

Patient Name : Mr.TANIKONDA MALLIKHARJUN	Collected : 09/Sep/2023 08:43AM
Age/Gender : 36 Y 0 M 30 D/M	Received : 09/Sep/2023 10:12AM
UHID/MR No : CMYS.0000057872	Reported : 09/Sep/2023 11:35AM
Visit ID : CMYSOPV117340	Status : Final Report
Ref Doctor : Dr.SELF	Sponsor Name : ARCOFEMI HEALTHCARE LIMITED
Emp/Auth/TPA ID : 167186	

DEPARTMENT OF HAEMATOLOGY

ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

PERIPHERAL SMEAR , WHOLE BLOOD EDTA

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SIN No:BED230217048

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**HEMOGRAM , WHOLE BLOOD EDTA**

<b>HAEMOGLOBIN</b>	16.1	g/dL	13-17	Spectrophotometer
PCV	48.70	%	40-50	Electronic pulse & Calculation
RBC COUNT	<b>5.6</b>	Million/cu.mm	4.5-5.5	Electrical Impedence
MCV	87	fL	83-101	Calculated
MCH	28.7	pg	27-32	Calculated
MCHC	33	g/dL	31.5-34.5	Calculated
R.D.W	13.4	%	11.6-14	Calculated
TOTAL LEUCOCYTE COUNT (TLC)	<b>11,900</b>	cells/cu.mm	4000-10000	Electrical Impedence

**DIFFERENTIAL LEUCOCYTIC COUNT (DLC)**

NEUTROPHILS	66.4	%	40-80	Electrical Impedence
LYMPHOCYTES	25.2	%	20-40	Electrical Impedence
EOSINOPHILS	<b>0.9</b>	%	1-6	Electrical Impedence
MONOCYTES	6.9	%	2-10	Electrical Impedence
BASOPHILS	0.6	%	<1-2	Electrical Impedence

**ABSOLUTE LEUCOCYTE COUNT**

NEUTROPHILS	<b>7901.6</b>	Cells/cu.mm	2000-7000	Electrical Impedence
LYMPHOCYTES	2998.8	Cells/cu.mm	1000-3000	Electrical Impedence
EOSINOPHILS	107.1	Cells/cu.mm	20-500	Electrical Impedence
MONOCYTES	821.1	Cells/cu.mm	200-1000	Electrical Impedence
BASOPHILS	71.4	Cells/cu.mm	0-100	Electrical Impedence

**PLATELET COUNT**

PLATELET COUNT	391000	cells/cu.mm	150000-410000	Electrical impedence
<b>ERYTHROCYTE SEDIMENTATION RATE (ESR)</b>	06	mm at the end of 1 hour	0-15	Modified Westergren

**PERIPHERAL SMEAR**

R.B.C: Majority are normocytic normochromic.  
W.B.C: Are increased in number with predominance of neutrophils.  
Platelets: Adequate and are seen in singles and clumps.  
Hemoparasites: Not seen.

**IMPRESSION: NORMOCYTIC NORMOCHROMIC BLOOD PICTURE WITH NEUTROPHILIC**

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**LEUCOCYTOSIS.**



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<b>BLOOD GROUP ABO AND RH FACTOR , WHOLE BLOOD EDTA</b>				
BLOOD GROUP TYPE	O			Forward & Reverse Grouping with Slide/Tube Aggluti
Rh TYPE	POSITIVE			Forward & Reverse Grouping with Slide/Tube Agglutination



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**DEPARTMENT OF BIOCHEMISTRY**

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<b>GLUCOSE, FASTING , NAF PLASMA</b>	<b>103</b>	mg/dL	70-100	GOD - POD
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**Comment:**

As per American Diabetes Guidelines, 2023

Fasting Glucose Values in mg/dL	Interpretation
70-100 mg/dL	Normal
100-125 mg/dL	Prediabetes
≥126 mg/dL	Diabetes
<70 mg/dL	Hypoglycemia

**Note:**

- The diagnosis of Diabetes requires a fasting plasma glucose of > or = 126 mg/dL and/or a random / 2 hr post glucose value of > or = 200 mg/dL on at least 2 occasions.
- Very high glucose levels (>450 mg/dL in adults) may result in Diabetic Ketoacidosis & is considered critical.

<b>GLUCOSE, POST PRANDIAL (PP), 2 HOURS , SODIUM FLUORIDE PLASMA (2 HR)</b>	<b>149</b>	mg/dL	70-140	GOD - POD
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**Comment:**

It is recommended that FBS and PPBS should be interpreted with respect to their Biological reference ranges and not with each other.

Conditions which may lead to lower postprandial glucose levels as compared to fasting glucose levels may be due to reactive hypoglycemia, dietary meal content, duration or timing of sampling after food digestion and absorption, medications such as insulin preparations, sulfonylureas, amylin analogues, or conditions such as overproduction of insulin.

Ref: Marks medical biochemistry and clinical approach

<b>HBA1C, GLYCATED HEMOGLOBIN ,</b>	<b>4.6</b>	%		HPLC
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Test Name	Result	Unit	Bio. Ref. Range	Method
WHOLE BLOOD EDTA				
<b>ESTIMATED AVERAGE GLUCOSE (eAG) ,</b> WHOLE BLOOD EDTA	85	mg/dL		Calculated

**Comment:**

Reference Range as per American Diabetes Association (ADA) 2023 Guidelines:

REFERENCE GROUP	HBA1C %
NON DIABETIC	<5.7
PREDIABETES	5.7 – 6.4
DIABETES	≥ 6.5
DIABETICS	
EXCELLENT CONTROL	6 – 7
FAIR TO GOOD CONTROL	7 – 8
UNSATISFACTORY CONTROL	8 – 10
POOR CONTROL	>10

**Note:** Dietary preparation or fasting is not required.

- HbA1C is recommended by American Diabetes Association for Diagnosing Diabetes and monitoring Glycemic Control by American Diabetes Association guidelines 2023.
- Trends in HbA1C values is a better indicator of Glycemic control than a single test.
- Low HbA1C in Non-Diabetic patients are associated with Anemia (Iron Deficiency/Hemolytic), Liver Disorders, Chronic Kidney Disease. Clinical Correlation is advised in interpretation of low Values.
- Falsely low HbA1c (below 4%) may be observed in patients with clinical conditions that shorten erythrocyte life span or decrease mean erythrocyte age. HbA1c may not accurately reflect glycemic control when clinical conditions that affect erythrocyte survival are present.
- In cases of Interference of Hemoglobin variants in HbA1C, alternative methods (Fructosamine) estimation is recommended for Glycemic Control
  - A: HbF >25%
  - B: Homozygous Hemoglobinopathy.
 (Hb Electrophoresis is recommended method for detection of Hemoglobinopathy)

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SIN No:PLF02024850,PLP1367197,EDT230082846

**APOLLO CLINICS NETWORK**

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**DEPARTMENT OF BIOCHEMISTRY**

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**LIPID PROFILE , SERUM**

TOTAL CHOLESTEROL	<b>213</b>	mg/dL	<200	CHE/CHO/POD
TRIGLYCERIDES	<b>155</b>	mg/dL	<150	Enzymatic
HDL CHOLESTEROL	<b>36</b>	mg/dL	>40	CHE/CHO/POD
NON-HDL CHOLESTEROL	<b>177</b>	mg/dL	<130	Calculated
LDL CHOLESTEROL	<b>146</b>	mg/dL	<100	Calculated
VLDL CHOLESTEROL	<b>31</b>	mg/dL	<30	Calculated
CHOL / HDL RATIO	<b>5.92</b>		0-4.97	Calculated

**Comment:**

Reference Interval as per National Cholesterol Education Program (NCEP) Adult Treatment Panel III Report.

	Desirable	Borderline High	High	Very High
TOTAL CHOLESTEROL	< 200	200 - 239	≥ 240	
TRIGLYCERIDES	<150	150 - 199	200 - 499	≥ 500
LDL	Optimal < 100 Near Optimal 100-129	130 - 159	160 - 189	≥ 190
HDL	≥ 60			
NON-HDL CHOLESTEROL	Optimal <130; Above Optimal 130-159	160-189	190-219	>220

1. Measurements in the same patient on different days can show physiological and analytical variations.
2. NCEP ATP III identifies non-HDL cholesterol as a secondary target of therapy in persons with high triglycerides.
3. Primary prevention algorithm now includes absolute risk estimation and lower LDL Cholesterol target levels to determine eligibility of drug therapy.
4. Low HDL levels are associated with Coronary Heart Disease due to insufficient HDL being available to participate in reverse cholesterol transport, the process by which cholesterol is eliminated from peripheral tissues.
5. As per NCEP guidelines, all adults above the age of 20 years should be screened for lipid status. Selective screening of children above the age of 2 years with a family history of premature cardiovascular disease or those with at least one parent with high total cholesterol is recommended.
6. VLDL, LDL Cholesterol Non HDL Cholesterol, CHOL/HDL RATIO, LDL/HDL RATIO are calculated parameters when Triglycerides are below 350 mg/dl. When Triglycerides are more than 350 mg/dl LDL cholesterol is a direct measurement.

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Test Name	Result	Unit	Bio. Ref. Range	Method
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SIN No:SE04476130

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LIVER FUNCTION TEST (LFT) , SERUM				
BILIRUBIN, TOTAL	<b>1.80</b>	mg/dL	0.20-1.20	DIAZO METHOD
BILIRUBIN CONJUGATED (DIRECT)	<b>0.50</b>	mg/dL	0.0-0.3	Calculated
BILIRUBIN (INDIRECT)	<b>1.30</b>	mg/dL	0.0-1.1	Dual Wavelength
ALANINE AMINOTRANSFERASE (ALT/SGPT)	<b>62</b>	U/L	<50	Visible with P-5-P
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	41.0	U/L	17-59	UV with P-5-P
ALKALINE PHOSPHATASE	122.00	U/L	38-126	p-nitrophenyl phosphate
PROTEIN, TOTAL	<b>8.60</b>	g/dL	6.3-8.2	Biuret
ALBUMIN	5.00	g/dL	3.5 - 5	Bromocresol Green
GLOBULIN	<b>3.60</b>	g/dL	2.0-3.5	Calculated
A/G RATIO	1.39		0.9-2.0	Calculated



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<b>RENAL PROFILE/KIDNEY FUNCTION TEST (RFT/KFT) , SERUM</b>				
CREATININE	0.90	mg/dL	0.66-1.25	Creatinine amidohydrolase
UREA	24.80	mg/dL	19-43	Urease
BLOOD UREA NITROGEN	11.6	mg/dL	8.0 - 23.0	Calculated
URIC ACID	7.80	mg/dL	3.5-8.5	Uricase
CALCIUM	9.90	mg/dL	8.4 - 10.2	Arsenazo-III
PHOSPHORUS, INORGANIC	3.00	mg/dL	2.5-4.5	PMA Phenol
SODIUM	141	mmol/L	135-145	Direct ISE
POTASSIUM	5.0	mmol/L	3.5-5.1	Direct ISE
CHLORIDE	106	mmol/L	98 - 107	Direct ISE



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Test Name	Result	Unit	Bio. Ref. Range	Method
<b>GAMMA GLUTAMYL TRANSPEPTIDASE (GGT) , SERUM</b>	68.00	U/L	15-73	Glycylglycine Nitoranalide



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**DEPARTMENT OF IMMUNOLOGY**

**ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324**

Test Name	Result	Unit	Bio. Ref. Range	Method
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**THYROID PROFILE TOTAL (T3, T4, TSH) , SERUM**

TRI-IODOTHYRONINE (T3, TOTAL)	1.50	ng/mL	0.64-1.52	CMIA
THYROXINE (T4, TOTAL)	9.93	µg/dL	4.87-11.72	CMIA
THYROID STIMULATING HORMONE (TSH)	1.750	µIU/mL	0.35-4.94	CMIA

**Comment:**

**Note:**

For pregnant females	Bio Ref Range for TSH in uIU/ml (As per American Thyroid Association)
First trimester	0.1 - 2.5
Second trimester	0.2 – 3.0
Third trimester	0.3 – 3.0

1. TSH is a glycoprotein hormone secreted by the anterior pituitary. TSH activates production of T3 (Triiodothyronine) and its prohormone T4 (Thyroxine). Increased blood level of T3 and T4 inhibit production of TSH.
2. TSH is elevated in primary hypothyroidism and will be low in primary hyperthyroidism. Elevated or low TSH in the context of normal free thyroxine is often referred to as sub-clinical hypo- or hyperthyroidism respectively.
3. Both T4 & T3 provides limited clinical information as both are highly bound to proteins in circulation and reflects mostly inactive hormone. Only a very small fraction of circulating hormone is free and biologically active.
4. Significant variations in TSH can occur with circadian rhythm, hormonal status, stress, sleep deprivation, medication & circulating antibodies.

TSH	T3	T4	FT4	Conditions
High	Low	Low	Low	Primary Hypothyroidism, Post Thyroidectomy, Chronic Autoimmune Thyroiditis
High	N	N	N	Subclinical Hypothyroidism, Autoimmune Thyroiditis, Insufficient Hormone Replacement Therapy.
N/Low	Low	Low	Low	Secondary and Tertiary Hypothyroidism
Low	High	High	High	Primary Hyperthyroidism, Goitre, Thyroiditis, Drug effects, Early Pregnancy
Low	N	N	N	Subclinical Hyperthyroidism
Low	Low	Low	Low	Central Hypothyroidism, Treatment with Hyperthyroidism

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Low	N	High	High	Thyroiditis, Interfering Antibodies
N/Low	High	N	N	T3 Thyrotoxicosis, Non thyroidal causes
High	High	High	High	Pituitary Adenoma; TSHoma/Thyrotropinoma



SIN No:SPL23128353

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**DEPARTMENT OF CLINICAL PATHOLOGY**

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**COMPLETE URINE EXAMINATION (CUE) , URINE**

**PHYSICAL EXAMINATION**

COLOUR	PALE YELLOW		PALE YELLOW	Visual
TRANSPARENCY	CLEAR		CLEAR	Visual
pH	6.5		5-7.5	Bromothymol Blue
SP. GRAVITY	1.020		1.002-1.030	Dipstick

**BIOCHEMICAL EXAMINATION**

URINE PROTEIN	NEGATIVE		NEGATIVE	PROTEIN ERROR OF INDICATOR
GLUCOSE	NEGATIVE		NEGATIVE	GOD-POD
URINE BILIRUBIN	NEGATIVE		NEGATIVE	AZO COUPLING
URINE KETONES (RANDOM)	NEGATIVE		NEGATIVE	NITROPRUSSIDE
UROBILINOGEN	NORMAL		NORMAL	EHRlich
BLOOD	NEGATIVE		NEGATIVE	Dipstick
NITRITE	NEGATIVE		NEGATIVE	Dipstick
LEUCOCYTE ESTERASE	NEGATIVE		NEGATIVE	PYRROLE HYDROLYSIS

**CENTRIFUGED SEDIMENT WET MOUNT AND MICROSCOPY**

PUS CELLS	2 - 3	/hpf	0-5	Microscopy
EPITHELIAL CELLS	1 - 2	/hpf	<10	MICROSCOPY
RBC	NIL	/hpf	0-2	MICROSCOPY
CASTS	NIL		0-2 Hyaline Cast	MICROSCOPY
CRYSTALS	ABSENT		ABSENT	MICROSCOPY



SIN No:UR2180756


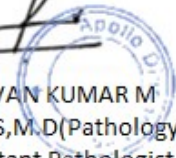
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Test Name	Result	Unit	Bio. Ref. Range	Method
URINE GLUCOSE(POST PRANDIAL)	NEGATIVE		NEGATIVE	Dipstick
URINE GLUCOSE(FASTING)	NEGATIVE		NEGATIVE	Dipstick

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

  
  
 Dr. PAVAN KUMAR M  
 M.B.B.S,M.D(Pathology)  
 Consultant Pathologist



SIN No:UPP015457,UF009418