



Name : **MASTER. SUSHANTH**
 Age/Gender : **4YEARS/MALE**
 Sample Type : **WB EDTA**
 Ref By : **C/O SRADDHA HOSPITAL**
 TypedBy : Bharat Saini

Bill Number : **M6211**
 Bill Date : 01-Sep-2024 07:52 PM
 Sample Collection : 01-Sep-2024 08:12 PM
 Sample Received : 01-Sep-2024 08:13 PM
 Reporting Date : 01-Sep-2024 09:22 PM

COMPLETE BLOOD PICTURE (CBP)

INVESTIGATION	RESULT	UNITS	NORMAL RANGE
HAEMOGRAM			
HAEMOGLOBIN (Method: Cell Counter)	10.0	g%	MALE: 13 -18 g% FEMALE: 11-13 g%
RBC Count (Method: Cell Counter)	4.2	Millions/Cumm	3.8 - 4.8
WBC Count (Method: Cell Counter)	8,300	Cells/Cumm	4,000 - 11,000
RDW-CV (Method: Cell Counter)	14.9	%	11.0 - 16.0
DIFFERENTIAL COUNT			
NEUTROPHILS (Method: Cell Counter)	44	%	Adults 40 - 75% Childrens 40 - 60 %
LYMPHOCYTES (Method: Cell Counter)	48	%	Adults 20 - 40 % Children 30 - 40 %
EOSINOPHILS (Method: Cell Counter)	01	%	Adult 01 - 06% Children 1 - 6%
MONOCYTES (Method: Cell Counter)	07	%	Adult 02 - 10% Children 6 - 10%
BASOPHILS (Method: Cell Counter)	00	%	Adults 0 - 0 % Children 0 - 0 %
PCV (Haematocrit) (Method: Cell Counter)	30	%	35.00 - 45.00 %
MCV (Method: Cell Counter)	71	FL	83 - 101 fl
MCH (Method: Cell Counter)	23	PG	27 - 32
MCHC (Method: Cell Counter)	32	%	32 - 35 %
PLATELET COUNT (Method: Cell Counter)	1.57	Lakhs/cumm	1.5 - 4.5
PERIPHERAL SMEAR			
RBCs	NORMOCYTIC HYPOCHROMIC		
WBCs	LYMPHOCYTOSIS		
PLATELETS	ADEQUATE		

Sugessted Clinical Correlation If necesarry Kindly Discuss.

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

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Authorized Signatory





LAB INCHARGE



Name : MASTER. SUSHANTH
Age/Gender : 4YEARS/MALE
Sample Type : SERUM
Reff By : C/O SRADDHA HOSPITAL
TypedBy : Bharat Saini

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C - REACTIVE PROTEINS (CRP)

INVESTIGATION	RESULT	UNITS	NORMAL RANGE
C – REACTIVE PROTEINS (Method: Immunoturbidimetry)	15.94	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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