However after 24-48 hrs a progressive increase in MCV and HCT is observed leading to a decrease in MCHC. A direct smear is recommended for an accurate differential count and for examination of RBC morphology. RBC AND PLATELET INDICES-Mentzer index (MCV/RBC) is an automated cell-counter based calculated screen tool to differentiate cases of Iron deficiency anaemia(>13) from Beta thalassaemia trait

(<13) in patients with microcytic anaemia. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HbA2 remains the gold standard for diagnosing a case of beta thalassaemia trait.

WBC DIFFERENTIAL COUNT-The optimal threshold of 3.3 for NLR showed a prognostic possibility of clinical symptoms to change from mild to severe in COVID positive patients. When age = 49.5 years old and NLR = 3.3, 46.1% COVID-19 patients with mild disease might become severe. By contrast, when age < 49.5 years old and NLR <

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3.3, COVID-19 patients tend to show mild disease. (Reference to - The diagnostic and predictive role of NLR, d-NLR and PLR in COVID-19 patients A.-P. Yang, et al. International Immunopharmacology 84 (2020) 106504 This ratio element is a calculated parameter and out of NABL scope.

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mm at 1 hr

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HAEMATOLOGY

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

ERYTHROCYTE SEDIMENTATION RATE (ESR), EDTA BLOOD

0 - 14E.S.R

METHOD: MODIFIED WESTERGREN

GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE **BLOOD**

HBA1C 6.0 High Non-diabetic: < 5.7 %

> Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5Therapeutic goals: < 7.0 Action suggested: > 8.0 (ADA Guideline 2021)

METHOD: HPLC TECHNOLOGY

ESTIMATED AVERAGE GLUCOSE(EAG) 125.5 High < 116.0 mg/dL

Interpretation(s)

ERYTHROCYTE SEDIMENTATION RATE (ESR),EDTA BLOOD-**TEST DESCRIPTION**:Erythrocyte sedimentation rate (ESR) is a test that indirectly measures the degree of inflammation present in the body. The test actually measures the rate of fall (sedimentation) of erythrocytes in a sample of blood that has been placed into a tall, thin, vertical tube. Results are reported as the millimetres of clear fluid (plasma) that are present at the top portion of the tube after one hour. Nowadays fully automated instruments are available to measure ESR.

ESR is not diagnostic it is a non-specific test that may be elevated in a number of different conditions. It provides general information about the presence of an inflammatory condition.CRP is superior to ESR because it is more sensitive and reflects a more rapid change

TEST INTERPRETATION

Increase in: Infections, Vasculities, Inflammatory arthritis, Renal disease, Anemia, Malignancies and plasma cell dyscrasias, Acute allergy Tissue injury, Pregnancy, Estrogen medication, Aging.
Finding a very accelerated ESR(>100 mm/hour) in patients with ill-defined symptoms directs the physician to search for a systemic disease (Paraproteinemias,

Disseminated malignancies, connective tissue disease, severe infections such as bacterial endocarditis).

In pregnancy BRI in first trimester is 0-48 mm/hr(62 if anemic) and in second trimester (0-70 mm /hr(95 if anemic). ESR returns to normal 4th week post partum. **Decreased** in: Polycythermia vera, Sickle cell anemia

LIMITATIONS

False elevated ESR: Increased fibrinogen, Drugs(Vitamin A, Dextran etc), Hypercholesterolemia

False Decreased: Poikilocytosis, (SickleCells, spherocytes), Microcytosis, Low fibrinogen, Very high WBC counts, Drugs (Quinine,

salicylates)

1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition 2. Paediatric reference intervals. AACC Press, 7th edition. Edited by S. Soldin 3. The reference for the adult reference range is "Practical Haematology by Dacie and Lewis,10th edition. GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD-**Used For**:

- 1. Evaluating the long-term control of blood glucose concentrations in diabetic patients.
- 2. Diagnosing diabetes.3. Identifying patients at increased risk for diabetes (prediabetes).

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The ADA recommends measurement of HbA1c (typically 3-4 times per year for type 1 and poorly controlled type 2 diabetic patients, and 2 times per year for well-controlled type 2 diabetic patients) to determine whether a patients metabolic control has remained continuously within the target range.

1. eAG (Estimated average glucose) converts percentage HbA1c to md/dl, to compare blood glucose levels.

- eAG gives an evaluation of blood glucose levels for the last couple of months.
 eAG is calculated as eAG (mg/dl) = 28.7 * HbA1c 46.7

HbA1c Estimation can get affected due to :

- 1. Shortened Erythrocyte survival: Any condition that shortens erythrocyte survival or decreases mean erythrocyte age (e.g. recovery from acute blood loss, hemolytic anemia) will falsely lower HbA1c test results. Fructosamine is recommended in these patients which indicates diabetes control over 15 days.
- 2.Vitamin C & E are reported to falsely lower test results. (possibly by inhibiting glycation of hemoglobin.
 3. Iron deficiency anemia is reported to increase test results. Hypertriglyceridemia, uremia, hyperbilirubinemia, chronic alcoholism, chronic ingestion of salicylates & opiates addiction are reported to interfere with some assay methods, falsely increasing results.

 4. Interference of hemoglobinopathies in HbA1c estimation is seen in
- a) Homozygous hemoglobinopathy. Fructosamine is recommended for testing of HbA1c. b) Heterozygous state detected (D10 is corrected for HbS & HbC trait.)
- c) HbF > 25% on alternate paltform (Boronate affinity chromatography) is recommended for testing of HbA1c.Abnormal Hemoglobin electrophoresis (HPLC method) is recommended for detecting a hemoglobinopathy

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IMMUNOHAEMATOLOGY

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

TYPE O **ABO GROUP**

METHOD: TUBE AGGLUTINATION

RH TYPE **POSITIVE**

METHOD: TUBE AGGLUTINATION

Interpretation(s)
ABO GROUP & RH TYPE, EDTA WHOLE BLOOD-Blood group is identified by antigens and antibodies present in the blood. Antigens are protein molecules found on the surface of red blood cells. Antibodies are found in plasma. To determine blood group, red cells are mixed with different antibody solutions to give A,B,O or AB.

Disclaimer: "Please note, as the results of previous ABO and Rh group (Blood Group) for pregnant women are not available, please check with the patient records for availability of the same.

The test is performed by both forward as well as reverse grouping methods.

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MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

GLUCOSE FASTING, FLUORIDE PLASMA

FBS (FASTING BLOOD SUGAR)

METHOD: HEXOKINASE

GLUCOSE, POST-PRANDIAL, PLASMA

PPBS(POST PRANDIAL BLOOD SUGAR)

125

94

Normal: < 140,

74 - 99

Impaired Glucose Tolerance: 140-199

Diabetic > or = 200

METHOD: HEXOKINASE

TRIGLYCERIDES

LIPID PROFILE WITH CALCULATED LDL

METHOD: OXIDASE, ESTERASE, PEROXIDASE

CHOLESTEROL, TOTAL

197

Desirable: <200

mg/dL

mg/dL

mg/dL

BorderlineHigh: 200-239 High: > or = 240

154 High

Desirable: < 150

mg/dL

Borderline High: 150 - 199 High: 200 - 499

Very High: > or = 500

METHOD: ENZYMATIC ASSAY

HDL CHOLESTEROL

VERY LOW DENSITY LIPOPROTEIN

40

< 40 Low

mg/dL

> or = 60 High

METHOD: DIRECT- NON IMMUNOLOGICAL

CHOLESTEROL LDL

126 High

30.8 High

Adult levels:

mg/dL

Optimal < 100

Near optimal/above optimal:

100-129

Borderline high: 130-159

High: 160-189

NON HDL CHOLESTEROL 157 High

Very high: = 190

Desirable: Less than 130 Above Desirable: 130 - 159

Borderline High: 160 - 189

High: 190 - 219

Very high: > or = 220

< or = 30

mg/dL

mg/dL

Dr. Arpita Pasari, MD **Consultant Pathologist**

METHOD: CALCULATED





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PATIENT NAME: CHINTAN TRIVEDI (PKG10000289) EC 85778 REF. DOCTOR: DR. BANK OF BARODA

CODE/NAME & ADDRESS: C000138355 ACCESSION NO: 0290WL001393 AGE/SEX :41 Years

ARCOFEMI HEALTHCARE LTD (MEDIWHEEL

F-703, LADO SARAI, MEHRAULISOUTH WEST

DELHI

NEW DELHI 110030

8800465156

PATIENT ID : CHINM310882290

CLIENT PATIENT ID: ABHA NO

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Test Report Status Results **Biological Reference Interval** Units <u>Final</u>

METHOD: CALCULATED

4.9 High 3.3 - 4.4CHOL/HDL RATIO

3.2 High LDL/HDL RATIO 0.5 - 3.0 Desirable/Low Risk

3.1 - 6.0 Borderline/Moderate

Risk >6.0 High Risk

Interpretation(s)

Serum lipid profile is measured for cardiovascular risk prediction. Lipid Association of India recommends LDL-C as primary target and Non HDL-C as co-primary treatment target.

Risk Stratification for ASCVD (Atherosclerotic cardiovascular disease) by Lipid Association of India

tion of municipality for	rise is principle curaniman un	sensely by expressions of enum		
Risk Category				
Extreme risk group	A.CAD with > 1 feature of high risk group			
	B. CAD with > 1 feature of Very high risk s	group or recurrent ACS (within 1 year) despite LDL-C < or =		
	50 mg/dl or polyvascular disease			
Very High Risk	1. Established ASCVD 2. Diabetes with 2:	major risk factors or evidence of end organ damage 3.		
	Familial Homozygous Hypercholesterolemi	a		
High Risk	1. Three major ASCVD risk factors. 2. Diabetes with 1 major risk factor or no evidence of end organ			
		90 mg/dl 5. Extreme of a single risk factor. 6. Coronary		
	Artery Calcium - CAC >300 AU. 7. Lipoprotein a >/= 50mg/dl 8. Non stenotic carotid plaque			
Moderate Risk	2 major ASCVD risk factors			
Low Risk	0-1 major ASCVD risk factors			
Major ASCVD (Ath	erosclerotic cardiovascular disease) Risk Fa	ectors		
Age > or = 45 years in males and > or = 55 years in females Current Cigarette smoking or tobacco use				
Family history of premature ASCVD 4. High blood pressure				
5. Low HDL				

Newer treatment goals and statin initiation thresholds based on the risk categories proposed by LAI in 2020.

Risk Group	Treatment Goals		Consider Drug Therapy	
	LDL-C (mg/dl)	Non-HDL (mg/dl)	LDL-C (mg/dl)	Non-HDL (mg/dl)
Extreme Risk Group Category A	<50 (Optional goal	< 80 (Optional goal	>OR = 50	>OR = 80
	< OR = 30)	<or 60)<="" =="" td=""><td></td><td></td></or>		
Extreme Risk Group Category B	<or 30<="" =="" td=""><td><or 60<="" =="" td=""><td>> 30</td><td>>60</td></or></td></or>	<or 60<="" =="" td=""><td>> 30</td><td>>60</td></or>	> 30	>60
Very High Risk	<50	<80	>OR= 50	>OR= 80
High Risk	<70	<100	>OR= 70	>OR= 100
Moderate Risk	<100	<130	>OR= 100	>OR= 130
Low Risk	<100	<130	>OR= 130*	>OR= 160

^{*}After an adequate non-pharmacological intervention for at least 3 months.

References: Management of Dyslipidaemia for the Prevention of Stroke: Clinical Practice Recommendations from the Lipid Association of India. Current Vascular Pharmacology, 2022, 20, 134-155.

LIVER FUNCTION PROFILE, SERUM

BILIRUBIN, TOTAL 0.41 0.0 - 1.2mg/dL

METHOD: JENDRASSIK AND GROFF

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Consultant Pathologist





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	İ	į	
Test Report Status <u>Final</u>	Results	Biological Reference Interv	al Units
	0.45		
BILIRUBIN, DIRECT METHOD: DIAZOTIZATION	0.15	0.0 - 0.2	mg/dL
BILIRUBIN, INDIRECT METHOD: CALCULATED	0.26	0.00 - 1.00	mg/dL
TOTAL PROTEIN METHOD: BIURET	7.0	6.4 - 8.3	g/dL
ALBUMIN METHOD: BROMOCRESOL GREEN	4.6	3.50 - 5.20	g/dL
GLOBULIN METHOD: CALCULATED	2.4	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO METHOD: CALCULATED	1.9	1.0 - 2.0	RATIO
ASPARTATE AMINOTRANSFERASE(AST/SGOT) METHOD: UV WITH P5P	16	UPTO 40	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT) METHOD: UV WITH P5P	16	UP TO 45	U/L
ALKALINE PHOSPHATASE METHOD: PNPP	93	40 - 129	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT) METHOD: G-GLUTAMYL-CARBOXY-NITROANILIDE	28	8 - 61	U/L
LACTATE DEHYDROGENASE METHOD: ENZYMATIC LACTATE - PYRUVATE(IFCC)	148	135 - 225	U/L
BLOOD UREA NITROGEN (BUN), SERUM			
BLOOD UREA NITROGEN METHOD: UREASE KINETIC	8	6 - 20	mg/dL
CREATININE, SERUM			
CREATININE METHOD: ALKALINE PICRATE KINETIC JAFFES	0.82	0.70 - 1.20	mg/dL
BUN/CREAT RATIO			
BUN/CREAT RATIO	9.76	5.0 - 15.0	
METHOD: CALCULATED			
URIC ACID, SERUM			
URIC ACID METHOD: URICASE/CATALASE UV	4.4	3.5 - 7.2	mg/dL

TOTAL PROTEIN, SERUM

Papite

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CLIENT PATIENT ID: ABHA NO : AGE/SEX :41 Years Male

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Test Report Status <u>Final</u>	Results	Biological Reference I	nterval Units
TOTAL PROTEIN METHOD: BIURET	7.0	6.4 - 8.3	g/dL
ALBUMIN, SERUM			
ALBUMIN METHOD: BROMOCRESOL GREEN	4.6	3.5 - 5.2	g/dL
GLOBULIN			
GLOBULIN	2.4	2.0 - 4.1	g/dL
ELECTROLYTES (NA/K/CL), SERUM			
SODIUM, SERUM METHOD: DIRECT ION SELECTIVE ELECTRODE	141.5	136.0 - 146.0	mmol/L
POTASSIUM, SERUM METHOD: DIRECT ION SELECTIVE ELECTRODE	4.45	3.50 - 5.10	mmol/L
CHLORIDE, SERUM METHOD: DIRECT ION SELECTIVE ELECTRODE	102.4	98.0 - 106.0	mmol/L

Interpretation(s)

Sodium	Potassium	Chloride
Decreased In:CCF,cirrhosis,	Decreased in: Low potassium	Decreased in: Vomiting, diarrhea,
vomiting, diarrhea, excessive	intake, prolonged vomiting or diarrhea,	renal failure combined with salt
sweating, salt-losing	RTA types I and II,	deprivation, over-treatment with
nephropathy, adrenal insufficiency,	hyperaldosteronism, Cushing's	diuretics, chronic respiratory acidosi
nephrotic syndrome, water	syndrome,osmotic diuresis (e.g.,	diabetic ketoacidosis, excessive
intoxication, SIADH. Drugs:	hyperglycemia), alkalosis, familial	sweating, SIADH, salt-losing
thiazides, diuretics, ACE inhibitors,	periodic paralysis,trauma	nephropathy, porphyria, expansion of
chlorpropamide,carbamazepine,anti	(transient).Drugs: Adrenergic agents,	extracellular fluid volume,
depressants (SSRI), antipsychotics.	diuretics.	adrenalinsufficiency,
		hyperaldosteronism, metabolic
		alkalosis. Drugs: chronic
		laxative, corticosteroids, diuretics.
Increased in: Dehydration	Increased in: Massive hemolysis,	Increased in: Renal failure, nephrotic
excessives weating, severe	severe tissue damage, rhabdomyolysis,	syndrome, RTA, dehydration,
vomiting or diarrhea), diabetes	acidosis, dehydration, renal failure,	overtreatment with
mellitus, diabetesinsipidus,	Addison's disease, RTA type IV,	saline, hyperparathyroidism, diabetes
hyperaldosteronism, inadequate	hyperkalemic familial periodic	insipidus, metabolic acidosis from
water intake. Drugs: steroids,	paralysis. Drugs: potassium salts,	diarrhea (Loss of HCO3-), respiratory
licorice, oral contraceptives.	potassium- sparing diuretics, NSAIDs,	alkalosis, hyperadrenocorticism.
	beta-blockers, ACE inhibitors, high-	Drugs: acetazolamide, androgens,
	dose trimethoprim-sulfamethoxazole.	hydrochlorothiazide, salicylates.
Interferences: Severe lipemia or	Interferences: Hemolysis of sample,	Interferences:Test is helpful in
hyperproteinemi, if sodium analysis	delayed separation of serum,	assessing normal and increased anion
involves a dilution step can cause	prolonged fist clenching during blood	gap metabolic acidosis and in
spurious results. The serum sodium	drawing, and prolonged tourniquet	distinguishing hypercalcemia due to
falls about 1.6 mEq/L for each 100	placement. Very high WBC/PLT counts	hyperparathyroidism (high serum
mg/dL increase in blood glucose.	may cause spurious. Plasma potassium	chloride) from that due to malignance
	levels are normal.	(Normal serum chloride)

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DELHI

NEW DELHI 110030

8800465156

ACCESSION NO: 0290WL001393

PATIENT ID : CHINM310882290

CLIENT PATIENT ID: ABHA NO

AGE/SEX DRAWN

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:41 Years

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Test Report Status Results **Biological Reference Interval** <u>Final</u> Units

Interpretation(s)
GLUCOSE FASTING, FLUORIDE PLASMA-TEST DESCRIPTION

Normally, the glucose concentration in extracellular fluid is closely regulated so that a source of energy is readily available to tissues and sothat no glucose is excreted in the

Increased in:Diabetes mellitus, Cushing's syndrome (10 - 15%), chronic pancreatitis (30%). Drugs:corticosteroids,phenytoin, estrogen, thiazides. **Decreased in**: Pancreatic islet cell disease with increased insulin, insulinoma, adrenocortical insufficiency, hypopituitarism, diffuse liver disease, malignancy (adrenocortical, stomach, fibrosarcoma), infant of a diabetic mother, enzyme deficiency diseases (e.g. galactosemia), Drugs-insulin, ethanol, propranolol sulfonylureas,tolbutamide,and other oral hypoglycemic agents.

NOTE: While random serum glucose levels correlate with home glucose monitoring results (weekly mean capillary glucose values),there is wide fluctuation within

individuals.Thus, glycosylated hemoglobin(HbA1c) levels are favored to monitor glycemic control.

High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glyosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc.

GLUCOSE, POST-PRANDIAL, PLASMA-High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glyosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc. Additional test HbA1c LIVER FUNCTION PROFILE, SERUM-

Bilirubin is a yellowish pigment found in bile and is a breakdown product of normal heme catabolism. Bilirubin is excreted in bile and urine, and elevated levels may give yellow discoloration in jaundice. Elevated levels results from increased bilirubin production (eg, hemolysis and ineffective erythropoiesis), decreased bilirubin excretion (eg, obstruction and hepatitis), and abnormal bilirubin metabolism (eg, hereditary and neonatal jaundice). Conjugated (direct) bilirubin is elevated more than unconjugated (indirect) bilirubin in Viral hepatitis, Drug reactions, Alcoholic liver disease Conjugated (direct) bilirubin is also elevated more than unconjugated (indirect) bilirubin when there is some kind of blockage of the bile ducts like in Gallstones getting into the bile ducts, tumors &scarring of the bile ducts. Increased unconjugated (indirect) bilirubin may be a result of Hemolytic or pernicious anemia, Transfusion reaction & a common metabolic condition termed Gilbert syndrome, due to low levels of the enzyme that attaches sugar molecules to bilirubin.

AST is an enzyme found in various parts of the body. AST is found in the liver, heart, skeletal muscle, kidneys, brain, and red blood cells, and it is commonly measured clinically as a marker for liver health. AST levels increase during chronic viral hepatitis, blockage of the bile duct, cirrhosis of the liver, liver cancer, kidney failure, hemolytic anemia, pancreatitis, hemochromatosis. AST levels may also increase after a heart attack or strenuous activity. ALT test measures the amount of this enzyme in the blood. ALT is found mainly in the liver, but also in smaller amounts in the kidneys, heart, muscles, and pancreas. It is commonly measured as a part of a diagnostic evaluation of hepatocellular injury, to determine liver health. AST levels increase during acute hepatitis, sometimes due to a viral infection, is chemia to the liver, chronic

hepatitis, obstruction of bile ducts, cirrhosis. **ALP** is a protein found in almost all body tissues. Tissues with higher amounts of ALP include the liver, bile ducts and bone. Elevated ALP levels are seen in Biliary obstruction, Osteoblastic bone tumors, osteomalacia, hepatitis, Hyperparathyroidism, Leukemia, Lymphoma, Pagets disease,Rickets,Sarcoidosis etc. Lower-than-normal ALP levels seen

GGT is an enzyme found in cell membranes of many tissues mainly in the liver, kidney and pancreas. It is also found in other tissues including intestine, spleen, heart, brain and seminal vesicles. The highest concentration is in the kidney, but the liver is considered the source of normal enzyme activity. Serum GGT has been widely used as an index of liver dysfunction. Elevated serum GGT activity can be found in diseases of the liver, billiary system and pancreas. Conditions that increase serum GGT are obstructive liver disease, high alcohol consumption and use of enzyme-inducing drugs etc.

Total Protein also known as total protein, is a biochemical test for measuring the total amount of protein in serum. Protein in the plasma is made up of albumin and globulin. Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenstroms disease.Lower-than-normal levels may be due to: Agammaglobulinemia,Bleeding (hemorrhage),Burns,Glomerulonephritis,Liver disease, Malabsorption,Malnutrition,Nephrotic

syndrome, Protein-losing enteropathy etc. **Albumin** is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by:Liver disease like cirrhosis of the liver, nephrotic syndrome,protein-losing enteropathy,Burns,hemodilution,increased vascular permeability or decreased lymphatic clearance,malnutrition and wasting etc

BLOOD UREA NITROGEN (BUN), SERUM-Causes of Increased levels include Pre renal (High protein diet, Increased protein catabolism, GI haemorrhage, Cortisol, Dehydration, CHF Renal), Renal Failure, Post Renal (Malignancy, Nephrolithiasis, Prostatism)

Causes of decreased level include Liver disease, SIADH.

CREATININE, SERUM-Higher than normal level may be due to:

• Blockage in the urinary tract, Kidney problems, such as kidney damage or failure, infection, or reduced blood flow, Loss of body fluid (dehydration), Muscle problems, such as breakdown of muscle fibers, Problems during pregnancy, such as seizures (eclampsia)), or high blood pressure caused by pregnancy (preeclampsia)

Lower than normal level may be due to:• Myasthenia Gravis, Muscuophy

URIC ACID, SERUM-Causes of Increased levels:-Dietary(High Protein Intake,Prolonged Fasting,Rapid weight loss),Gout,Lesch nyhan syndrome,Type 2 DM,Metabolic syndrome Causes of decreased levels-Low Zinc intake,OCP,Multiple Sclerosis

TOTAL PROTEIN, SERUM-is a biochemical test for measuring the total amount of protein in serum. Protein in the plasma is made up of albumin and globulin. Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenstroms disease Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage), Burns, Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic

syndrome,Protein-losing enteropathy etc. ALBUMIN, SERUM-Human serum albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. **Low blood albumin levels (hypoalbuminemia) can be caused by:** Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance,malnutrition and wasting etc.

Dr. Arpita Pasari, MD Consultant Pathologist



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V<u>iew Details</u>

View Report





PATIENT NAME: CHINTAN TRIVEDI (PKG10000289) EC 85778 REF. DOCTOR: DR. BANK OF BARODA

ABHA NO

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8800465156

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Test Report Status Results **Biological Reference Interval** Units <u>Final</u>

CLINICAL PATH - URINALYSIS

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

PHYSICAL EXAMINATION, URINE

COLOR PALE YELLOW

APPEARANCE CLEAR

CHEMICAL EXAMINATION, URINE

PH 5.5 4.7 - 7.51.003 - 1.035 SPECIFIC GRAVITY 1.010 **PROTEIN** NOT DETECTED NOT DETECTED **GLUCOSE** NOT DETECTED NOT DETECTED **KETONES** NOT DETECTED NOT DETECTED **BLOOD** NOT DETECTED NOT DETECTED **BILIRUBIN** NOT DETECTED NOT DETECTED UROBILINOGEN **NORMAL NORMAL NITRITE** NOT DETECTED NOT DETECTED LEUKOCYTE ESTERASE NOT DETECTED NOT DETECTED

MICROSCOPIC EXAMINATION, URINE

/HPF RED BLOOD CELLS NOT DETECTED NOT DETECTED /HPF PUS CELL (WBC'S) 2-3 0-5 EPITHELIAL CELLS 2-3 0-5 /HPF

NOT DETECTED **CASTS** CRYSTALS NOT DETECTED

BACTERIA NOT DETECTED NOT DETECTED YEAST NOT DETECTED NOT DETECTED

Please note that all the urinary findings are confirmed manually as well. **REMARKS**

Interpretation(s)

The following table describes the probable conditions, in which the analytes are present in urine

Presence of	Conditions
Proteins	Inflammation or immune illnesses
Pus (White Blood Cells)	Urinary tract infection, urinary tract or kidney stone, tumors or any kind
	of kidney impairment

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Test Report Status Final Results Biological Reference Interval Units

Glucose	Diabetes or kidney disease
Ketones	Diabetic ketoacidosis (DKA), starvation or thirst
Urobilinogen	Liver disease such as hepatitis or cirrhosis
Blood	Renal or genital disorders/trauma
Bilirubin	Liver disease
Erythrocytes	Urological diseases (e.g. kidney and bladder cancer, urolithiasis), urinary tract infection and glomerular diseases
Leukocytes	Urinary tract infection, glomerulonephritis, interstitial nephritis either acute or chronic, polycystic kidney disease, urolithiasis, contamination by genital secretions
Epithelial cells	Urolithiasis, bladder carcinoma or hydronephrosis, ureteric stents or bladder catheters for prolonged periods of time
Granular Casts	Low intratubular pH, high urine osmolality and sodium concentration, interaction with Bence-Jones protein
Hyaline casts	Physical stress, fever, dehydration, acute congestive heart failure, renal diseases
Calcium oxalate	Metabolic stone disease, primary or secondary hyperoxaluria, intravenous infusion of large doses of vitamin C, the use of vasodilator naftidrofuryl oxalate or the gastrointestinal lipase inhibitor orlistat, ingestion of ethylene glycol or of star fruit (Averrhoa carambola) or its juice
Uric acid	arthritis
Bacteria	Urinary infectionwhen present in significant numbers & with pus cells.
Trichomonas vaginalis	Vaginitis, cervicitis or salpingitis

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Test Report Status <u>Final</u> Results Biological Reference Interval Units

CLINICAL PATH - STOOL ANALYSIS

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

PHYSICAL EXAMINATION, STOOL

COLOUR BROWN

CONSISTENCY WELL FORMED

MUCUS ABSENT NOT DETECTED

VISIBLE BLOOD ABSENT ABSENT ABSENT

ADULT PARASITE NOT DETECTED

CHEMICAL EXAMINATION, STOOL

STOOL PH ALKALINE

OCCULT BLOOD NOT DETECTED NOT DETECTED

MICROSCOPIC EXAMINATION, STOOL

PUS CELLS 1-2 /hpf

RED BLOOD CELLS NOT DETECTED NOT DETECTED /HPF

CYSTS NOT DETECTED NOT DETECTED

OVA NOT DETECTED

LARVAE NOT DETECTED NOT DETECTED

TROPHOZOITES NOT DETECTED NOT DETECTED

FAT ABSENT

VEGETABLE CELLS ABSENT
CHARCOT LEYDEN CRYSTALS ABSENT

Interpretation(s)

Stool routine analysis is only a screening test for disorders of gastrointentestinal tract like infection, malabsorption, etc. The following table describes the probable conditions, in which the analytes are present in stool.

PRESENCE OF	CONDITION
Pus cells	Pus in the stool is an indication of infection
Red Blood cells	Parasitic or bacterial infection or an inflammatory bowel condition such as
	ulcerative colitis

Dr.Arpita Pasari, MD Consultant Pathologist - ment -

Dr.Meena Jinwah ,MBBS . MD Consultant Microbiologist





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AGE/SEX :41 Years Male

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Test Report Status Results **Biological Reference Interval** Units **Final**

Parasites	Infection of the digestive system. Stool examination for ova and parasite detects presence of parasitic infestation of gastrointestinal tract. Various forms of parasite that can be detected include cyst, trophozoite and larvae. One negative result does not rule out the possibility of parasitic infestation. Intermittent shedding of parasites warrants examinations of multiple specimens tested on consecutive days. Stool specimens for parasitic examination should be collected before initiation of antidiarrheal therapy or antiparasitic therapy. This test does not detect presence of opportunistic parasites like Cyclospora, Cryptosporidia and Isospora species. Examination of Ova and Parasite has been carried out by direct and concentration techniques.				
Mucus	Mucus is a protective layer that lubricates, protects& reduces damage due to bacteria or viruses.				
Charcot-Leyden crystal	Parasitic diseases.				
Ova & cyst	Ova & cyst indicate parasitic infestation of intestine.				
Frank blood	Bleeding in the rectum or colon.				
Occult blood	Occult blood indicates upper GI bleeding.				
Macrophages	Macrophages in stool are an indication of infection as they are protective cells.				
Epithelial cells	Epithelial cells that normally line the body surface and internal organs show up in stool when there is inflammation or infection.				
Fat	Increased fat in stool maybe seen in conditions like diarrhoea or malabsorption.				
pH	Normal stool pH is slightly acidic to neutral. Breast-fed babies generally has acidic stool.				

ADDITIONAL STOOL TESTS:

- Stool Culture: This test is done to find cause of GI infection, make decision about best treatment for GI infection & to find out if 1. treatment for GI infection worked.
- Fecal Calprotectin: It is a marker of intestinal inflammation. This test is done to differentiate Inflammatory Bowel Disease (IBD) 2. from Irritable Bowel Syndrome (IBS).
- 3. Fecal Occult Blood Test(FOBT): This test is done to screen for colon cancer & to evaluate possible cause of unexplained anaemia.
- 4. Clostridium Difficile Toxin Assay: This test is strongly recommended in healthcare associated bloody or waterydiarrhoea, due to overuse of broad spectrum antibiotics which alter the normal GI flora.
- Biofire (Film Array) GI PANEL: In patients of Diarrhoea, Dysentry, Rice watery Stool, FDA approved, Biofire Film Array Test, (Real Time Multiplex PCR) is strongly recommended as it identifies organisms, bacteria, fungi, virus , parasite and other opportunistic pathogens, Vibrio cholera infections only in 3 hours. Sensitivity 96% & Specificity 99%.
- Rota Virus Immunoassay: This test is recommended in severe gastroenteritis in infants & children associated with watery diarrhoea, vomitting& abdominal cramps. Adults are also affected. It is highly contagious in nature.

Dr. Arpita Pasari, MD **Consultant Pathologist** Mr. Oak

Dr. Meena Jinwah, MBBS. MD **Consultant Microbiologist**





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Units

PATIENT NAME: CHINTAN TRIVEDI (PKG10000289) EC 85778 REF. DOCTOR: DR. BANK OF BARODA

CODE/NAME & ADDRESS : C000138355
ARCOFEMI HEALTHCARE LTD (MEDIWHEEL

F-703, LADO SARAI, MEHRAULISOUTH WEST

DELHI

NEW DELHI 110030

Test Report Status

8800465156

ACCESSION NO : 0290WL001393

PATIENT ID : CHINM310882290

CLIENT PATIENT ID:

ABHA NO :

AGE/SEX :41 Years

DRAWN

Biological Reference Interval

RECEIVED : 09/12/2023 13:17:27 REPORTED :11/12/2023 16:24:52

SPECIALISED CHEMISTRY - HORMONE

Results

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

THYROID PANEL, SERUM

T3 127.60 80.0 - 200.0 ng/dL

METHOD: CHEMILUMINESCENCE TECHNOLOGY

Final

T4 7.31 5.10 - 14.10 $\mu g/dL$

METHOD: CHEMILUMINESCENCE TECHNOLOGY

TSH (ULTRASENSITIVE) 1.460 0.270 - 4.200 μIU/mL

METHOD: CHEMILUMINESCENCE TECHNOLOGY

Interpretation(s)

Triiodothyronine T3, Thyroxine T4, and Thyroid Stimulating Hormone TSH are thyroid hormones which affect almost every physiological process in the body, including growth, development, metabolism, body temperature, and heart rate.

Production of T3 and its prohormone thyroxine (T4) is activated by thyroid-stimulating hormone (TSH), which is released from the pituitary gland. Elevated concentrations of T3, and T4 in the blood inhibit the production of TSH.

Excessive secretion of thyroxine in the body is hyperthyroidism, and deficient secretion is called hypothyroidism.

In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hyperthyroidism, TSH levels are low. Below mentioned are the guidelines for Pregnancy related reference ranges for Total T4, TSH & Total T3. Measurement of the serum TT3 level is a more sensitive test for the diagnosis of hyperthyroidism, and measurement of TT4 is more useful in the diagnosis of hypothyroidism. Most of the thyroid hormone in blood is bound to transport proteins. Only a very small fraction of the circulating hormone is free and biologically active. It is advisable to detect Free T3, FreeT4 along with TSH, instead of testing for albumin bound Total T3, Total T4.

Sr. No.	TSH	Total T4	FT4	Total T3	Possible Conditions
1	High	Low	Low	Low	(1) Primary Hypothyroidism (2) Chronic autoimmune Thyroiditis (3)
					Post Thyroidectomy (4) Post Radio-Iodine treatment
2	High	Normal	Normal	Normal	(1)Subclinical Hypothyroidism (2) Patient with insufficient thyroid
					hormone replacement therapy (3) In cases of Autoimmune/Hashimoto
					thyroiditis (4). Isolated increase in TSH levels can be due to Subclinical
					inflammation, drugs like amphetamines, Iodine containing drug and
					dopamine antagonist e.g. domperidone and other physiological reasons.
3	Normal/Low	Low	Low	Low	(1) Secondary and Tertiary Hypothyroidism
4	Low	High	High	High	(1) Primary Hyperthyroidism (Graves Disease) (2) Multinodular Goitre
		-	-	-	(3)Toxic Nodular Goitre (4) Thyroiditis (5) Over treatment of thyroid
					hormone (6) Drug effect e.g. Glucocorticoids, dopamine, T4
					replacement therapy (7) First trimester of Pregnancy
5	Low	Normal	Normal	Normal	(1) Subclinical Hyperthyroidism
6	High	High	High	High	(1) TSH secreting pituitary adenoma (2) TRH secreting tumor
7	Low	Low	Low	Low	(1) Central Hypothyroidism (2) Euthyroid sick syndrome (3) Recent
					treatment for Hyperthyroidism

Dr.Arpita Pasari, MD Consultant Pathologist



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View Details

View Repor

PERFORMED AT :

Patient Ref. No. 775000005703035



REF. DOCTOR: DR. BANK OF BARODA PATIENT NAME: CHINTAN TRIVEDI (PKG10000289) EC 85778

CODE/NAME & ADDRESS: C000138355 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, LADO SARAI, MEHRAULISOUTH WEST

DELHI

NEW DELHI 110030 8800465156

ACCESSION NO: 0290WL001393

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AGE/SEX DRAWN

RECEIVED: 09/12/2023 13:17:27

:41 Years

REPORTED :11/12/2023 16:24:52

Test Report Status Results **Biological Reference Interval** Units **Final**

8	Normal/Low	Normal	Normal	High	(1) T3 thyrotoxicosis (2) Non-Thyroidal illness
9	Low	High	High	Normal	(1) T4 Ingestion (2) Thyroiditis (3) Interfering Anti TPO antibodies

REF: 1. TIETZ Fundamentals of Clinical chemistry 2.Guidlines of the American Thyroid association during pregnancy and Postpartum, 2011. NOTE: It is advisable to detect Free T3, FreeT4 along with TSH, instead of testing for albumin bound Total T3, Total T4.TSH is not affected by variation in thyroid - binding protein. TSH has a diurnal rhythm, with peaks at 2:00 - 4:00 a.m. And troughs at 5:00 - 6:00 p.m. With ultradian variations.

> **End Of Report** Please visit www.agilusdiagnostics.com for related Test Information for this accession

CONDITIONS OF LABORATORY TESTING & REPORTING

- 1. It is presumed that the test sample belongs to the patient named or identified in the test requisition form.
- 2. All tests are performed and reported as per the turnaround time stated in the AGILUS Directory of Services.
- 3. Result delays could occur due to unforeseen circumstances such as non-availability of kits / equipment breakdown / natural calamities / technical downtime or any other unforeseen event.
- 4. A requested test might not be performed if:
 - i. Specimen received is insufficient or inappropriate
 - ii. Specimen quality is unsatisfactory
 - iii. Incorrect specimen type
 - iv. Discrepancy between identification on specimen container label and test requisition form

- AGILUS Diagnostics confirms that all tests have been performed or assayed with highest quality standards, clinical safety & technical integrity.
- 6. Laboratory results should not be interpreted in isolation; it must be correlated with clinical information and be interpreted by registered medical practitioners only to determine final diagnosis.
- 7. Test results may vary based on time of collection, physiological condition of the patient, current medication or nutritional and dietary changes. Please consult your doctor or call us for any clarification.
- 8. Test results cannot be used for Medico legal purposes.
- 9. In case of queries please call customer care (91115 91115) within 48 hours of the report.

Agilus Diagnostics Ltd

Fortis Hospital, Sector 62, Phase VIII, Mohali 160062

Dr. Arpita Pasari, MD **Consultant Pathologist**





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