



Name : **MASTER. ADITHYA**
 Age/Gender : **5YEARS/MALE**
 Sample Type : **WB EDTA**
 Ref By : **DR.P.GIRI**
 TypedBy : Bharat Saini

Bill Number : **M6317**
 Bill Date : 02-Sep-2024 07:36 PM
 Sample Collection : 02-Sep-2024 07:53 PM
 Sample Received : 02-Sep-2024 07:53 PM
 Reporting Date : 02-Sep-2024 08:23 PM

COMPLETE BLOOD PICTURE (CBP)

INVESTIGATION	RESULT	UNITS	NORMAL RANGE
HAEMOGRAM			
HAEMOGLOBIN (Method: Cell Counter)	12.0	gm/dL	13.0 - 18.0
RBC Count (Method: Cell Counter)	5.2	Millions/Cumm	4.0 - 5.5
WBC Count (Method: Cell Counter)	17,600	Cells/Cumm	4,000 - 12,000
RDW - CV (Method: Cell Counter)	17.4	%	11.0 - 16.0
DIFFERENTIAL COUNT			
NEUTROPHILS (Method: Cell Counter)	69	%	40 - 60
LYMPHOCYTES (Method: Cell Counter)	24	%	20 - 40
EOSINOPHILS (Method: Cell Counter)	01	%	01 - 06
MONOCYTES (Method: Cell Counter)	06	%	06 - 10
BASOPHILS (Method: Cell Counter)	00	%	00 - 00
PCV (Haematocrit) (Method: Cell Counter)	35	%	34 - 38
MCV (Method: Cell Counter)	76	FL	83 - 101
MCH (Method: Cell Counter)	27	PG	27 - 32
MCHC (Method: Cell Counter)	34	%	32 - 35
PLATELET COUNT (Method: Cell Counter)	5.15	Lakhs/Cumm	1.5 - 4.5
PERIPHERAL SMEAR			
RBCs	NORMOCYTIC NORMOCHROMIC		
WBCs	LEUCOCYTOSIS WITH NEUTROPHILIA		
PLATELETS	THROMBOCYTOSIS		

Sugessted Clinical Correlation If necesarry Kindly Discuss.

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

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Authorized Signatory



LAB INCHARGE



Name : MASTER. ADITHYA
Age/Gender : 5YEARS/MALE
Sample Type : SERUM
Reff By : DR.P.GIRI
TypedBy : Bharat Saini

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C - REACTIVE PROTEINS (CRP)

INVESTIGATION	RESULT	UNITS	NORMAL RANGE
C – REACTIVE PROTEINS (Method: Immunoturbidimetry)	17.05	mg/L	0.0 - 6.0
INTERPRETATION	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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