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 Register On
 : 23/03/2024 9:56 AM

 SID No.
 : 124005319
 Collection On
 : 23/03/2024 10:51 AM

 Age / Sex
 : 33 Year(s) / Female
 Report On
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Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
BLOOD GROUPING AND Rh TYPING (EDTA Blood/Agglutination) INTERPRETATION: Reconfirm the Blood group	'O' 'Negative' o and Typing before	blood transfusion	
Complete Blood Count With - ESR	71 0		
Haemoglobin (Whole Blood - W/Spectrophotometry)	12.7	g/dL	12.5 - 16.0
Packed Cell Volume(PCV)/Haematocrit (Whole Blood - W/Derived from Impedance)	38.7	%	37 - 47
RBC Count (Whole Blood - W/Impedance Variation)	4.49	mill/cu.mm	4.2 - 5.4
Mean Corpuscular Volume(MCV) (Whole Blood - W/Derived from Impedance)	86.4	fL	78 - 100
Mean Corpuscular Haemoglobin(MCH) (Whole Blood - W/Derived from Impedance)	28.3	pg	27 - 32
Mean Corpuscular Haemoglobin concentration(MCHC) (Whole Blood - W/Derived from Impedance)	32.8	g/dL	32 - 36
RDW-CV (Whole Blood - W/Derived from Impedance)	14.0	%	11.5 - 16.0
RDW-SD (Whole Blood - W/Derived from Impedance)	42.34	fL	39 - 46
Total Leukocyte Count (TC) (Whole Blood - W/Impedance Variation)	5700	cells/cu.mm	4000 - 11000
Neutrophils (EDTA Blood/Impedance Variation & Flow Cytometry)	52.9	%	40 - 75
Lymphocytes (EDTA Blood/Impedance Variation & Flow Cytometry)	37.3	%	20 - 45







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Eosinophils (EDTA Blood/Impedance Variation & Flow Cytometry)	0.3	%	01 - 06
Monocytes (EDTA Blood/Impedance Variation & Flow Cytometry)	8.8	%	01 - 10
Basophils (EDTA Blood/Impedance Variation & Flow Cytometry)	0.7	%	00 - 02
INTERPRETATION: Tests done on Automated F	ive Part cell counte	er. All abnormal results are revi	ewed and confirmed microscopically.
Absolute Neutrophil count (Whole Blood - W/Impedance Variation & Flow Cytometry)	3.02	10^3 / μl	1.5 - 6.6
Absolute Lymphocyte Count (Whole Blood - W/Impedance Variation & Flow Cytometry)	2.13	10^3 / μ1	1.5 - 3.5
Absolute Eosinophil Count (AEC) (Whole Blood - W/Impedance Variation & Flow Cytometry)	0.02	10^3 / μl	0.04 - 0.44
Absolute Monocyte Count (Whole Blood - W/Impedance Variation & Flow Cytometry)	0.50	10^3 / μl	< 1.0
Absolute Basophil count (Whole Blood - W/Impedance Variation & Flow Cytometry)	0.04	10^3 / μl	< 0.2
Platelet Count (Whole Blood - W/Impedance Variation)	162	10^3 / μΙ	150 - 450
MPV (Whole Blood - W/Derived from Impedance)	8.9	fL	8.0 - 13.3
PCT (Whole Blood - W/Automated Blood cell Counter)	0.14	%	0.18 - 0.28
ESR (Erythrocyte Sedimentation Rate) (Whole Blood - W/Automated - Westergren method)	14	mm/hr	< 20

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Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> Reference Interval
BUN / Creatinine Ratio	7.77		6.0 - 22.0
Glucose Fasting (FBS) (Plasma - F/GOD-PAP)	70.8	mg/dL	Normal: < 100 Pre Diabetic: 100 - 125 Diabetic: >= 126

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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		Negative
Glucose Postprandial (PPBS) (Plasma - PP/GOD-PAP)	80.2	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		Negative
Blood Urea Nitrogen (BUN) (Serum/Urease UV / derived)	7.1	mg/dL	7.0 - 21
Creatinine (Serum/Modified Jaffe)	0.90	mg/dL	0.6 - 1.1

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-acetylcysteine, chemotherapeutic agent such as flucytosine

Uric Acid (Serum/Enzymatic)	4.4	mg/dL	2.6 - 6.0
Liver Function Test			
Bilirubin(Total) (Serum/DCA with ATCS)	0.46	mg/dL	0.1 - 1.2
Bilirubin(Direct) (Serum/Diazotized Sulfanilic Acid)	0.16	mg/dL	0.0 - 0.3







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Bilirubin(Indirect) (Serum/Derived)	0.30	mg/dL	0.1 - 1.0
SGOT/AST (Aspartate Aminotransferase) (Serum/Modified IFCC)	16.8	U/L	5 - 40
SGPT/ALT (Alanine Aminotransferase) (Serum/ <i>Modified IFCC</i>)	18.3	U/L	5 - 41
GGT(Gamma Glutamyl Transpeptidase) (Serum/IFCC / Kinetic)	19.2	U/L	< 38
Alkaline Phosphatase (SAP) (Serum/Modified IFCC)	68.8	U/L	42 - 98
Total Protein (Serum/Biuret)	7.33	gm/dl	6.0 - 8.0
Albumin (Serum/Bromocresol green)	4.04	gm/dl	3.5 - 5.2
Globulin (Serum/Derived)	3.29	gm/dL	2.3 - 3.6
A : G RATIO (Serum/Derived)	1.23		1.1 - 2.2
<u>Lipid Profile</u>			
Cholesterol Total (Serum/CHOD-PAP with ATCS)	189.5	mg/dL	Optimal: < 200 Borderline: 200 - 239 High Risk: >= 240
Triglycerides (Serum/GPO-PAP with ATCS)	72.6	mg/dL	Optimal: < 150 Borderline: 150 - 199 High: 200 - 499 Very High: >= 500







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	Value		Reference Interval

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

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part of the day.			
HDL Cholesterol (Serum/Immunoinhibition)	44.9	mg/dL	Optimal(Negative Risk Factor): >= 60 Borderline: 50 - 59 High Risk: < 50
LDL Cholesterol (Serum/Calculated)	130.1	mg/dL	Optimal: < 100 Above Optimal: 100 - 129 Borderline: 130 - 159 High: 160 - 189 Very High: >= 190
VLDL Cholesterol (Serum/Calculated)	14.5	mg/dL	< 30
Non HDL Cholesterol (Serum/Calculated)	144.6	mg/dL	Optimal: < 130 Above Optimal: 130 - 159 Borderline High: 160 - 189 High: 190 - 219

Very High: ≥ 220

INTERPRETATION: 1. Non-HDL Cholesterol is now proven to be a better cardiovascular risk marker than LDL Cholesterol. 2.It is the sum of all potentially atherogenic proteins including LDL, IDL, VLDL and chylomicrons and it is the "new bad cholesterol" and is a co-primary target for cholesterol lowering therapy.

Total Cholesterol/HDL Cholesterol	4.2	Optimal: < 3.3
Ratio		Low Risk: 3.4 - 4.4
(Serum/Calculated)		Average Risk: 4.5 - 7.1
		Moderate Risk: 7.2 - 11.0
		High Risk: > 11.0

Triglyceride/HDL Cholesterol Ratio Optimal: < 2.51.6 Mild to moderate risk: 2.5 - 5.0 (TG/HDL) High Risk: > 5.0(Serum/Calculated)







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Investigation	Observed <u>Value</u>	<u>Unit</u>	<u>Biological</u> Reference Interval
LDL/HDL Cholesterol Ratio (Serum/Calculated)	2.9		Optimal: 0.5 - 3.0 Borderline: 3.1 - 6.0 High Risk: > 6.0
Glycosylated Haemoglobin (HbA1c)			
HbA1C (Whole Blood/ <i>HPLC</i>)	5.6	%	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 114.02 mg/dL

(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.

THYROID PROFILE / TFT

T3 (Triiodothyronine) - Total 0.84 ng/ml 0.7 - 2.04

(Serum/Chemiluminescent Immunometric Assay

(CLIA))

INTERPRETATION:

Comment:

Total T3 variation can be seen in other condition like pregnancy, drugs, nephrosis etc. In such cases, Free T3 is recommended as it is Metabolically active.

T4 (Tyroxine) - Total 8.43 μg/dl 4.2 - 12.0

(Serum/Chemiluminescent Immunometric Assay

(CLIA))

INTERPRETATION:

Comment:

Total T4 variation can be seen in other condition like pregnancy, drugs, nephrosis etc. In such cases, Free T4 is recommended as it is Metabolically active.







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<u>Investigation</u>	Observed Value	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
TSH (Thyroid Stimulating Hormone) (Serum/Chemiluminescent Immunometric Assay (CLIA))	30.1	μIU/mL	0.35 - 5.50

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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 Iodine intake, TPO status, Serum HCG concentration, race, Ethnicity and BMI. 2.TSH Levels are subject to circadian variation, reaching peak levels between 2-4am and at a minimum between 6-10PM. The variation can

be of the order of 50%,hence time of the day has influence on the measured serum TSH concentrations.

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

Stool Analysis - ROUTINE

Colour (Stool)	Brown	Brown
Blood (Stool)	Absent	Absent
Mucus (Stool)	Absent	Absent
Reaction (Stool)	Acidic	Acidic

Urine Analysis - Routine

COLOUR (Urine)	Pale yellow	Yellow to Amber
APPEARANCE (Urine)	Slightly turbid	Clear
Protein (Urine/Protein error of indicator)	Negative	Negative
Glucose (Urine/GOD - POD)	Negative	Negative







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Investigation	Observed <u>Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
Pus Cells (Urine/Automated - Flow cytometry)	3 - 4	/hpf	NIL
Epithelial Cells (Urine/Automated - Flow cytometry)	1 - 2	/hpf	NIL
RBCs (Urine/Automated - Flow cytometry)	NIL	/hpf	NIL
Casts (Urine/Automated - Flow cytometry)	NIL	/hpf	NIL
Crystals (Urine/Automated - Flow cytometry)	NIL	/hpf	NIL
Others (Urine)	NIL		

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

Consistency (Stool)	Semi Solid	Semi Solid
Ova (Stool)	NIL	NIL
Others (Stool)	NIL	NIL
Cysts (Stool)	NIL	NIL
Trophozoites (Stool)	NIL	NIL
RBCs (Stool)	NIL /hpf	Nil
Pus Cells (Stool)	1 - 2 /hpf	NIL
Macrophages (Stool)	NIL	NIL
Epithelial Cells (Stool)	NIL /hpf	NIL







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-- End of Report --





Name	Mrs.JAMUNA	ID	MED122518704
Age & Gender	33/FEMALE	Visit Date	23/03/2024
Ref Doctor Name	MediWheel		

ULTRA SOUND SCAN

WHOLE ABDOMEN

Liver is normal in size and shows uniform echotexture with no focal abnormality. There is no intra or extra hepatic biliary ductal dilatation. Portal vein and IVC are normal.

Gall bladder is normal sized and smooth walled. No evidence of calculi. Wall thickness is normal.

Pancreas shows a normal configuration and echotexture. Pancreatic duct is normal.

Spleen is normal in size and echotexture.

Bilateral kidneys are normal in size, shape and position. Cortical echoes are normal bilaterally. There is no calculus or calyceal dilatation.

Right kidney measures 9.3 x 4.1 cm.

Left kidney measures 9.0 x 3.6 cm.

Ureters are not dilated.

Urinary bladder is smooth walled and uniformly transonic. No intravesical mass or calculus.

Uterus is anteverted, and measures 5.9 x 3.8 cm.

Uterus shows 1.3×1.2 cm of size hypoechoic fibroid in fundal region partly subserosal, partly intramural.

Endometrial thickness is 4.5 mm.

Bilateral ovaries are normal in size.

No significant mass or cyst is seen in the ovaries.

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.
- 2. The results reported here in are subject to interpretation by qualified medical professionals only.
- 3.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.
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- $10. \\ Reports are subject to interpretation in their entirety, partial or selective interpretation may lead to false opinion.$
- 11.Disputes, if any , with regard to the report findings are subject to the exclusive jurisdiction of the competent courts chennai only.





Name	Mrs.JAMUNA	ID	MED122518704
Age & Gender	33/FEMALE	Visit Date	23/03/2024
Ref Doctor Name	MediWheel		

Parametria are free.

Iliac fossae are normal.

There is no free or loculated peritoneal fluid.

IMPRESSION:

> Small fundal fibroid.

Dr.PRASHANT MOORTHY, MBBS., MD., Consultant Radiologist

Dr. M. JAYAPRABA. Consultant Sonologist

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