

**DEPARTMENT OF LABORATORY MEDICINE**

Patient Name	: Mrs. MONIKA SAKLANI	Order No	: 1000104120
UHID	: UHJ A24008240	Registered On	: 18/11/2024 08:25:12 AM
Age/Sex	: 32/Years Female	Collected On	: 18/11/2024 08:37:55 AM
Ward / Bed No	:	Reported On	: 18/11/2024 01:27:46 PM
Reference	: Dr. Ashmitha Padma	Bill No	: OPBJ A240011116
Station	: At Hospital	Mobile No	: 9870932003
Payer Name	: Mediwheel	Report Status	: Final Report

Test Name	Result	Unit	Bio. Ref. Interval
<b><u>BIOCHEMISTRY</u></b>			
<b>FASTING GLUCOSE</b> (Method: Hexokinase)	138	mg/dL	ADA Guidelines < 100 mg/dl - Normal 100 to 125 mg/dl - Prediabetes ≥ 126 mg/dl - Diabetes
<b>POST PRANDIAL GLUCOSE</b> (Method: Hexokinase)	269	mg/dL	70-140
<b>GLYCOSYLATED HAEMOGLOBIN (HBA1C)</b>			Sample: Whole blood (EDTA)
<b>HBA1C</b> (Method: HPLC)	6.7	%	ADA Guidelines < 5.7% - Normal 5.7 to 6.4% - Prediabetes ≥ 6.5% - Diabetes
<b>Estimated Average Glucose (eAG)</b> (Method: Calculated)	146	mg/dL	
<b>THYROID PROFILE (TOTAL T3, TOTAL T4 &amp; TSH)</b>			Sample: Serum
<b>TOTAL T3</b> (Method: CLIA)	0.94	ng/mL	0.87-1.78
<b>TOTAL T4</b> (Method: CLIA)	11.59	ng/dL	5.1-14.1
<b>THYROID STIMULATING HORMONE (TSH)</b> (Method: CLIA: Ultra-sensitive)	2.36	μIU/mL	0.34 - 5.60 μIU/mL (Non Pregnant) 0.3 - 4.5 μIU/mL (I trimester) 0.5 - 5.2 μIU/mL (II & III trimester)
<b>LIPID PROFILE</b>			Sample: Serum
<b>TOTAL CHOLESTEROL</b> (Method: CHOD-POD)	147	mg/dL	ATP III Guidelines < 200 - Desirable 200-239 - Borderline high ≥ 240 - High
<b>TRIGLYCERIDES</b> (Method: Enzymatic GPO-POD)	70	mg/dL	< 150 - Normal 150-199 - Borderline High 200-499 - High ≥ 500 - Very High
<b>HDL CHOLESTEROL</b> (Method: ENZYMATIC METHOD)	39.6	mg/dL	< 40 - Low ≥ 60 - High

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Test Name	Result	Unit	Bio. Ref. Interval
<b>LDL CHOLESTEROL</b> (Method: Calculated)	93.40	mg/dL	<100 - Optimal 100-129 - Near or above optimal 130-159 - Borderline high 160-189 - High ≥190 - Very high
<b>VLDL CHOLESTEROL</b> (Method: Calculated)	14.00	mg/dL	< 30
<b>TOTAL CHOLESTEROL : HDL RATIO</b> (Method: Calculated)	3.71		Low Risk: 3.3 - 4.4 Average Risk: 4.5 - 7.1 Moderate Risk: 7.2 - 11.0
<b>LDL/HDL CHOLESTEROL RATIO</b> (Method: Calculated)	2.36		< 2.5 Optimal
<b>NON HDL CHOLESTEROL</b> (Method: Calculated)	107.40	mg/dL	< 130
<b>URIC ACID</b> (Method:Uricase - POD(Enzymatic))	5.4	mg/dL	2.6-6.0
<b>BLOOD UREA NITROGEN(BUN)</b> (Method:Urease GLDH - Kinetic)	9	mg/dL	7.93-20.07
<b>CREATININE</b> (Method:Modified Jaffe, Kinetic)	0.62	mg/dL	0.6-1.1
<b>LIVER FUNCTION TEST</b>			
<b>TOTAL BILIRUBIN</b> (Method:Dichlorophenyl Diazotization)	0.78	mg/dL	0.3-1.2
<b>DIRECT BILIRUBIN</b> (Method:Dichlorophenyl Diazotization)	0.21	mg/dL	0.0-0.2
<b>INDIRECT BILIRUBIN</b> (Method: Calculated)	0.57	mg/dL	0.2-1.0
<b>TOTAL PROTEIN</b> (Method:BIURET)	7.5	g/dL	6.6-8.3
<b>ALBUMIN</b> (Method:BCG)	3.98	g/dL	3.5-5.2
<b>GLOBULIN</b> (Method: Calculated)	3.52	g/dL	2.3-3.5

Sample: Serum

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Test Name	Result	Unit	Bio. Ref. Interval
AG RATIO (Method: Calculated)	1.13		2:1
SERUM SGOT (Method:IFCC without P5P)	24	U/L	< 35
SERUM SGPT (Method:IFCC without P5P)	22	U/L	< 35
ALKALINE PHOSPHATASE, SERUM (Method:PNPP AMP Buffer)	63	U/L	46-122
GGT (Method:IFCC)	28	U/L	< 38



**Dr. Varsha Shree R**  
M.D(Pathology)  
CONSULTANT PATHOLOGIST  
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**HAEMATOLOGY**
**COMPLETE BLOOD COUNT(CBC)**

Sample: Whole blood (EDTA)

<b>HAEMOGLOBIN</b> (Method:Photometric Measurement: Oxyhemoglobin method)	12.70	g/dL	12-16
<b>PACKED CELL VOLUME/HEMATOCRIT (PCV/HCT)</b> (Method: Calculated)	40.2	%	37-47
<b>TOTAL WBC COUNT (TLC)</b> (Method:Coulter Principle)	6480	Cells/Cum	4000-11000
<b>DIFFERENTIAL COUNT</b>			
<b>NEUTROPHILS</b> (Method:Optical/Impedance)	65.04	%	40-75
<b>LYMPHOCYTES</b> (Method:Optical/Impedance)	27.86	%	20-45
<b>EOSINOPHILS</b> (Method:Optical/Impedance)	1.34	%	0-6
<b>MONOCYTES</b> (Method:Optical/Impedance)	5.54	%	2-10
<b>BASOPHILS</b> (Method:Optical/Impedance)	0.22	%	0-2
<b>RED BLOOD CORPUSCLES(RBC)</b> (Method:Coulter Principle)	4.54	million/cum	4.0-5.2
<b>MCV</b> (Method:Derived from RBC Histogram)	88.5	fL	78-100
<b>MCH</b> (Method: Calculated)	28.0	pg	27-31
<b>MCHC</b> (Method: Calculated)	31.6	g/dL	31-37
<b>RDW - CV</b> (Method: Calculated)	13.5	%	11.5-14.5
<b>PLATELET COUNT</b> (Method:Electrical Impedance)	2.21	Lakhs/Cum	1.5-4.5


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Test Name	Result	Unit	Bio. Ref. Interval
MEAN PLATELET VOLUME(MPV) (Method:Derived from PLT Histogram)	10.44	fl	9-13
PLATELET DISTRIBUTION WIDTH (PDW) (Method: Calculated)	19.5	fl	9-19
ABSOLUTE NEUTROPHIL COUNT (ANC) (Method: Calculated)	4210	Cells/Cum	1500-7500
ABSOLUTE EOSINOPHIL COUNT (AEC) (Method: Calculated)	90	Cells/Cum	40-440
ABSOLUTE LYMPHOCYTE COUNT (ALC) (Method: Calculated)	1810	Cells/Cum	1000-4000
ABSOLUTE MONOCYTE COUNT (AMC) (Method: Calculated)	360	Cells/Cum	200-1000
ABSOLUTE BASOPHIL COUNT (ABC) (Method: Calculated)	10	Cells/Cum	20-100
<b>ERYTHROCYTE SEDIMENTATION RATE(ESR)</b> (Method:Modified Westergren Method)	18	mm/hour	1-20
<b>BLOOD GROUPING &amp; RH TYPING</b>			Sample: Whole blood (EDTA)
ABO Group (Method:Agglutination Method)	AB		
Rh Factor (Method:Agglutination Method)	Positive		

Interpretation Notes

Note: Both forward and reverse grouping performed



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<b><u>CLINICAL PATHOLOGY</u></b>			
<b>URINE EXAMINATION, ROUTINE</b>			
Sample: Urine			
<b>PHYSICAL EXAMINATION</b>			
VOLUME	25 mL	mL	
COLOUR	Pale Yellow		
APPEARANCE	Clear		
PH	5.0		5.0-8.0
SPECIFIC GRAVITY	1.020		1.005-1.030
<b>CHEMICAL EXAMINATION</b>			
PROTEIN	Absent		Absent
(Method:Protein Error of pH Indicator)			
GLUCOSE	Absent		Absent
(Method:GOD-POD)			
KETONE BODIES	Absent		Absent
(Method:Nitroprusside method/ Rothera's test)			
BILIRUBIN	Negative		Negative
(Method:DIAZO/FOUCHET'S TEST)			
BILE SALT	Absent		Absent
(Method:Hay's sulfur test)			
NITRITE	Negative		Negative
(Method:Griess method)			
UROBILINOGEN	Normal		
(Method:Azo coupling method)			
LEUKOCYTE ESTERASE	Negative		Negative
(Method:Leukocyte Esterase activity)			
BLOOD	Negative		Negative
(Method:Peroxidase Reaction)			
<b>MICROSCOPIC EXAMINATION</b>			

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EPITHELIAL CELLS	0-2	/HPF	0-5
PUS CELLS	0-2	/HPF	0-5
RBCs	Nil	/HPF	0-2
CASTS	Nil	/LPF	
CRYSTALS	Nil		
OTHERS	Nil		
<b>URINE SUGAR, FASTING</b> (Method:GOD-POD)	Absent		
<b>URINE SUGAR (POST PRANDIAL)</b>	Absent		

Verified By  
T S Abhishek

---End of Report---



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<b>HBA1C</b> (Method: HPLC)	6.7	%	ADA Guidelines < 5.7% - Normal 5.7 to 6.4% - Prediabetes ≥ 6.5% - Diabetes
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<b>THYROID PROFILE (TOTAL T3, TOTAL T4 &amp; TSH)</b>			Sample: Serum
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<b>TOTAL CHOLESTEROL : HDL RATIO</b> (Method: Calculated)	3.71		Low Risk: 3.3 - 4.4 Average Risk: 4.5 - 7.1 Moderate Risk: 7.2 - 11.0
<b>LDL/HDL CHOLESTEROL RATIO</b> (Method: Calculated)	2.36		< 2.5 Optimal
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<b>CREATININE</b> (Method:Modified Jaffe, Kinetic)	0.62	mg/dL	0.6-1.1
<b>LIVER FUNCTION TEST</b>			
<b>TOTAL BILIRUBIN</b> (Method:Dichlorophenyl Diazotization)	0.78	mg/dL	0.3-1.2
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<b>INDIRECT BILIRUBIN</b> (Method: Calculated)	0.57	mg/dL	0.2-1.0
<b>TOTAL PROTEIN</b> (Method:BIURET)	7.5	g/dL	6.6-8.3
<b>ALBUMIN</b> (Method:BCG)	3.98	g/dL	3.5-5.2
<b>GLOBULIN</b> (Method: Calculated)	3.52	g/dL	2.3-3.5

Sample: Serum

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SERUM SGPT (Method:IFCC without P5P)	22	U/L	< 35
ALKALINE PHOSPHATASE, SERUM (Method:PNPP AMP Buffer)	63	U/L	46-122
GGT (Method:IFCC)	28	U/L	< 38



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**HAEMATOLOGY**
**COMPLETE BLOOD COUNT(CBC)**

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<b>TOTAL WBC COUNT (TLC)</b> (Method:Coulter Principle)	6480	Cells/Cum	4000-11000
<b>DIFFERENTIAL COUNT</b>			
<b>NEUTROPHILS</b> (Method:Optical/Impedance)	65.04	%	40-75
<b>LYMPHOCYTES</b> (Method:Optical/Impedance)	27.86	%	20-45
<b>EOSINOPHILS</b> (Method:Optical/Impedance)	1.34	%	0-6
<b>MONOCYTES</b> (Method:Optical/Impedance)	5.54	%	2-10
<b>BASOPHILS</b> (Method:Optical/Impedance)	0.22	%	0-2
<b>RED BLOOD CORPUSCLES(RBC)</b> (Method:Coulter Principle)	4.54	million/cum	4.0-5.2
<b>MCV</b> (Method:Derived from RBC Histogram)	88.5	fL	78-100
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ABSOLUTE MONOCYTE COUNT (AMC) (Method: Calculated)	360	Cells/Cum	200-1000
ABSOLUTE BASOPHIL COUNT (ABC) (Method: Calculated)	10	Cells/Cum	20-100
<b>ERYTHROCYTE SEDIMENTATION RATE(ESR)</b> (Method:Modified Westergren Method)	18	mm/hour	1-20
<b>BLOOD GROUPING &amp; RH TYPING</b>			Sample: Whole blood (EDTA)
ABO Group (Method:Agglutination Method)	AB		
Rh Factor (Method:Agglutination Method)	Positive		

Interpretation Notes

Note: Both forward and reverse grouping performed



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CLINICAL PATHOLOGY
**URINE EXAMINATION, ROUTINE**

Sample: Urine

**PHYSICAL EXAMINATION**

VOLUME	25 mL	mL	
COLOUR	Pale Yellow		
APPEARANCE	Clear		
PH	5.0		5.0-8.0
SPECIFIC GRAVITY	1.020		1.005-1.030

**CHEMICAL EXAMINATION**

PROTEIN (Method:Protein Error of pH Indicator)	Absent		Absent
GLUCOSE (Method:GOD-POD)	Absent		Absent
KETONE BODIES (Method:Nitroprusside method/ Rothera's test)	Absent		Absent
BILIRUBIN (Method:DIAZO/FOUCHET'S TEST)	Negative		Negative
BILE SALT (Method:Hay's sulfur test)	Absent		Absent
NITRITE (Method:Griess method)	Negative		Negative
UROBILINOGEN (Method:Azo coupling method)	Normal		
LEUKOCYTE ESTERASE (Method:Leukocyte Esterase activity)	Negative		Negative
BLOOD (Method:Peroxidase Reaction)	Negative		Negative

**MICROSCOPIC EXAMINATION**


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Payer Name	: Mediwheel	Report Status	: Final Report

Test Name	Result	Unit	Bio. Ref. Interval
EPITHELIAL CELLS	0-2	/HPF	0-5
PUS CELLS	0-2	/HPF	0-5
RBCs	Nil	/HPF	0-2
CASTS	Nil	/LPF	
CRYSTALS	Nil		
OTHERS	Nil		
<b>URINE SUGAR, FASTING</b> (Method:GOD-POD)	Absent		
<b>URINE SUGAR (POST PRANDIAL)</b>	Absent		

Verified By  
T S Abhishek

---End of Report---



**Dr. Varsha Shree R**  
M.D(Pathology)  
CONSULTANT PATHOLOGIST  
KMC No : 103567