Patient Name : MR. VEERANKI. VASU Age / Gender : 35 years / Male

Patient ID: 30807

Source : Corporate

Referral : Mediwhile

Sample ID :

Collection Time : Dec 09, 2023, 08:28 a.m.

Reporting Time : Dec 09, 2023, 10:03 a.m.

Test Description	Value(s)	Reference Range	Unit
CBC; Complete Blood Count			
Hemoglobin (Hb)* Method : Cynmeth Photometric Measurement	15.1	13.5 - 18.0	gm/dL
Erythrocyte (RBC) Count* Method : Electrical Impedence	5.67	4.7 - 6.0	mil/cu.mm
Packed Cell Volume (PCV)* Method : Calculated	46.5	42 - 52	%
Mean Cell Volume (MCV)* Method : Electrical Impedence	82.01	78 - 100	fL
Mean Cell Haemoglobin (MCH)* Method : Calculated	26.63	27 - 31	pg
Mean Corpuscular Hb Concn. (MCHC)* Method : Calculated	32.47	32 - 36	gm/dL
Red Cell Distribution Width (RDW)* Method : Electrical Impedence	13.8	11.5 - 14.0	%
Total Leucocytes (WBC) Count* Method : Electrical Impedence	6600	4000-10000	cell/cu.mm
Neutrophils* Method : VCSn Technology	62	40 - 80	%
Lymphocytes* Method : VCSn Technology	30	20 - 40	%
Monocytes* Method : VCSn Technology	7	2 - 10	%
Eosinophils* Method : VCSn Technology	1	1 - 6	%
Basophils	0	0 - 1	
Platelet Count* Method : Electrical Impedence	1.94	1.5 - 4.5	Lakhs/cu.mm
Mean Platelet Volume (MPV)*	7.8	7.2 - 11.7	fL

Method : Electrical Impedence

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Test Description	Value(s)	Reference Range	Unit	
DCT*	0 151	0.2 - 0.5	0/.	
Method : Calculated	0.151	0.2 - 0.5	20	
PDW*	16.1	9.0 - 17.0	%	
Method : Calculated				

Tests done on Automated Three Part Cell Counter. (WBC, RBC, Platelet count by impedance method, colorimetric method for Hemoglobin, WBC differential by flow cytometry using laser technology other parameters are calculated). All Abnormal Haemograms are reviewed confirmed microscopically.

Esr, Erythrocyte Sedimentation Rate			
Esr, Erythrocyte Sedimentation Rate	05	0-10	mm/hr
(Westergren)			
Interpretation:			

- It indicates presence and intensity of an inflammatory process. It does not diagnose a specific disease. Changes in the ESR are more significant than the abnormal results of a single test.
- It is a prognostic test and used to monitor the course or response to treatment of diseases like tuberculosis, bacterial endocarditis, acute rheumatic fever, rheumatoid arthritis, SLE, Hodgkins disease, temporal arteritis and polymyalgia rheumatica.
- It is also increased in pregnancy, multiple myeloma, menstruation, and hypothyroidism.

Urine Routine		
Colour*	Pale Yellow	
Transparency (Appearance)*	Clear	Clear
Reaction (pH)*	6.5	4.5 - 8
Specific Gravity*	1.020	1.010 - 1.030
Chemical Examination (Automated Dipstic	k Method) Urine	
Urine Glucose*	Negative	Negative
Urine Protein*	Negative	Negative
Urine Ketone*	Negative	Negative
Blood*	Negative	Negative
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Test Description	Value(s)	Reference Range	Unit
Bilirubin*	Negative	Negative	
Nitrite*	Negative	Negative	
Leucocytes*	Negative	Negative	
Urobilinogen*	Normal	Normal	
Microscopic Examination Urine			
Pus Cells (WBCs)*	2-3	0 - 5	/hpf
Epithelial Cells*	1-2	0 - 4	/hpf
Red blood Cells*	Absent	Absent	/hpf
Crystals*	Absent	Absent	
Cast*	Absent	Absent	
Bacteria*	Absent	Absent	

Stool Complete Exam

Blood Group & Rh Type

Blood Grouping & Rh Typing

Method : Forward and Reverse By Tube Method

"B" POSITIVE (+VE)

Methodology

This is done by forward and reverse grouping by tube Agglutination method.

Interpretation

Newborn baby does not produce ABO antibodies until 3 to 6 months of age. So the blood group of the Newborn baby is done by ABO antigen grouping (forward grouping) only, antibody grouping (reverse grouping) is not required.Confirmation of the New-born's blood group is indicatedwhen the A and B antigen expression and the isoagglutinins are fully developed (2–4 years).

Fasting - Glucose

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Sample ID :

		00	0000001
Test Description	Value(s)	Reference Range	Unit
Clucoso Eacting*	07.54	Normal: 70,100	ma/dl
Method - Plasma Hevokinase	97.54	Impaired Easting Glucose (IEG):	ilig/dL
method . Flasma, nexocinase		101-125	
		Diabetes Mellitus: >125	
Post Prandial Blood Sugar			
Blood Glucose-Post Prandial*	116 34	80-140	ma/dl
Method : Plasma - P, Hexokinase			
Fasting Urine Sugar			
Fasting Urine Glucose	NEGATIVE	Negative	
Post Prandial Urine Sugar			
HBA1C (Glycosylated Haemoglobin)			
Glyco Hb (HbA1C)	6.73	Non-Diabetic: <=5.9	%
Method : EDTA Whole blood, HPLC		Pre Diabetic:6.0-6.4	
		Diabetic: >=6.5	
Estimated Average Glucose :	146.45		mg/dL
Interpretations			

1. HbA1C has been endorsed by clinical groups and American Diabetes Association guidelines 2017 for diagnosing diabetes using a cut off point of 6.5%

2. Low glycated haemoglobin in a non diabetic individual are often associated with systemic inflammatory diseases, chronic anaemia (especially severe iron deficiency and haemolytic), chronic renal failure and liver diseases. Clinical correlation suggested.

3. In known diabetic patients, following values can be considered as a tool for monitoring the glycemic control.

Excellent control-6-7 % Fair to Good control - 7-8 %

Coupter

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Sample ID :



		668055697		
Test Description	Value(s)	Reference Range	Unit	
Unsatisfactory control – 8 to 10 %				
Poor Control – More than 10 %				
Thyroid Function Test (TFT)				
TRI-IODO THYRONINE (T3)	1.341	0.60 - 1.81	ng/mL	
Method : CLIA				
TOTAL THYROXINE (T4)	7.084	4.2 - 12.0	ug/dL	
Method : CLIA				
THYROID STIMULATING HORMONE (TSH)	1.953	0.46 – 8.10 : 1 Yrs – 5 Yrs	ulU/mL	
Method : CLIA		0.36 – 5.80 : 6 Yrs – 18 Yrs		
		0.35 – 5.50 : >18 Yrs		
		Pregnancy Ranges		
		1st Trimester :0.1 - 2.5		
		2nd Trimester :0.2 - 3.0		
		3rd Trimester:0.3 - 3.0		

Comments:

IF NOT ON DRUGS SUGGESTED FT3 & FT4 ESTIMATION

Please correlate with clinical conditions.

Note : Serum T3, T4 and TSH form the three components of thyroid screening panel, useful in diagnosing various disorders of the thyroid gland. Primary Hypothyroidism is accompanied by depressed serum T3 and T4 values and elevated serum TSH levels. Although elevated TSH levels are nearly always indicative of Primary Hypothyroidism, rarely they can from TSH secreting pituitary tumors (Secondary hyperthyroidism)To confirm diagnosis - evaluate FT3 and FT4.

Lipid Profile

Couptin

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Test Description	Value(s)	Reference Range	Unit
Cholesterol-Total Method : Serum, Cholesterol oxidase esterase, peroxidase	179.36	Desirable: <= 200 Borderline High: 201-239 High: > 239 Ref: The National Cholesterol Education Program (NCEP) Adult	mg/dL
Triglycerides Method : Serum, Enzymatic, endpoint	152.18	Treatment Panel III Report. Normal: < 150 Borderline High: 150-199 High: 200-499 Verv High: >= 500	mg/dL
Cholesterol-HDL Direct Method : Serum, Direct measure-PEG	35.15	<40: Low 40 - 60: Optimal > 60: Desirable	mg/dL
LDL Cholesterol Method : Serum	113.77	Optimal: < 100 Near optimal/above optimal: 100-129 Borderline high: 130-159 High: 160-189 Very High: >= 190	mg/dL
Non - HDL Cholesterol, Serum Method : calculated	144.21	Desirable: < 130 mg/dL Borderline High: 130-159mg/dL High: 160-189 mg/dL Very High: > or = 190 mg/dL	mg/dL
VLDL Cholesterol	30.44	6 - 38	mg/dL
CHOL/HDL RATIO Method : calculated	5.10	3.5 - 5.0	ratio
LDL/HDL RATIO Method : calculated	3.24	Desirable / low risk - 0.5 -3.0 Low/ Moderate risk - 3.0- 6.0 Elevated / High risk - > 6.0	ratio

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668055697

Test Description	Value(s)	Reference Range	Unit
Note: 8-10 hours fasting sample is required.			
KIDNEY FUNCTION TEST			
Urea * Method : Serum	35.4	15- 50	mg/dL
Blood Urea Nitrogen-BUN* Method : Serum, Urease	16.54	7 - 24	mg/dL
Uric Acid* Method : Serum, Uricase/POD	6.21	3.5 - 7.2	mg/dL
Creatinine* Method : Serum, Jaffe IDMS	1.18	0.7 - 1.3	mg/dL
Liver Funtion Test (LFT) with GGT			
Bilirubin - Total Method : Serum, Jendrassik Grof	0.65	0.3 - 1.2	mg/dL
Bilirubin - Direct Method : Serum, Diazotization	0.28	Adults and Children: < 0.5	mg/dL
Bilirubin - Indirect Method : Serum, Calculated	0.37	0.1 - 1.0	mg/dL
SGOT Method : Serum, UV with P5P, IFCC 37 degree	13.61	< 50	U/L
SGPT Method : Serum, UV with P5P, IFCC 37 degree	22.03	< 50	U/L
Total Protein Method : Serum. Biuret, reagent blank end point	6.79	6.6 - 8.3	g/dL
Albumin Method : Serum Bromcresol purple	4.41	Adults: 3.5 - 5.2	g/dL
Globulin Method : Calculated	2.38	1.8 - 3.6	g/dL

Chapter

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Source : Corporate		Sample ID :	668055697	
Test Description	Value(s)	Reference Range	Unit	
A/G Ratio Method : Calculated	1.85	1.2 - 2.2	ratio	

END OF REPORT

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