Name	: Ms. Bhuvaneshwari G		
PID No.	: MED130038156	Register On : 12/11/2022 8:39 AM	M
SID No.	: 422076937	Collection On : 12/11/2022 9:19 AM	
Age / Sex	: 49 Year(s) / Female	Report On : 12/11/2022 5:44 PM	MEDALL
Туре	: OP	Printed On : 05/12/2022 5:49 PM	
Ref. Dr	: MediWheel		

	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> Reference Interval
HAEMATOLOGY			
Complete Blood Count With - ESR			
Haemoglobin (EDTA Blood/Spectrophotometry)	14.4	g/dL	12.5 - 16.0
Packed Cell Volume(PCV)/Haematocrit (EDTA Blood)	44.8	%	37 - 47
RBC Count (EDTA Blood)	5.30	mill/cu.mm	4.2 - 5.4
Mean Corpuscular Volume(MCV) (EDTA Blood)	84.4	fL	78 - 100
Mean Corpuscular Haemoglobin(MCH) (EDTA Blood)	27.2	pg	27 - 32
Mean Corpuscular Haemoglobin concentration(MCHC) (EDTA Blood)	32.3	g/dL	32 - 36
RDW-CV (EDTA Blood)	14.4	%	11.5 - 16.0
RDW-SD (EDTA Blood)	42.54	fL	39 - 46
Total Leukocyte Count (TC) (EDTA Blood)	10600	cells/cu.mm	4000 - 11000
Neutrophils (EDTA Blood)	51.6	%	40 - 75
Lymphocytes (EDTA Blood)	31.4	%	20 - 45
Eosinophils (EDTA Blood)	10.8	%	01 - 06
Monocytes (EDTA Blood)	5.5	%	01 - 10



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Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> Reference Interval
Basophils (Blood)	0.7	%	00 - 02
INTERPRETATION: Tests done on Automated Five F	Part cell counter. All	abnormal results are	reviewed and confirmed microscopically.
Absolute Neutrophil count (EDTA Blood)	5.47	10^3 / µl	1.5 - 6.6
Absolute Lymphocyte Count (EDTA Blood)	3.33	10^3 / µl	1.5 - 3.5
Absolute Eosinophil Count (AEC) (EDTA Blood)	1.14	10^3 / µl	0.04 - 0.44
Absolute Monocyte Count (EDTA Blood)	0.58	10^3 / µl	< 1.0
Absolute Basophil count (EDTA Blood)	0.07	10^3 / µl	< 0.2
Platelet Count (EDTA Blood)	273	10^3 / µl	150 - 450
MPV (EDTA Blood)	8.1	fL	8.0 - 13.3
PCT (EDTA Blood/Automated Blood cell Counter)	0.22	%	0.18 - 0.28
ESR (Erythrocyte Sedimentation Rate) (Citrated Blood)	30	mm/hr	< 20

Dr.Arjun C.P MBBS,MD Pathology Reg No:KMC \$9655 APPROVED BY

The results pertain to sample tested.

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Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> Reference Interval
BIOCHEMISTRY			
Liver Function Test			
Bilirubin(Total) (Serum/DCA with ATCS)	0.56	mg/dL	0.1 - 1.2
Bilirubin(Direct) (Serum/Diazotized Sulfanilic Acid)	0.20	mg/dL	0.0 - 0.3
Bilirubin(Indirect) (Serum/Derived)	0.36	mg/dL	0.1 - 1.0
SGOT/AST (Aspartate Aminotransferase) (Serum/ <i>Modified IFCC</i>)	24.82	U/L	5 - 40
SGPT/ALT (Alanine Aminotransferase) (Serum/ <i>Modified IFCC</i>)	26.37	U/L	5 - 41
GGT(Gamma Glutamyl Transpeptidase) (Serum/IFCC / Kinetic)	47.77	U/L	< 38
Alkaline Phosphatase (SAP) (Serum/Modified IFCC)	117.8	U/L	42 - 98
Total Protein (Serum/Biuret)	7.16	gm/dl	6.0 - 8.0
Albumin (Serum/Bromocresol green)	4.15	gm/dl	3.5 - 5.2
Globulin (Serum/Derived)	3.01	gm/dL	2.3 - 3.6
A : G RATIO	1.38		1.1 - 2.2

(Serum/Derived)



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Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> Reference Interval
Lipid Profile			
Cholesterol Total (Serum/CHOD-PAP with ATCS)	195.72	mg/dL	Optimal: < 200 Borderline: 200 - 239 High Risk: >= 240
Triglycerides (Serum/ <i>GPO-PAP with ATCS</i>)	143.96	mg/dL	Optimal: < 150 Borderline: 150 - 199 High: 200 - 499 Very High: >=500

INTERPRETATION: The reference ranges are based on fasting condition. Triglyceride levels change drastically in response to food, increasing as much as 5 to 10 times the fasting levels, just a few hours after eating. Fasting triglyceride levels show considerable diurnal variation too. There is evidence recommending triglycerides estimation in non-fasting condition for evaluating the risk of heart disease and screening for metabolic syndrome, as non-fasting sample is more representative of the `usual_circulating level of triglycerides during most part of the day.

HDL Cholesterol (Serum/Immunoinhibition)	42.25	mg/dL	Optimal(Negative Risk Factor): >= 60 Borderline: 50 - 59 High Risk: < 50
LDL Cholesterol (Serum/ <i>Calculated</i>)	124.7	mg/dL	Optimal: < 100 Above Optimal: 100 - 129 Borderline: 130 - 159 High: 160 - 189 Very High: >=190
VLDL Cholesterol (Serum/Calculated)	28.8	mg/dL	< 30
Non HDL Cholesterol (Serum/ <i>Calculated</i>)	153.5	mg/dL	Optimal: < 130 Above Optimal: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very High: >=220



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Investigation	<u>Observed</u> <u>Value</u>	Unit Biological Reference Interval	
INTERPRETATION: 1.Non-HDL Cholesterol is now 2.It is the sum of all potentially atherogenic proteins in co-primary target for cholesterol lowering therapy.	1	r cardiovascular risk marker than LDL Cholesterol. 'LDL and chylomicrons and it is the "new bad cholesterol" and is	a
Total Cholesterol/HDL Cholesterol Ratio (Serum/ <i>Calculated</i>)	4.6	Optimal: < 3.3 Low Risk: 3.4 - 4.4 Average Risk: 4.5 - 7.1 Moderate Risk: 7.2 - 11.0 High Risk: > 11.0	
Triglyceride/HDL Cholesterol Ratio (TG/HDL) (Serum/ <i>Calculated</i>)	3.4	Optimal: < 2.5 Mild to moderate risk: 2.5 - 5.0 High Risk: > 5.0	
LDL/HDL Cholesterol Ratio (Serum/Calculated)	3	Optimal: 0.5 - 3.0 Borderline: 3.1 - 6.0 High Risk: > 6.0	



APPROVED BY

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<u>Investigation</u> <u>Glycosylated Haemoglobin (HbA1c)</u>	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
HbA1C (Whole Blood/ <i>HPLC</i>)	10.5	%	Normal: 4.5 - 5.6 Prediabetes: 5.7 - 6.4 Diabetic: >= 6.5
INTERPRETATION: If Diabetes - Good control : 6.1 - 7.0 %, Fair control : 7.1 - 8.0 %, Poor control >= 8.1 %			

Estimated Average Glucose	254.65	mg/dL
8		e

(Whole Blood)

INTERPRETATION: Comments

HbA1c provides an index of Average Blood Glucose levels over the past 8 - 12 weeks and is a much better indicator of long term glycemic control as compared to blood and urinary glucose determinations.

Conditions that prolong RBC life span like Iron deficiency anemia, Vitamin B12 & Folate deficiency,

hypertriglyceridemia, hyperbilirubinemia, Drugs, Alcohol, Lead Poisoning, Asplenia can give falsely elevated HbA1C values. Conditions that shorten RBC survival like acute or chronic blood loss, hemolytic anemia, Hemoglobinopathies, Splenomegaly, Vitamin E ingestion, Pregnancy, End stage Renal disease can cause falsely low HbA1c.



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Investigation	<u>Observed</u> Value	<u>Unit</u>	<u>Biological</u> Reference Interval
IMMUNOASSAY	<u> </u>		
<u>THYROID PROFILE / TFT</u>			
T3 (Triiodothyronine) - Total (Serum/ECLIA) INTERPRETATION: Comment : Total T3 variation can be seen in other condition like preg	1.21 gnancy, drugs, neph	ng/ml 1rosis etc. In such cas	0.7 - 2.04 es, Free T3 is recommended as it is
Metabolically active. T4 (Tyroxine) - Total (Serum/ <i>ECLIA</i>)	9.41	μg/dl	4.2 - 12.0
INTERPRETATION: Comment : Total T4 variation can be seen in other condition like preg Metabolically active.	gnancy, drugs, nepł	nrosis etc. In such cas	es, Free T4 is recommended as it is
TSH (Thyroid Stimulating Hormone) (Serum/ECLIA)	0.935	µIU/mL	0.35 - 5.50
INTERPRETATION: Reference range for cord blood - upto 20 1 st trimester: 0.1-2.5 2 nd trimester 0.2-3.0 3 rd trimester : 0.3-3.0 (Indian Thyroid Society Guidelines) Comment : 1.TSH reference range during pregnancy depends on Iodi 2.TSH Levels are subject to circadian variation, reaching of the order of 50%, hence time of the day has influence o	peak levels betwee	n 2-4am and at a min	imum between 6-10PM. The variation can be

3.Values&lt,0.03 µIU/mL need to be clinically correlated due to presence of rare TSH variant in some individuals.



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Investigation	<u>Observed</u> <u>Unit</u> <u>Value</u>	<u>Biological</u> <u>Reference Interval</u>
CLINICAL PATHOLOGY		
<u>PHYSICAL EXAMINATION (URINE</u> <u>COMPLETE)</u>		
Colour (Urine)	Yellow	Yellow to Amber
Appearance (Urine)	Clear	Clear
Volume(CLU) (Urine)	20	
<u>CHEMICAL EXAMINATION (URINE</u> <u>COMPLETE)</u>		
pH (Urine)	5.0	4.5 - 8.0
Specific Gravity (Urine)	1.029	1.002 - 1.035
Ketone (Urine)	Positive(+)	Negative
Remark: Rechecked		
Urobilinogen (Urine)	Normal	Normal
Blood (Urine)	Negative	Negative
Nitrite (Urine)	Negative	Negative
Bilirubin (Urine)	Negative	Negative
Protein (Urine)	Trace	Negative



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Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> Reference Interval
Glucose (Urine/GOD - POD)	Positive(+)		Negative
Leukocytes(CP) (Urine)	Negative		
<u>MICROSCOPIC EXAMINATION</u> (URINE COMPLETE)			
Pus Cells (Urine)	0-2	/hpf	NIL
Epithelial Cells (Urine)	0-2	/hpf	NIL
RBCs (Urine)	NIL	/hpf	NIL
Others (Urine)	NIL		

INTERPRETATION: Note: Done with Automated Urine Analyser & Automated urine sedimentation analyser. All abnormal reports are reviewed and confirmed microscopically.

Casts (Urine)	NIL	/hpf	NIL
Crystals (Urine)	NIL	/hpf	NIL



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	:



Biological Reference Interval

Investigation

IMMUNOHAEMATOLOGY

BLOOD GROUPING AND Rh TYPING (EDTA Blood/Agglutination)

'B' 'Positive'

<u>Observed</u> <u>Value</u>

Dr Anusha.K.S

<u>Unit</u>

Sr.Consultant Pathologist Reg No : 100674 APPROVED BY

The results pertain to sample tested.

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Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
BIOCHEMISTRY			
BUN / Creatinine Ratio	10.8		6.0 - 22.0
Glucose Fasting (FBS) (Plasma - F/GOD-PAP)	202.84	mg/dL	Normal: < 100 Pre Diabetic: 100 - 125 Diabetic: >= 126

INTERPRETATION: Factors such as type, quantity and time of food intake, Physical activity, Psychological stress, and drugs can influence blood glucose level.

Glucose, Fasting (Urine) (Urine - F/GOD - POD)	+		Negative
Glucose Postprandial (PPBS) (Plasma - PP/GOD-PAP)	335.09	mg/dL	70 - 140

INTERPRETATION:

Factors such as type, quantity and time of food intake, Physical activity, Psychological stress, and drugs can influence blood glucose level. Fasting blood glucose level may be higher than Postprandial glucose, because of physiological surge in Postprandial Insulin secretion, Insulin resistance, Exercise or Stress, Dawn Phenomenon, Somogyi Phenomenon, Anti- diabetic medication during treatment for Diabetes.

Urine Glucose(PP-2 hours) (Urine - PP)	++++		Negative
Blood Urea Nitrogen (BUN) (Serum/Urease UV / derived)	8.9	mg/dL	7.0 - 21
Creatinine	0.82	mg/dL	0.6 - 1.1

(Serum/Modified Jaffe)

INTERPRETATION: Elevated Creatinine values are encountered in increased muscle mass, severe dehydration, Pre-eclampsia, increased ingestion of cooked meat, consuming Protein/ Creatine supplements, Diabetic Ketoacidosis, prolonged fasting, renal dysfunction and drugs such as cefoxitin ,cefazolin, ACE inhibitors ,angiotensin II receptor antagonists,N-acetylcyteine , chemotherapeutic agent such as flucytosine etc.

Uric Acid	3.58	mg/dL	2.6 - 6.0
(Serum/Enzymatic)			



-- End of Report --



Name	Ms.Bhuvaneshwari G	ID	MED130038156
Age & Gender	49/FEMALE	Visit Date	12/11/2022
Ref Doctor Name	MediWheel		

X-ray mammogram (mediolateral oblique and craniocaudal views) followed by Sonomammography was performed. MAMMOGRAPHY OF BOTH BREASTS

Both breasts show mixed fibro glandular fatty tissue.

No evidence of focal soft tissue lesion.

No evidence of cluster micro calcification.

Subcutaneous fat deposition is within normal limits.

SONOMAMMOGRAPHY OF BOTH BREASTS

Both breasts show normal echopattern.

No evidence of focal solid / cystic areas in either breast.

No evidence of ductal dilatation.

Few lymphnodes with maintained fatty hilum are noted in both axillae.

IMPRESSION:

> NO SIGNIFICANT ABNORMALITY.

ASSESSMENT: BI-RADS CATEGORY -1

DR. APARNA CONSULTANT RADIOLOGIST A/VP

BI-RADS CLASSIFICATION CATEGORY RESULT

Assessment incomplete. Need additional imaging evaluation REPORT DISCLAIMER

1. This is only a radiologincal imperssion. Like other investigations, radiological investication also have

- limitation. Therefore radiologincal reports should be interpreted in correlation with clinical and pathological findings.
- The results reported here in are subject to interpretation by qualified medical professionals only.
 Customer identities are accepted provided by the customer or their representative.
- 4.information about the customer's condition at the time of sample collection such as fasting, food
- consumption, medication, etc are accepted as provided by the customer or representative and shall not be investigated for its truthfulness.

5.If any specimen/sample is received from any others laboratory/hospital,its is presumed that the sample belongs to the patient identified or named.

- 6.Test results should be interpreted in context of clinical and other findings if any. In case of any clarification /doubt , the refrering doctor/patient can contact the respective section head of the laboratory.
- 7.Results of the test are influenced by the various factors such as sensitivity, specificity of the procedures of the tests, quality of the samples and drug interactions etc.,

8.If the test results are found not to be correlating clinically can contact the lab in charge for clarification or retesting where practicable within 24 hours from the time of issue of results.

- 9.Liability is limited to the extend of amount billed.
- 10.Reports are subject to interpretation in their entirety.partial or selective interpretation may lead to false opinion.
- 11.Disputes, if any , with regard to the report findings are subject to the exclusive jurisdiction of the competent courts chennai only.



Name	Ms.Bhuvaneshwari G	ID	MED130038156
Age & Gender	49/FEMALE	Visit Date	12/11/2022
Ref Doctor Name	MediWheel		

1	Negative. Routine mammogram in 1 year recommended.
2	Benign finding. Routine mammogram in 1 year recommended.
3	Probably benign finding. Short interval follow-up suggested.
4	Suspicious. Biopsy should be considered.
5	Highly suggestive of malignancy. Appropriate action should be taken.

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