



LAB REPORT

Name : MR. MR. MALEKAR SHASHI KANTH
Age / Gender : 33 years / Male
Sample ID : 3808
Source : MEDIWHEEL

Referral : Dr. MEDIWHEEL
Collection Time : Sep 16, 2024, 12:23 p.m.
Receiving Time : Sep 16, 2024, 03:30 p.m.
Reporting Time : Sep 16, 2024, 03:51 p.m.

Lab Code :



HEMATOLOGY

Test Description	Value(s)	Reference Range	
CBC - Complete Blood Count			
Hemoglobin (Hb)* Method : Cynmeth Photometric Measurement	16.1	13 - 17	gm/dL
Erythrocyte (RBC) Count* Method : Electrical Impedence	5.30	4.7 - 6.0	mil/cu.mm
Packed Cell Volume (PCV)* Method : Calculated	47.2	42 - 52	%
Mean Cell Volume (MCV)* Method : Electrical Impedence	88.9	80-100	fL
Mean Cell Haemoglobin (MCH)* Method : Calculated	30.4	27 - 32	pg
Mean Corpuscular Hb Conc. (MCHC)* Method : Calculated	34.2	30-35	gm/dL
Total Leucocytes (WBC) Count* Method : Electrical Impedence	8100	4000-10000	cell/cu.mm
Neutrophils* Method : VCSn Technology	59	40 - 75	%
Lymphocytes* Method : VCSn Technology	34	20 - 40	%
Monocytes* Method : VCSn Technology	06	2 - 10	%
Eosinophils* Method : VCSn Technology	01	1 - 6	%
Basophils* Method : VCSn Technology	00	0-1	%
Platelet Count* Method : Electrical Impedence	2.11	1.50 - 4.50	lakhs/cumm

Tests done on Automated Five 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.

****END OF REPORT****

Dr. Guruprasad
Consultant Pathologist
Kmc 96510





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HEMATOLOGY

Test Description	Value(s)	Reference Range
Erythrocyte Sedimentation Rate		
Erythrocyte Sedimentation Rate	05	<15 mm/hr
Method : EDTA Whole blood, modified westergren		

Interpretation:

It indicates presence and intensity of an inflammatory process. It is a prognostic test and used to monitor the course or response to treatment of diseases like tuberculosis, acute rheumatic fever,. It is also increased in multiple myeloma, hypothyroidism.

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IMMUNOLOGY

Test Description	Value(s)	Reference Range
TFT - THYROID FUNCTION TEST		
T3 -Total Method : (Serum,CLia)	1.48	Newborn: 0.7-2.0 < 1 Year: 1.0-2.4 1 - 5 Years: 1.0-2.4 6 - 10 Years: 0.9-2.4 11 - 50 Years: 0.7-2.0 > 50 Years: 0.4-1.8 First Trimester: 0.8-1.9 Second Trimester: 1.0-2.6 ng/ml
T4-Total* Method : Serum,CLIA	8.51	Newborn: > 7.5 7 Days - 1 Year: 5.9-13.7 1 - 9 Years: 5.5-10.3 9 - 12 Years: 5.5-9.3 12 - 14 Years: 5.0-8.3 Male 15 - 60 Years: 4.6-10.5 > 60 Years: 5.0-10.7 Female 15 - 60 Years: 5.5-11.0 ug/dL
TSH-Ultrasensitive* Method : (serum/CLIA)	1.88	Newborn: < 20 4 Days - 6 Months: 0.7-4.8 6 Months - 4 Years: 0.7-4.2 5 - 20 Years: 0.5-3.4 21 - 54 Years: 0.4-4.2 55 - 87 Years: 0.5-8.9 First Trimester: 0.1-2.5 Second Trimester: 0.2-3.0 μIU/mL

Interpretation

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BIOCHEMISTRY

Test Description	Value(s)	Reference Range	
LIPID PROFILE			
Cholesterol-Total Method : Serum, Cholesterol oxidase esterase, peroxidase	148.0	Desirable: <= 200 Borderline High: 201-239 High: > 239 Ref: The National Cholesterol Education Program (NCEP) Adult Treatment Panel III Report.	mg/dL
Triglycerides Method : Serum, Enzymatic, endpoint	107.0	Normal: < 150 Borderline High: 150-199 High: 200-499 Very High: >= 500	mg/dL
Cholesterol-HDL Direct Method : Serum, Direct measure-PEG	33.7	Normal: > 40 Major Heart Risk: < 40	mg/dL
LDL Cholesterol Method : Serum	92.90	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	114.30	Desirable: < 130 mg/dL Borderline High: 130-159mg/dL High: 160-189 mg/dL Very High: > or = 190 mg/dL	mg/dL
VLDL Cholesterol Method : calculated	21.40	6 - 38	mg/dL
CHOL/HDL RATIO Method : calculated	4.39	3.5 - 5.0	ratio
LDL/HDL RATIO Method : calculated	2.76	Desirable / low risk - 0.5 -3.0 Low/ Moderate risk - 3.0- 6.0 Elevated / High risk - > 6.0	ratio
HDL/LDL RATIO Method : calculated	0.36	Desirable / low risk - 0.5 -3.0 Low/ Moderate risk - 3.0- 6.0 Elevated / High risk - > 6.0	ratio

Note: 8-10 hours fasting sample is required.

END OF REPORT

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BIOCHEMISTRY

Test Description	Value(s)	Reference Range
Glycosylated Haemoglobin (HbA1c)		
Glyco Hb (HbA1C) Method : EDTA Whole blood,HPLC	5.1	Non-Diabetic: <=5.6 % Pre Diabetic:5.7-6.4 Diabetic: >=6.5
Estimated Average Glucose :	99.67	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 glyated 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 %		
Unsatisfactory control – 8 to 10 %		
Poor Control – More than 10 %		

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Reporting Time : Sep 16, 2024, 03:46 p.m.

Lab Code :



000226024

PATHOLOGY

Test Description	Value(s)	Reference Range
<u>Creatinine, Serum</u>		
Creatinine Method : Serum, Jaffe	0.9	Children(1 yrs - 14 yrs) : 0.30 - 0.70 Adult Male : 0.72 - 1.25 Adult Female : 0.57 - 1.11

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Collection Time : Sep 16, 2024, 12:23 p.m.

Receiving Time : Sep 16, 2024, 03:30 p.m.

Reporting Time : Sep 16, 2024, 03:38 p.m.

Lab Code :



000226024

HEMATOLOGY

Investigations	Result(s)
Blood Group & Rh Typing	
Blood Group Method : Forward and Reverse By Tube Method	"B"
RH Factor	Positive
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 indicated when the A and B antigen expression and the isoagglutinins are fully developed (2-4 years).	

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CLINICAL PATHOLOGY

Test Description	Value(s)	Reference Range
Urine Complete Analysis		
Volume*	15	- ml
Colour*	Pale Yellow	Pale Yellow
Transparency (Appearance)*	Clear	Clear
Deposit*	Absent	Absent
Reaction (pH)*	5.5	4.5 - 8
Specific Gravity*	1.015	1.010 - 1.030
Chemical Examination (Automated Dipstick Method) Urine		
Urine Glucose (sugar)*	Absent	Absent
Urine Protein (Albumin)*	Absent	Absent
Urine Ketones (Acetone)*	Absent	Absent
Blood*	Absent	Absent
Bile pigments*	Absent	Absent
Nitrite*	Absent	Absent
Urobilinogen*	Normal	Normal
Microscopic Examination Urine		
Pus Cells (WBCs)*	2-4	0 - 5 /hpf
Epithelial Cells*	2-3	0 - 4 /hpf
Red blood Cells*	Absent	Absent /hpf
Crystals*	Absent	Absent
Cast*	Absent	Absent
Trichomonas Vaginalis*	Absent	Absent
Yeast Cells*	Absent	Absent
Amorphous deposits*	Absent	Absent
Bacteria*	Absent	Absent

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BIOCHEMISTRY

Test Description	Value(s)	Reference Range
Fasting Glucose- FBS		
Glucose fasting Method : Fluoride Plasma-F, Hexokinase	77	Normal: 70 - 99 Impaired Tolerance: 100-125 Diabetes mellitus: \geq 126 (on more than one occassion) (American diabetes association guidelines 2018)
Urine Fasting	Absent	

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CLINICAL BIOCHEMISTRY

Test Description	Value(s)	Reference Range	
Glucose - Post Prandial(PP)			
Blood Glucose-Post Prandial* Method : Plasma - P, Hexokinase	131	70-140	mg/dL

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CLINICAL BIOCHEMISTRY

Test Description	Value(s)	Reference Range
Urea - Serum		
Urea*	18.1	17 - 43
Method : Urease		mg/dL

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CLINICAL BIOCHEMISTRY

Test Description	Value(s)	Reference Range	
Blood Urea Nitrogen (BUN)			
UREA*	18.1	17 - 43	mg/dL
Method : Serum,Urease			
BUN*	8.46	Children 1-14yrs: 5.1- 16.8, 14-19yrs: 8.4-21, Adult Male < 50yrs : 8.9-20.6, > 50yrs: 8.4-25.7, Adult Female < 50yrs: 7.0-18.7, > 50yrs: 9.8- 20.1	mg/dL
Method : Serum,Calculated			

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CLINICAL BIOCHEMISTRY

Test Description	Value(s)	Reference Range	
Uric Acid - Serum			
Uric Acid*	7.15	3.5 - 7.2	mg/dL
Method : Uricase, POD			

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BIOCHEMISTRY

Test Description	Value(s)	Reference Range	
LFT - LIVER FUNCTION TEST			
Bilirubin - Total Method : Serum, Jendrassik Grof	0.62	0.3 - 1.2	mg/dL
Bilirubin - Direct Method : Serum, Diazotization	0.23	0.0 - 0.5	mg/dL
Bilirubin - Indirect Method : Serum, Calculated	0.39	0.1 - 1.0	mg/dL
SGOT Method : Serum, UV with P5P, IFCC 37 degree	44.4	<55	U/L
SGPT Method : Serum, UV with P5P, IFCC 37 degree	49.9	<55	U/L
SGOT/SGPT Method : calculated	0.89	0.7 - 1.4	ratio
GGT-Gamma Glutamyl Transpeptidase Method : Serum, G-glutamyl-carboxy-nitroanilide	36.3	< 55	U/L
Alkaline Phosphatase-ALPI Method : Serum, PNPP, AMP Buffer, IFCC 37 degree	78.2	30-120	U/L
Total Protein Method : Serum, Biuret, reagent blank end point	7.3	6.4 - 8.3	g/dL
Albumin Method : Serum, Bromocresol purple	4.5	3.5 - 5.2	g/dL
Globulin Method : Calculated	2.80	2.3 - 3.5	g/dL
A/G Ratio Method : Calculated	1.61	1.0 - 1.8	ratio

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CLINICAL BIOCHEMISTRY

Test Description	Value(s)	Reference Range
PSA - Prostate Specific Antigen(Total)		
PSA Method : CLIA	0.643	0-4 ng/ml

Interpretation:
Increased levels are noted in prostate cancer, benign prostatic hypertrophy, prostatitis

END OF REPORT

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