

**PATIENT NAME : HEMLATA JOSHI****REF. DOCTOR : SELF****CODE/NAME & ADDRESS :** C000138394ARCOFEMI HEALTHCARE LTD (MEDIWHEEL  
F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST  
DELHI  
NEW DELHI 110030  
8800465156**ACCESSION NO :** 0181WL001806**PATIENT ID :** HEMLF200387181**CLIENT PATIENT ID:****ABHA NO :****AGE/SEX :** 36 Years Female**DRAWN :****RECEIVED :** 23/12/2023 10:42:50**REPORTED :** 28/12/2023 16:07:44**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

WITHIN NORMAL LIMITS

**MEDICAL HISTORY**

RELEVANT PRESENT HISTORY

NOT SIGNIFICANT

RELEVANT PAST HISTORY

NOT SIGNIFICANT

RELEVANT PERSONAL HISTORY

MARRIED / VEG DIET / NO ALLERGIES / NO SMOKING / NO ALCOHOL.

MENSTRUAL HISTORY (FOR FEMALES)

REGULAR

LMP (FOR FEMALES)

03/12/2023.

RELEVANT FAMILY HISTORY

NOT SIGNIFICANT

HISTORY OF MEDICATIONS

NOT SIGNIFICANT

**ANTHROPOMETRIC DATA & BMI**

HEIGHT IN METERS

1.58

mts

WEIGHT IN KGS.

62

Kgs

BMI

25

BMI &amp; 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

PHYSICAL ATTITUDE

NORMAL



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S.K. Tower, Hari Niwas, Lbs Marg  
Thane, 400602  
Maharashtra, India  
Tel : 9111591115, Fax : CIN - U74899PB1995PLC045956  
Email : customercare.thane@agilus.in**Patient Ref. No. 77500005852721**

**PATIENT NAME : HEMLATA JOSHI**

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GENERAL APPEARANCE / NUTRITIONAL STATUS	HEALTHY			
BUILT / SKELETAL FRAMEWORK	AVERAGE			
FACIAL APPEARANCE	NORMAL			
SKIN	NORMAL			
UPPER LIMB	NORMAL			
LOWER LIMB	NORMAL			
NECK	NORMAL			
NECK LYMPHATICS / SALIVARY GLANDS	NOT ENLARGED OR TENDER			
THYROID GLAND	NOT ENLARGED			
CAROTID PULSATION	NORMAL			
TEMPERATURE	NORMAL			
PULSE	84/MIN.REGULAR, ALL PERIPHERAL PULSES WELL FELT, NO CAROTID BRUIT			
RESPIRATORY RATE	NORMAL			

**CARDIOVASCULAR SYSTEM**

BP	133/71 MM HG (SUPINE)	mm/Hg
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



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**PER ABDOMEN**

APPEARANCE	NORMAL
VENOUS PROMINENCE	ABSENT
LIVER	NOT PALPABLE
SPLEEN	NOT PALPABLE
HERNIA	ABSENT

**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**

CONJUNCTIVA	NORMAL
EYELIDS	NORMAL
EYE MOVEMENTS	NORMAL
CORNEA	NORMAL
DISTANT VISION RIGHT EYE WITHOUT GLASSES	WITHIN NORMAL LIMIT



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DISTANT VISION LEFT EYE WITHOUT GLASSES	WITHIN NORMAL LIMIT
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
REMARKS / RECOMMENDATIONS	DRINK 2-3 LITER WATER DAILY. REPEAT URINE ROUTINE AFTER 15 DAYS. LOW FAT,LOW CALORIE, LOW CARBOHYDRATE, HIGH FIBRE DIET. REGULAR EXERCISE.REGULAR WALK FOR 30-40 MIN DAILY. REPEAT LIPID PROFILE AFTER 3 MONTHS OF DIET AND EXERCISE.



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

**TMT OR ECHO**

**CLINICAL PROFILE**

2D ECHO : NORMAL

**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.

\*\*\*\*\*

**\*\*End Of Report\*\***

Please visit [www.agilusdiagnostics.com](http://www.agilusdiagnostics.com) for related Test Information for this accession

**CONDITIONS OF LABORATORY TESTING & REPORTING**

- |  |  |
|--|--|
| <ol style="list-style-type: none"> <li>1. It is presumed that the test sample belongs to the patient named or identified in the test requisition form.</li> <li>2. All tests are performed and reported as per the turnaround time stated in the AGILUS Directory of Services.</li> <li>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.</li> <li>4. A requested test might not be performed if:                     <ol style="list-style-type: none"> <li>i. Specimen received is insufficient or inappropriate</li> <li>ii. Specimen quality is unsatisfactory</li> <li>iii. Incorrect specimen type</li> <li>iv. Discrepancy between identification on specimen container label and test requisition form</li> </ol> </li> </ol> | <ol style="list-style-type: none"> <li>5. AGILUS Diagnostics confirms that all tests have been performed or assayed with highest quality standards, clinical safety &amp; technical integrity.</li> <li>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.</li> <li>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.</li> <li>8. Test results cannot be used for Medico legal purposes.</li> <li>9. In case of queries please call customer care (91115 91115) within 48 hours of the report.</li> </ol> |
|--|--|

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 Fortis Hospital, Sector 62, Phase VIII,  
 Mohali 160062



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Results

Biological Reference Interval Units

## HAEMATOLOGY - CBC

**MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE****BLOOD COUNTS, EDTA WHOLE BLOOD**

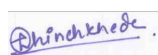
Test Name	Result	Biological Reference Interval	Units
HEMOGLOBIN (HB) METHOD : SLS- HEMOGLOBIN DETECTION METHOD	12.0	12.0 - 15.0	g/dL
RED BLOOD CELL (RBC) COUNT METHOD : HYDRODYNAMIC FOCUSING BY DC DETECTION	4.21	3.8 - 4.8	mil/ $\mu$ L
WHITE BLOOD CELL (WBC) COUNT METHOD : FLUORESCENCE FLOW CYTOMETRY	6.53	4.0 - 10.0	thou/ $\mu$ L
PLATELET COUNT METHOD : HYDRODYNAMIC FOCUSING BY DC DETECTION	358	150 - 410	thou/ $\mu$ L

**RBC AND PLATELET INDICES**

Test Name	Result	Biological Reference Interval	Units
HEMATOCRIT (PCV) METHOD : CUMULATIVE PULSE HEIGHT DETECTION METHOD	38.8	36.0 - 46.0	%
MEAN CORPUSCULAR VOLUME (MCV) METHOD : CALCULATED FROM RBC & HCT	92.2	83.0 - 101.0	fL
MEAN CORPUSCULAR HEMOGLOBIN (MCH) METHOD : CALCULATED FROM THE RBC & HGB	28.5	27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC) METHOD : CALCULATED FROM THE HGB & HCT	<b>30.9 Low</b>	31.5 - 34.5	g/dL
RED CELL DISTRIBUTION WIDTH (RDW) METHOD : CALCULATED FROM RBC SIZE DISTRIBUTION CURVE	<b>14.4 High</b>	11.6 - 14.0	%
MENTZER INDEX	21.9		
MEAN PLATELET VOLUME (MPV) METHOD : CALCULATED FROM PLATELET COUNT & PLATELET HEMATOCRIT	<b>11.0 High</b>	6.8 - 10.9	fL

**WBC DIFFERENTIAL COUNT**

Test Name	Result	Biological Reference Interval	Units
NEUTROPHILS METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING	54	40 - 80	%
LYMPHOCYTES METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING	32	20 - 40	%
MONOCYTES	9	2 - 10	%



**Dr. Priyal Chinchkhede, MD**  
Consultant Pathologist

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METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING <b>EOSINOPHILS</b>	5	1 - 6	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING <b>BASOPHILS</b>	0	0 - 1	%
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING <b>ABSOLUTE NEUTROPHIL COUNT</b>	3.53	2.0 - 7.0	thou/ $\mu$ L
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING <b>ABSOLUTE LYMPHOCYTE COUNT</b>	2.09	1.0 - 3.0	thou/ $\mu$ L
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING <b>ABSOLUTE MONOCYTE COUNT</b>	0.56	0.2 - 1.0	thou/ $\mu$ L
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING <b>ABSOLUTE EOSINOPHIL COUNT</b>	0.33	0.02 - 0.50	thou/ $\mu$ L
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING <b>ABSOLUTE BASOPHIL COUNT</b>	<b>0 Low</b>	0.02 - 0.10	thou/ $\mu$ L
METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING <b>NEUTROPHIL LYMPHOCYTE RATIO (NLR)</b>	1.7		

**MORPHOLOGY**

RBC	NORMOCYTIC NORMOCHROMIC
WBC	NORMAL MORPHOLOGY
METHOD : MICROSCOPIC EXAMINATION	
PLATELETS	ADEQUATE

**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.

**Dr. Priyal Chinchkhede, MD**  
**Consultant Pathologist**



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### HAEMATOLOGY

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

#### ERYTHROCYTE SEDIMENTATION RATE (ESR),EDTA BLOOD

E.S.R	5	0 - 20	mm
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METHOD : MODIFIED WESTERGRN

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

HBA1C	5.4	Non-diabetic Adult < 5.7 Pre-diabetes 5.7 - 6.4 Diabetes diagnosis: > or = 6.5 Therapeutic goals: < 7.0 Action suggested : > 8.0 (ADA Guideline 2021)	%
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METHOD : HPLC

ESTIMATED AVERAGE GLUCOSE(EAG)	108.3	< 116.0	mg/dL
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METHOD : CALCULATED PARAMETER

#### 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, Vasculitides, 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.

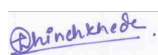
**Decreased** in: Polycythemia vera, Sickle cell anemia

##### 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)



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**REFERENCE :**

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).

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 patient's metabolic control has remained continuously within the target range.

1. eAG (Estimated average glucose) converts percentage HbA1c to mg/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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### IMMUNOHAEMATOLOGY

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

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

ABO GROUP

TYPE O

METHOD : GEL COLUMN AGGLUTINATION METHOD.

RH TYPE

POSITIVE

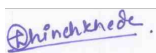
METHOD : GEL COLUMN AGGLUTINATION METHOD.

#### **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.



**Dr. Priyal Chinchkhede, MD**  
Consultant Pathologist

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Agilus Diagnostics Ltd.  
Mulund Goregoan Link Road  
Mumbai, 400078  
Maharashtra, India  
Fax :  
CIN - U74899PB1995PLC045956



Patient Ref. No. 77500005852721

<b>PATIENT NAME : HEMLATA JOSHI</b>		<b>REF. DOCTOR : SELF</b>	
<b>CODE/NAME &amp; ADDRESS : C000138394</b>		<b>ACCESSION NO : 0181WL001806</b>	<b>AGE/SEX : 36 Years Female</b>
ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST DELHI NEW DELHI 110030 8800465156		<b>PATIENT ID : HEMLF200387181</b>	<b>DRAWN :</b>
		<b>CLIENT PATIENT ID:</b>	<b>RECEIVED : 23/12/2023 10:42:50</b>
		<b>ABHA NO :</b>	<b>REPORTED : 28/12/2023 16:07:44</b>

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**

<b>FBS (FASTING BLOOD SUGAR)</b>	<b>87</b>	Normal 75 - 99 Pre-diabetics: 100 - 125 Diabetic: > or = 126	mg/dL
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METHOD : ENZYMATIC REFERENCE METHOD WITH HEXOKINASE

**GLUCOSE, POST-PRANDIAL, PLASMA**

<b>PPBS(POST PRANDIAL BLOOD SUGAR)</b>	<b>81</b>	70 - 139	mg/dL
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METHOD : ENZYMATIC REFERENCE METHOD WITH HEXOKINASE

**LIPID PROFILE WITH CALCULATED LDL**

<b>CHOLESTEROL, TOTAL</b>	<b>219 High</b>	Desirable : < 200 Borderline : 200 - 239 High : > / = 240	mg/dL
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METHOD : ENZYMATIC COLORIMETRIC ASSAY

<b>TRIGLYCERIDES</b>	<b>105</b>	Normal: < 150 Borderline high: 150 - 199 High: 200 - 499 Very High: >/= 500	mg/dL
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METHOD : ENZYMATIC COLORIMETRIC ASSAY

<b>HDL CHOLESTEROL</b>	<b>62 High</b>	At Risk: < 40 Desirable: > or = 60	mg/dL
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METHOD : ENZYMATIC, COLORIMETRIC

<b>CHOLESTEROL LDL</b>	<b>136 High</b>	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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METHOD : ENZYMATIC COLORIMETRIC ASSAY

**Dr. Priyal Chinchkhede, MD**  
Consultant Pathologist

**Dr. Ushma Wartikar, MD**  
Consultant Pathologist

**Dr. (Mrs) Neelu K Bhojani**  
Lab Head



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NON HDL CHOLESTEROL		<b>157 High</b>	Desirable : < 130 Above Desirable : 130 -159 Borderline High : 160 - 189 High : 190 - 219 Very high : > / = 220	mg/dL
VERY LOW DENSITY LIPOPROTEIN CHOL/HDL RATIO		21 3.5	< OR = 30.0 Low Risk : 3.3 - 4.4 Average Risk : 4.5 - 7.0 Moderate Risk : 7.1 - 11.0 High Risk : > 11.0	mg/dL
LDL/HDL RATIO		2.2	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**

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)

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Lab Head



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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**

<b>BILIRUBIN, TOTAL</b> METHOD : COLORIMETRIC DIAZO	0.33	Upto 1.2	mg/dL
<b>BILIRUBIN, DIRECT</b> METHOD : DIAZO METHOD	0.27	< 0.30	mg/dL
<b>BILIRUBIN, INDIRECT</b>	<b>0.06 Low</b>	0.1 - 1.0	mg/dL
<b>TOTAL PROTEIN</b> METHOD : COLORIMETRIC	<b>8.6 High</b>	6.0 - 8.0	g/dL
<b>ALBUMIN</b> METHOD : COLORIMETRIC	4.8	3.97 - 4.94	g/dL
<b>GLOBULIN</b>	<b>3.8 High</b>	2.0 - 3.5	g/dL
<b>ALBUMIN/GLOBULIN RATIO</b>	1.3	1.0 - 2.1	RATIO
<b>ASPARTATE AMINOTRANSFERASE(AST/SGOT)</b> METHOD : UV ABSORBANCE	26	< OR = 35	U/L
<b>ALANINE AMINOTRANSFERASE (ALT/SGPT)</b> METHOD : UV ABSORBANCE	24	< OR = 35	U/L
<b>ALKALINE PHOSPHATASE</b> METHOD : COLORIMETRIC	71	35 - 104	U/L
<b>GAMMA GLUTAMYL TRANSFERASE (GGT)</b> METHOD : ENZYMATIC, COLORIMETRIC	26	0 - 40	U/L
<b>LACTATE DEHYDROGENASE</b> METHOD : UV ABSORBANCE	205	125 - 220	U/L

**BLOOD UREA NITROGEN (BUN), SERUM**

<b>BLOOD UREA NITROGEN</b> METHOD : ENZYMATIC ASSAY	7	6 - 20	mg/dL
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**Consultant Pathologist**

**Dr. Ushma Wartikar, MD**  
**Consultant Pathologist**

**Dr. (Mrs) Neelu K Bhojani**  
**Lab Head**



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REF. DOCTOR : SELF

CODE/NAME &amp; ADDRESS : C000138394

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

CREATININE	0.51	0.5 - 0.9	mg/dL
METHOD : COLORIMETRIC			

**BUN/CREAT RATIO**

BUN/CREAT RATIO	13.73	8.0 - 15.0	
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**URIC ACID, SERUM**

URIC ACID	3.0	2.4 - 5.7	mg/dL
METHOD : ENZYMATIC COLORIMETRIC ASSAY			

**TOTAL PROTEIN, SERUM**

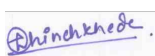
TOTAL PROTEIN	<b>8.6 High</b>	6.0 - 8.0	g/dL
METHOD : COLORIMETRIC			

**ALBUMIN, SERUM**

ALBUMIN	4.8	3.97 - 4.94	g/dL
METHOD : COLORIMETRIC			

**GLOBULIN**

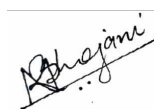
GLOBULIN	<b>3.8 High</b>	2.0 - 3.5	g/dL
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**ELECTROLYTES (NA/K/CL), SERUM**


 Dr. Priyal Chinchkhede, MD  
 Consultant Pathologist



 Dr. Ushma Wartikar, MD  
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SODIUM, SERUM		144	136 - 145	mmol/L
METHOD : ION SELECTIVE ELECTRODE TECHNOLOGY				
POTASSIUM, SERUM		4.87	3.5 - 5.1	mmol/L
METHOD : ION SELECTIVE ELECTRODE TECHNOLOGY				
CHLORIDE, SERUM		107	98 - 107	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,hyppopituitarism,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.

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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 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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**CLINICAL PATH - URINALYSIS**

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

**PHYSICAL EXAMINATION, URINE**

<b>COLOR</b> METHOD : MICROSCOPIC EXAMINATION	PALE YELLOW
<b>APPEARANCE</b> METHOD : MICROSCOPIC EXAMINATION	<b>SLIGHTLY HAZY</b>

**CHEMICAL EXAMINATION, URINE**

<b>PH</b> METHOD : METHYL RED & BROMOTHYMOL BLUE	6.0	5.00 - 7.50
<b>SPECIFIC GRAVITY</b>	1.015	1.010 - 1.030
<b>PROTEIN</b> METHOD : TETRA BROMOPHENOL BLUE/SULFOSALICYLIC ACID	NOT DETECTED	NOT DETECTED
<b>GLUCOSE</b> METHOD : GLUCOSE OXIDASE / PEROXIDASE (GOD - POD) METHOD	NOT DETECTED	NOT DETECTED
<b>KETONES</b> METHOD : SODIUM NITROPRUSSIDE REACTION	NOT DETECTED	NOT DETECTED
<b>BLOOD</b> METHOD : STRIP TEST - DIAZONIUM SALT COUPLING	NOT DETECTED	NOT DETECTED
<b>UROBILINOGEN</b> METHOD : CAFFEINE BENZOATE	NORMAL	NORMAL
<b>NITRITE</b> METHOD : STRIP NAPHTHOETHYLENEDIAMINE HYDROCHLORIDE,TATTANIC ACID	NOT DETECTED	NOT DETECTED
<b>LEUKOCYTE ESTERASE</b> METHOD : STRIP HETROCYCLIC CARBOXYLIC ACID ESTER ,DIAZONIUM SALT	<b>DETECTED (+)</b>	NOT DETECTED

**MICROSCOPIC EXAMINATION, URINE**

<b>RED BLOOD CELLS</b> METHOD : MICROSCOPIC EXAMINATION	NOT DETECTED	NOT DETECTED	/HPF
<b>PUS CELL (WBC'S)</b> METHOD : MICROSCOPIC EXAMINATION	<b>8-10</b>	0-5	/HPF
<b>EPITHELIAL CELLS</b> METHOD : MICROSCOPIC EXAMINATION	<b>8-10</b>	0-5	/HPF

**Dr. (Mrs) Neelu K Bhojani**  
Lab Head

**Dr. Ushma Wartikar, MD**  
Consultant Pathologist

**Dr. Priyal Chinchkhede, MD**  
Consultant Pathologist



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**PATIENT NAME : HEMLATA JOSHI**

**REF. DOCTOR : SELF**

**CODE/NAME & ADDRESS :** C000138394  
 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL  
 F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST  
 DELHI  
 NEW DELHI 110030  
 8800465156

**ACCESSION NO :** 0181WL001806  
**PATIENT ID :** HEMLF200387181  
**CLIENT PATIENT ID:**  
**ABHA NO :**

**AGE/SEX :** 36 Years Female  
**DRAWN :**  
**RECEIVED :** 23/12/2023 10:42:50  
**REPORTED :** 28/12/2023 16:07:44

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CASTS METHOD : MICROSCOPIC EXAMINATION	NOT DETECTED			
CRYSTALS METHOD : MICROSCOPIC EXAMINATION	NOT DETECTED			
BACTERIA METHOD : MICROSCOPIC EXAMINATION	NOT DETECTED	NOT DETECTED		
YEAST	NOT DETECTED	NOT DETECTED		

**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

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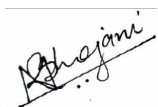
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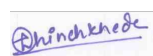
Uric acid	arthritis
Bacteria	Urinary infection when present in significant numbers & with pus cells.
Trichomonas vaginalis	Vaginitis, cervicitis or salpingitis



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Consultant Pathologist

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Patient Ref. No. 77500005852721

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<b>CODE/NAME &amp; ADDRESS</b> : C000138394 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST DELHI NEW DELHI 110030 8800465156	<b>ACCESSION NO</b> : <b>0181WL001806</b>	<b>AGE/SEX</b> : 36 Years	Female
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**CYTOLOGY**

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

**PAPANICOLAOU SMEAR**

<b>TEST METHOD</b>	CONVENTIONAL GYNEC CYTOLOGY
METHOD : MICROSCOPIC EXAMINATION	
<b>SPECIMEN TYPE</b>	P-1894/23
METHOD : MICROSCOPIC EXAMINATION	
<b>REPORTING SYSTEM</b>	TWO UNSTAINED CERVICAL SMEARS RECEIVED
<b>SPECIMEN ADEQUACY</b>	2014 BETHESDA SYSTEM FOR REPORTING CERVICAL CYTOLOGY
METHOD : PAP STAIN & MICROSCOPIC EXAMINATION	
<b>MICROSCOPY</b>	SATISFACTORY
METHOD : PAP STAIN	
<b>INTERPRETATION / RESULT</b>	THE SMEARS SHOW MAINLY SUPERFICIAL SQUAMOUS CELLS, FEW INTERMEDIATE SQUAMOUS CELLS, MANY CLUSTERS OF ENDOCERVICAL CELLS IN THE BACKGROUND OF FEW POLYMORPHS & RBC'S.
METHOD : PAP STAIN & MICROSCOPIC EXAMINATION	
	NEGATIVE FOR INTRAEPITHELIAL LESION OR MALIGNANCY

**Comments**

PLEASE NOTE PAPANICOLAOU SMEAR STUDY IS A SCREENING PROCEDURE FOR CERVICAL CANCER WITH INHERENT FALSE NEGATIVE RESULTS HENCE SHOULD BE INTERPRETED WITH CAUTION. NO CYTOLOGICAL EVIDENCE OF HPV INFECTION IN THE SMEARS STUDIED. SMEARS WILL BE PRESERVED FOR 5 YEARS ONLY.

**Dr. Priyal Chinchkhede, MD**  
Consultant Pathologist



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**CLINICAL PATH - STOOL ANALYSIS**
**MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE**
**PHYSICAL EXAMINATION,STOOL**

COLOUR

SAMPLE NOT RECEIVED

METHOD : VISUAL


**Dr. Sheetal Sawant, MD**  
**Consultant Microbiologist**

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### SPECIALISED CHEMISTRY - HORMONE

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

##### THYROID PANEL, SERUM

T3	140.0	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 : ELECTROCHEMILUMINESCENCE			
T4	6.83	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 : ELECTROCHEMILUMINESCENCE			
TSH (ULTRASENSITIVE)	1.330	Non Pregnant Women 0.27 - 4.20 Pregnant Women (As per American Thyroid Association) 1st Trimester 0.100 - 2.500 2nd Trimester 0.200 - 3.000 3rd Trimester 0.300 - 3.000	µIU/mL
METHOD : ELECTROCHEMILUMINESCENCE			

#### Interpretation(s)

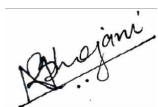
**Triiodothyronine T3**, **Thyroxine T4**, and **Thyroid Stimulating Hormone TSH** are thyroid hormones which affect almost every physiological process in the body, including growth, development, metabolism, body temperature, and heart rate.

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.

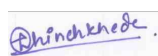
Excessive secretion of thyroxine in the body is hyperthyroidism, and deficient secretion is called hypothyroidism.

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



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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
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.

**Dr. (Mrs) Neelu K Bhojani**  
Lab Head

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