

PATIENT NAME : JAIMINI PRANAVKUMAR RANA

REF. DOCTOR : SELF

CODE/NAME &amp; ADDRESS : C000138364

ARCOFEMI HEALTHCARE LTD (MEDIWHEEL  
F-703, LADO SARAI, MEHRAULISOUTH WEST  
DELHI  
NEW DELHI 110030  
8800465156

ACCESSION NO : 0321WG001444

PATIENT ID : JAIMF281195321

CLIENT PATIENT ID:

ABHA NO :

AGE/SEX : 27 Years Female

DRAWN :

RECEIVED : 08/07/2023 08:53:14

REPORTED : 10/07/2023 17:59:09

Test Report Status **Final**

Results

Biological Reference Interval Units

**MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE****XRAY-CHEST**

IMPRESSION

NO ABNORMALITY DETECTED

**ECG**

ECG

NORMAL SINUS RHYTHM

**MEDICAL HISTORY**

RELEVANT PRESENT HISTORY

C/O LEFT KNEE PAIN AND BACK PAIN FOR 1 MONTH

RELEVANT PAST HISTORY

COLD ON TREATMENT SINCE 1 WEEK

P/H/O JAUNDICE DURING CHILDHOOD

RELEVANT PERSONAL HISTORY

NOT SIGNIFICANT

MENSTRUAL HISTORY (FOR FEMALES)

REGULAR

LMP (FOR FEMALES)

21/06/2023

RELEVANT FAMILY HISTORY

DIABETES

OCCUPATIONAL HISTORY

NOT SIGNIFICANT

HISTORY OF MEDICATIONS

NOT SIGNIFICANT

**ANTHROPOMETRIC DATA & BMI**

HEIGHT IN METERS

1.64

mts

WEIGHT IN KGS.

62.1

Kgs

BMI

23

BMI &amp; Weight Status as follows

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

PHYSICAL ATTITUDE

NORMAL

GENERAL APPEARANCE / NUTRITIONAL STATUS

HEALTHY

BUILT / SKELETAL FRAMEWORK

AVERAGE

FACIAL APPEARANCE

NORMAL

SKIN

NORMAL

UPPER LIMB

NORMAL

LOWER LIMB

NORMAL

NECK

NORMAL

P. V. Kapadia

Jinal

Dr. Priyank Kapadia  
PhysicianDr. Jinal kamodia  
Consultant Radiology

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NECK LYMPHATICS / SALIVARY GLANDS

NOT ENLARGED OR TENDER

THYROID GLAND

NOT ENLARGED

TEMPERATURE

NORMAL

PULSE

70/MIN

RESPIRATORY RATE

NORMAL

**CARDIOVASCULAR SYSTEM**

BP

122/84 MM HG  
(SITTING)

mm/Hg

PERICARDIUM

NORMAL

APEX BEAT

NORMAL

HEART SOUNDS

S1, S2 HEARD NORMALLY

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

LIVER

NOT PALPABLE

SPLEEN

NOT PALPABLE

**CENTRAL NERVOUS SYSTEM**

HIGHER FUNCTIONS

NORMAL

CRANIAL NERVES

NORMAL

CEREBELLAR FUNCTIONS

NORMAL

SENSORY SYSTEM

NORMAL

MOTOR SYSTEM

NORMAL

REFLEXES

NORMAL

**MUSCULOSKELETAL SYSTEM**

SPINE

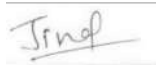
NORMAL

JOINTS

NORMAL

**BASIC EYE EXAMINATION**


Dr. Priyank Kapadia  
Physician



Dr. Jinal kamodia  
Consultant Radiology

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DISTANT VISION RIGHT EYE WITH GLASSES	WITH GLASSES NORMAL
DISTANT VISION LEFT EYE WITH GLASSES	WITH GLASSES NORMAL
NEAR VISION RIGHT EYE WITHOUT GLASSES	WITHIN NORMAL LIMIT
NEAR VISION LEFT EYE WITHOUT GLASSES	WITHIN NORMAL LIMIT
COLOUR VISION	NORMAL

**SUMMARY**

RELEVANT HISTORY	NOT SIGNIFICANT
RELEVANT GP EXAMINATION FINDINGS	NOT SIGNIFICANT
RELEVANT LAB INVESTIGATIONS	ESR:- HIGH
RELEVANT NON PATHOLOGY DIAGNOSTICS	NO ABNORMALITIES DETECTED
REMARKS / RECOMMENDATIONS	ESR:- HIGH
	ADV:- PHYSICIAN OPINION

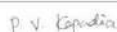
**Comments**

OUR PANEL DOCTORS FOR NON-PATHOLOGY TESTS:-

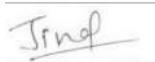
CHECK UP DONE BY:- DR. NAMRATA AGRAWAL (M.B.B.S)

REPORT REVIEWED BY:- DR. PRIYANK KAPADIYA (M.B.B.S DNB MEDICINE)

RADIOLOGIST:- DR. SAHIL N SHAH (M.D.RADIOLOGY) / DR. J. S. KAMODIA (M. D. RADIOLOGY)



**Dr.Priyank Kapadia**  
Physician



**Dr.Jinal kamodia**  
Consultant Radiology

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**Results**

**Units**

**MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE**

**ULTRASOUND ABDOMEN**

**ULTRASOUND ABDOMEN**

NO ABNORMALITIES DETECTED

**TMT OR ECHO**

**TMT OR ECHO**

2D ECHO:-

- 1) NORMAL CHAMBERS AND VALVES.
- 2) GOOD LV SYSTOLIC FUNCTION. LVEF 60%. NO RWMA AT REST.
- 3) NO MR, AR, TR.
- 4) NORMAL LV COMPLIANCE.
- 5) NO PAH.
- 6) NO LV CLOT, VEGETATION OR PERICARDIAL EFFUSION.
- 7) IAS/IVS INTACT.

**Interpretation(s)**

MEDICAL

HISTORY.\*\*\*\*\*

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

\*\*\*\*\*

*P. V. Kapadia*

**Dr. Priyank Kapadia  
Physician**

*Jinal*

**Dr. Jinal kamodia  
Consultant Radiology**



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## HAEMATOLOGY - CBC

## MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE

## BLOOD COUNTS, EDTA WHOLE BLOOD

Parameter	Result	Reference Interval	Units
HEMOGLOBIN (HB) METHOD : PHOTOMETRIC MEASUREMENT	11.8 Low	12.0 - 15.0	g/dL
RED BLOOD CELL (RBC) COUNT METHOD : COULTER PRINCIPLE	4.19	3.8 - 4.8	mil/ $\mu$ L
WHITE BLOOD CELL (WBC) COUNT METHOD : COULTER PRINCIPLE	6.35	4.0 - 10.0	thou/ $\mu$ L
PLATELET COUNT METHOD : COULTER PRINCIPLE	417 High	150 - 410	thou/ $\mu$ L

## RBC AND PLATELET INDICES

HEMATOCRIT (PCV) METHOD : CALCULATED	36.3	36.0 - 46.0	%
MEAN CORPUSCULAR VOLUME (MCV) METHOD : DERIVED PARAMETER FROM RBC HISTOGRAM	86.7	83.0 - 101.0	fL
MEAN CORPUSCULAR HEMOGLOBIN (MCH) METHOD : CALCULATED	28.2	27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC) METHOD : CALCULATED	32.6	31.5 - 34.5	g/dL
RED CELL DISTRIBUTION WIDTH (RDW) METHOD : DERIVED PARAMETER FROM RBC HISTOGRAM	14.5 High	11.6 - 14.0	%
MENTZER INDEX METHOD : CALCULATED PARAMETER	20.7		
MEAN PLATELET VOLUME (MPV) METHOD : DERIVED PARAMETER FROM PLATELET HISTOGRAM	8.3	6.8 - 10.9	fL

## WBC DIFFERENTIAL COUNT

NEUTROPHILS METHOD : OPTICAL IMPEDENCE & MICROSCOPY	61	40 - 80	%
LYMPHOCYTES METHOD : OPTICAL IMPEDENCE & MICROSCOPY	28	20 - 40	%
MONOCYTES METHOD : OPTICAL IMPEDENCE & MICROSCOPY	9	2.0 - 10.0	%
EOSINOPHILS	1	1.0 - 6.0	%



**Dr. Miral Gajera**  
Consultant Pathologist

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Test Report Status	Final	Results	Biological Reference Interval	Units
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METHOD : OPTICAL IMPEDENCE & MICROSCOPY

BASOPHILS	1	0 - 1	%
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METHOD : IMPEDANCE

ABSOLUTE NEUTROPHIL COUNT	3.87	2.0 - 7.0	thou/μL
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METHOD : CALCULATED

ABSOLUTE LYMPHOCYTE COUNT	1.78	1.0 - 3.0	thou/μL
---------------------------	------	-----------	---------

METHOD : CALCULATED PARAMETER

ABSOLUTE MONOCYTE COUNT	0.57	0.2 - 1.0	thou/μL
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METHOD : CALCULATED PARAMETER

ABSOLUTE EOSINOPHIL COUNT	0.06	0.02 - 0.50	thou/μL
---------------------------	------	-------------	---------

METHOD : CALCULATED

ABSOLUTE BASOPHIL COUNT	0.06	0.02 - 0.10	thou/μL
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METHOD : CALCULATED

NEUTROPHIL LYMPHOCYTE RATIO (NLR)	2.2		
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METHOD : CALCULATED PARAMETER

**MORPHOLOGY**

RBC	NORMOCYTIC NORMOCHROMIC
-----	-------------------------

METHOD : MICROSCOPIC EXAMINATION

WBC	NORMAL MORPHOLOGY
-----	-------------------

METHOD : MICROSCOPIC EXAMINATION

PLATELETS	ADEQUATE
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METHOD : MICROSCOPIC EXAMINATION

REMARKS	NO PREMATURE CELLS ARE SEEN. MALARIAL PARASITE NOT DETECTED.
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METHOD : MICROSCOPIC EXAMINATION

**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 < 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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## HAEMATOLOGY

## MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE

## ERYTHROCYTE SEDIMENTATION RATE (ESR),WHOLE BLOOD

E.S.R	35 High	0 - 20	mm at 1 hr
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METHOD : WESTERGREN METHOD

## GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD

HBA1C	5.1	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 Therapeutic goals: < 7.0 Action suggested : > 8.0 (ADA Guideline 2021)	%
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METHOD : HPLC

ESTIMATED AVERAGE GLUCOSE(EAG)	99.7	< 116.0	mg/dL
--------------------------------	------	---------	-------

## Interpretation(s)

## ERYTHROCYTE SEDIMENTATION RATE (ESR),WHOLE 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, Vasculitias, 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: Polycythemia 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)

## REFERENCE :

1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition 2. Paediatric reference intervals. AACCC 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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Consultant Pathologist

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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.
2. eAG gives an evaluation of blood glucose levels for the last couple of months.
3. 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 addition 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 platform (Boronate affinity chromatography) is recommended for testing of HbA1c. Abnormal Hemoglobin electrophoresis (HPLC method) is recommended for detecting a hemoglobinopathy



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ACCESSION NO : **0321WG001444**

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### IMMUNOHAEMATOLOGY

#### MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE

#### ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP

TYPE O

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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**AGE/SEX : 27 Years Female**

**DRAWN :**

**RECEIVED : 08/07/2023 08:53:14**

**REPORTED : 10/07/2023 17:59:09**

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

**MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE**

**GLUCOSE FASTING,FLUORIDE PLASMA**

FBS (FASTING BLOOD SUGAR)	91	74 - 99	mg/dL
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METHOD : HEXOKINASE

**GLUCOSE, POST-PRANDIAL, PLASMA**

PPBS(POST PRANDIAL BLOOD SUGAR)	105	70 - 140	mg/dL
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METHOD : HEXOKINASE

**LIPID PROFILE, SERUM**

CHOLESTEROL, TOTAL	164	Desirable: < 200 BorderlineHigh: 200 - 239 High: > or = 240	mg/dL
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METHOD : ENZYMATIC, COLORIMETRIC

TRIGLYCERIDES	73	Desirable: < 150 BorderlineHigh: 150 - 199 High: 200 - 499 Very High: > or = 500	mg/dL
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METHOD : ENZYMATIC, COLORIMETRIC

HDL CHOLESTEROL	49	< 40 Low > or = 60 High	mg/dL
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CHOLESTEROL LDL	100	Adult levels: Optimal < 100 Near optimal/above optimal: 100-129 Borderline high : 130-159 High : 160-189 Very high : = 190	mg/dL
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NON HDL CHOLESTEROL	115	Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL
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VERY LOW DENSITY LIPOPROTEIN	14.6	< or = 30	mg/dL
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CHOL/HDL RATIO	3.4	3.3 - 4.4	
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LDL/HDL RATIO	<b>-1.3 Low</b>	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	
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Consultant Pathologist



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Ahmedabad, 380015  
Gujrat, India  
Tel : 079-48912999,079-48913999,079-48914999  
Email : customercare.ahmedabad@agilus.in



**Patient Ref. No. 775000003830037**

PATIENT NAME : JAIMINI PRANAVKUMAR RANA

REF. DOCTOR : SELF

CODE/NAME &amp; ADDRESS : C000138364

ARCOFEMI HEALTHCARE LTD (MEDIWHEEL  
F-703, LADO SARAI, MEHRAULISOUTH WEST  
DELHI  
NEW DELHI 110030  
8800465156

ACCESSION NO : 0321WG001444

PATIENT ID : JAIMF281195321

CLIENT PATIENT ID:

ABHA NO :

AGE/SEX : 27 Years Female

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Test Report Status **Final**

Results

Biological Reference Interval Units

**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**

Risk Category	
Extreme risk group	A.CAD with > 1 feature of high risk group
	B. CAD with > 1 feature of Very high risk 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 Hypercholesterolemia
High Risk	1. Three major ASCVD risk factors. 2. Diabetes with 1 major risk factor or no evidence of end organ damage. 3. CKD stage 3B or 4. 4. LDL >190 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 (Atherosclerotic cardiovascular disease) Risk Factors	
1. Age > or = 45 years in males and > or = 55 years in females	3. Current Cigarette smoking or tobacco use
2. 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 < OR = 30)	< 80 (Optional goal <OR = 60)	>OR = 50	>OR = 80
Extreme Risk Group Category B	<OR = 30	<OR = 60	> 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	0.62	Upto 1.2	mg/dL
BILIRUBIN, DIRECT	<b>0.23 High</b>	Upto 0.2	mg/dL
METHOD : DIAZO COLORIMETRIC			
BILIRUBIN, INDIRECT	0.39	0.00 - 1.00	mg/dL
TOTAL PROTEIN	7.1	6.4 - 8.3	g/dL
METHOD : COLORIMETRIC			
ALBUMIN	3.9	3.5 - 5.2	g/dL
METHOD : BROMOCRESOL GREEN			



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Consultant Pathologist

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GLOBULIN	3.2	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO	1.2	1.0 - 2.0	RATIO
ASPARTATE AMINOTRANSFERASE(AST/SGOT) METHOD : IFCC WITHOUT PYRIDOXAL PHOSPHATE	15	0 - 32	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT) METHOD : IFCC WITHOUT PYRIDOXAL PHOSPHATE	6	0 - 33	U/L
ALKALINE PHOSPHATASE METHOD : COLORIMETRIC	<b>109 High</b>	35 - 104	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT) METHOD : ENZYMATIC, COLORIMETRIC	13	5 - 36	U/L
LACTATE DEHYDROGENASE METHOD : UV ASSAY METHOD	184	135 - 214	U/L
<b>BLOOD UREA NITROGEN (BUN), SERUM</b>			
BLOOD UREA NITROGEN	7	6 - 20	mg/dL
<b>CREATININE, SERUM</b>			
CREATININE METHOD : JAFFE ALKALINE PICRATE	<b>0.51 Low</b>	0.60 - 1.10	mg/dL
<b>BUN/CREAT RATIO</b>			
BUN/CREAT RATIO	13.73	5.0 - 15.0	
<b>URIC ACID, SERUM</b>			
URIC ACID	3.8	2.4 - 5.7	mg/dL
<b>TOTAL PROTEIN, SERUM</b>			
TOTAL PROTEIN METHOD : COLORIMETRIC	7.1	6.4 - 8.3	g/dL
<b>ALBUMIN, SERUM</b>			
ALBUMIN METHOD : BROMOCRESOL GREEN	3.9	3.5 - 5.2	g/dL
<b>GLOBULIN</b>			
GLOBULIN	3.2	2.0 - 4.1	g/dL
<b>ELECTROLYTES (NA/K/CL), SERUM</b>			
SODIUM, SERUM METHOD : ISE	138.5	136 - 145	mmol/L
POTASSIUM, SERUM METHOD : ISE	4.50	3.3 - 5.1	mmol/L

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Consultant Pathologist



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 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL  
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**CHLORIDE, SERUM**      **104.8**      **98 - 106**      **mmol/L**

METHOD : ION SELECTIVE ELECTRODE TECHNOLOGY

**Interpretation(s)**

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

**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 urine.

**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 glycemc control.

High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glycosuria,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 Glycosuria, 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

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**Consultant Pathologist**



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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, ischemia 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 in Hypophosphatasia, Malnutrition, Protein deficiency, Wilsons disease.

**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, biliary 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.

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Consultant Pathologist



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**CLINICAL PATH - URINALYSIS**

**MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE**

**PHYSICAL EXAMINATION, URINE**

COLOR Yellow  
APPEARANCE Clear

**CHEMICAL EXAMINATION, URINE**

PH	6.0	4.7 - 7.5	
METHOD : REFLECTANCE SPECTROPHOTOMETRY			
SPECIFIC GRAVITY	1.015	1.003 - 1.035	
METHOD : REFLECTANCE SPECTROPHOTOMETRY			
PROTEIN	NOT DETECTED	NEGATIVE	
METHOD : REFLECTANCE SPECTROPHOTOMETRY			
GLUCOSE	NOT DETECTED	NOT DETECTED	
METHOD : REFLECTANCE SPECTROPHOTOMETRY			
KETONES	NOT DETECTED	NOT DETECTED	
METHOD : REFLECTANCE SPECTROPHOTOMETRY			
BLOOD	NOT DETECTED	NEGATIVE	
METHOD : REFLECTANCE SPECTROPHOTOMETRY			
BILIRUBIN	NOT DETECTED	NOT DETECTED	
METHOD : REFLECTANCE SPECTROPHOTOMETRY			
UROBILINOGEN	NORMAL	NORMAL	
METHOD : REFLECTANCE SPECTROPHOTOMETRY			
NITRITE	NOT DETECTED	NOT DETECTED	
METHOD : REFLECTANCE SPECTROPHOTOMETRY			
LEUKOCYTE ESTERASE	NOT DETECTED	NOT DETECTED	
METHOD : REFLECTANCE SPECTROPHOTOMETRY			

**MICROSCOPIC EXAMINATION, URINE**

RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
METHOD : MICROSCOPIC EXAMINATION			
PUS CELL (WBC'S)	0-1	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION			
EPITHELIAL CELLS	2-3	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION			
CASTS	NOT DETECTED		
METHOD : MICROSCOPIC EXAMINATION			

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**CRYSTALS**

METHOD : MICROSCOPIC EXAMINATION

NOT DETECTED

**BACTERIA**

METHOD : MICROSCOPIC EXAMINATION

NOT DETECTED

NOT DETECTED

**YEAST**

METHOD : MICROSCOPIC EXAMINATION

NOT DETECTED

NOT DETECTED

**REMARKS**

MICROSCOPIC EXAMINATION OF URINE IS CARRIED OUT ON CENTRIFUGED URINARY SEDIMENT.

**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
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

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CODE/NAME &amp; ADDRESS : C000138364

ARCOFEMI HEALTHCARE LTD (MEDIWHEEL  
F-703, LADO SARAI, MEHRAULISOUTH WEST  
DELHI  
NEW DELHI 110030  
8800465156

ACCESSION NO : **0321WG001444**

PATIENT ID : JAIMF281195321

CLIENT PATIENT ID:

ABHA NO :

AGE/SEX : 27 Years Female

DRAWN :

RECEIVED : 08/07/2023 08:53:14

REPORTED : 10/07/2023 17:59:09

Test Report Status	Final	Results	Biological Reference Interval	Units
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Bacteria	Urinary infection when present in significant numbers & with pus cells.
Trichomonas vaginalis	Vaginitis, cervicitis or salpingitis



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Consultant Pathologist

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Email : customercare.ahmedabad@agilus.in



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Test Report Status **Final**

Results

Biological Reference Interval Units

## CYTOLOGY

**MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE****PAPANICOLAOU SMEAR**

TEST METHOD

CONVENTIONAL GYNEC CYTOLOGY

SPECIMEN TYPE

TWO UNSTAINED CERVICAL SMEARS RECEIVED

REPORTING SYSTEM

2014 BETHESDA SYSTEM FOR REPORTING CERVICAL CYTOLOGY

SPECIMEN ADEQUACY

SATISFACTORY

MICROSCOPY

SMEARS SHOW PREDOMINANTLY SUPERFICIAL AND INTERMEDIATE  
SQUAMOUS CELLS AGAINST BACKGROUND OF MILD ACUTE  
INFLAMMATION. ENDOCERVICAL CELLS NOT SEEN ON SMEAR. NO  
EVIDENCE OF DYSPLASIA OR MALIGNANT CELLS SEEN.

INTERPRETATION / RESULT

NEGATIVE FOR INTRAEPITHELIAL LESION OR MALIGNANCY

**Comments**

PAP SMEAR IS ASCRENING PROCEDURE FOR CERVICAL CANCER WITH INHERENT FALSE NEGATIVE RESULTS HENCE RESULTS SHOULD BE INTERPRETED WITH CAUTION.



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**CLINICAL PATH - STOOL ANALYSIS**

**MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE**

**PHYSICAL EXAMINATION,STOOL**

COLOUR	BROWN	
CONSISTENCY	WELL FORMED	
MUCUS	NOT DETECTED	NOT DETECTED
VISIBLE BLOOD	ABSENT	ABSENT
ADULT PARASITE	NOT DETECTED	

METHOD : MICROSCOPIC EXAMINATION

**CHEMICAL EXAMINATION,STOOL**

STOOL PH	NEGATIVE	
OCCULT BLOOD	NOT DETECTED	NOT DETECTED

METHOD : HEMOSPOT

**MICROSCOPIC EXAMINATION,STOOL**

PUS CELLS	NOT DETECTED		/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		

METHOD : MICROSCOPIC EXAMINATION

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METHOD : MICROSCOPIC EXAMINATION

**Interpretation(s)**

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

PRESENCE OF	CONDITION
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<b>Pus cells</b>	Pus in the stool is an indication of infection
<b>Red Blood cells</b>	Parasitic or bacterial infection or an inflammatory bowel condition such as ulcerative colitis
<b>Parasites</b>	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.
<b>Mucus</b>	Mucus is a protective layer that lubricates, protects & reduces damage due to bacteria or viruses.
<b>Charcot-Leyden crystal</b>	Parasitic diseases.
<b>Ova &amp; cyst</b>	Ova & cyst indicate parasitic infestation of intestine.
<b>Frank blood</b>	Bleeding in the rectum or colon.
<b>Occult blood</b>	Occult blood indicates upper GI bleeding.
<b>Macrophages</b>	Macrophages in stool are an indication of infection as they are protective cells.
<b>Epithelial cells</b>	Epithelial cells that normally line the body surface and internal organs show up in stool when there is inflammation or infection.
<b>Fat</b>	Increased fat in stool maybe seen in conditions like diarrhoea or malabsorption.
<b>pH</b>	Normal stool pH is slightly acidic to neutral. Breast-fed babies generally have an acidic stool.

**ADDITIONAL STOOL TESTS :**

- 1. Stool Culture:-** This test is done to find cause of GI infection, make decision about best treatment for GI infection & to find out if treatment for GI infection worked.
- 2. Fecal Calprotectin:** It is a marker of intestinal inflammation. This test is done to differentiate Inflammatory Bowel Disease (IBD) 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 watery diarrhoea, due to overuse of broad spectrum antibiotics which alter the normal GI flora.
- 5. Biofire (Film Array) GI PANEL:** In patients of Diarrhoea, Dysentery, 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%.
- 6. Rota Virus Immunoassay:** This test is recommended in severe gastroenteritis in infants & children associated with watery diarrhoea, vomiting & abdominal cramps. Adults are also affected. It is highly contagious in nature.

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**SPECIALISED CHEMISTRY - HORMONE**

**MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE**

**THYROID PANEL, SERUM**

T3	165.40	Non-Pregnant Women 80.0 - 200.0 Pregnant Women 1st Trimester:105.0 - 230.0 2nd Trimester:129.0 - 262.0 3rd Trimester:135.0 - 262.0	ng/dL
METHOD : ECLIA			
T4	11.41	Non-Pregnant Women 5.10 - 14.10 Pregnant Women 1st Trimester: 7.33 - 14.80 2nd Trimester: 7.93 - 16.10 3rd Trimester: 6.95 - 15.70	µg/dL
METHOD : ECLIA			
TSH (ULTRASENSITIVE)	1.410	Non Pregnant Women 0.27 - 4.20 Pregnant Women 1st Trimester: 0.33 - 4.59 2nd Trimester: 0.35 - 4.10 3rd Trimester: 0.21 - 3.15	µIU/mL
METHOD : ECLIA			

**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
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Consultant Pathologist



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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
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\*\***

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Results

Biological Reference Interval Units

**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
5. 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.

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