





Name : MR. SAIF SHAIKH Bill Number : M6054

Age/Gender : 31YEARS/MALE Bill Date : 30-Aug-2024 08:38 PM Sample Type : WB EDTA Sample Collection : 30-Aug-2024 08:45 PM Reff By : DR.P.GIRI Sample Received : 30-Aug-2024 08:53 PM TypedBy : Bharat Saini : 30-Aug-2024 09:03 PM Reporting Date

COMPLETE BLOOD PICTURE (CBP)

INVESTIGATION	RESULT	<u>UNITS</u>	NORMAL RANGE	
HAEMOGRAM HAEMOGLOBIN (Method: Cell Counter)	13.0	g%	MALE: 13 -18 g% FEMALE: 11-13 g%	
RBC Count (Method: Cell Counter)	4.4	Millions/Cumm	3.8 - 4.8	
WBC Count (Method: Cell Counter)	2,700	Cells/Cumm	4,000 - 11,000	
RDW-CV (Method: Cell Counter)	12.2	%	11.0 - 16.0	
DIFFERENTIAL COUNT				
NEUTROPHILS (Method: Cell Counter)	73	%	Adults 40 - 75% Childrens 40 - 60 %	
LYMPHOCYTES (Method: Cell Counter)	17	%	Adults 20 - 40 % Children 30 - 40 %	
EOSINOPHILS (Method: Cell Counter)	01	%	Adult 01 - 06% Children 1 - 6%	
MONOCYTES (Method: Cell Counter)	09	%	Adult 02 - 10% Children 6 - 10%	
BASOPHILS (Method: Cell Counter)	00	%	Adults 0 - 0 % Children 0 - 0 %	
PCV (Haematocrit) (Method: Cell Counter)	39	%	35.00 - 45.00 %	
MCV (Method: Cell Counter)	88	FL	83 - 101 fl	
MCH (Method: Cell Counter)	29	PG	27 - 32	
MCHC (Method: Cell Counter)	33	%	32 - 35 %	
PLATELET COUNT (Method: Cell Counter)	0.63	Lakhs/cumm	1.5 - 4.5	
PERIPHERAL SMEAR				
RBCs	NORMOCYTIC NORMOCHROMIC			
WBCs	LEUCOCY	LEUCOCYTOPENIA		

PLATELETS THROMBOCYTOPENIA

Sugessted Clinical Correlation If necesarry Kindly Discuss.







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Authorized Signatory

LAB INCHARGE





Name : MR. SAIF SHAIKH Bill Number :

Age/Gender : 31YEARS/MALE Bill Date : 30-Aug-2024 08:38 PM

Sample Type : SERUM Sample Collection : 30-Aug-2024 08:45 PM
Reff By : DR.P.GIRI Sample Received : 30-Aug-2024 08:53 PM

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C - REACTIVE PROTEINS (CRP)

INVESTIGATION RESULT UNITS NORMAL RANGE

C – REACTIVE PROTEINS 18.27 mg/L 0.0 - 6.0 (Method: Immunoturbidimetry)

INERPRETAION NEGATIVE

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 Corelate With Clinical Findings If Necessary Discuss

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

Authorized Signatory

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