



Name : MASTER. MAYANK YADAV
Age/Gender : 2YEARS/MALE
Sample Type : WB EDTA
Reff By : C/O.SRADDHA HOSPITAL
TypedBy : Tanveer Fathima

Bill Number : M6188
Bill Date : 01-Sep-2024 02:13 PM
Sample Collection : 01-Sep-2024 02:14 PM
Sample Received : 01-Sep-2024 02:14 PM
Reporting Date : 01-Sep-2024 02:30 PM

COMPLETE BLOOD PICTURE (CBP)

INVESTIGATION	RESULT	UNITS	NORMAL RANGE
HAEMOGRAM			
HAEMOGLOBIN (Method: Cell Counter)	10.4	gm/dL	13 - 18
RBC Count (Method: Cell Counter)	5.2	Millions/Cumm	3.8 - 4.8
WBC Count (Method: Cell Counter)	6,500	Cells/Cumm	4,000 - 11,000
RDW - CV (Method: Cell Counter)	16.4	%	11.0 - 16.0
DIFFERENTIAL COUNT			
NEUTROPHILS (Method: Cell Counter)	54	%	40 - 75
LYMPHOCYTES (Method: Cell Counter)	38	%	20 - 40
EOSINOPHILS (Method: Cell Counter)	01	%	01 - 06
MONOCYTES (Method: Cell Counter)	07	%	02 - 10
BASOPHILS (Method: Cell Counter)	00	%	00 - 00
PCV (Haematocrit) (Method: Cell Counter)	31	%	35 - 45
MCV (Method: Cell Counter)	60	FL	83 - 101
MCH (Method: Cell Counter)	20	PG	27 - 32
MCHC (Method: Cell Counter)	32	%	32 - 35
PLATELET COUNT (Method: Cell Counter)	2.04	Lakhs/Cumm	1.5 - 4.5
PERIPHERAL SMEAR			
RBCs	MICROCYTIC HYPOCHROMIC		
WBCs	WITHIN NORMAL LIMITS		
PLATELETS	ADEQUATE		

Sugessted Clinical Correlation If necesarry Kindly Discuss.



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-----End of the Report-----



Authorized Signatory



LAB INCHARGE



Name : MASTER. MAYANK YADAV
Age/Gender : 2YEARS/MALE
Sample Type : SERUM
Reff By : C/O.SRADDHA HOSPITAL
TypedBy : Md Masud Ansari

Bill Number : M6188
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Sample Received : 01-Sep-2024 02:14 PM
Reporting Date : 01-Sep-2024 04:59 PM

C - REACTIVE PROTEINS (CRP)

INVESTIGATION	RESULT	UNITS	NORMAL RANGE
C – REACTIVE PROTEINS (Method: Immunoturbidimetry)	16.38	mg/L	0.0 - 6.0
INTERPRETAION	POSITIVE		

Note:

1. The CRP test is a sensitive indicator of inflammatory processes.
2. The determination of the CRP level can be used in therapy control
3. As with all the diagnostic methods, the final diagnosis should not be based on the result of a single test, but on a correlation of test results with other clinical findings.
4. The strength of agglutination is not indicative of the CRP concentration in the samples tested.

Please Correlate With Clinical Findings If Necessary Discuss

-----End of the Report-----

Authorized Signatory



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