

Lab Address:- # Plot No. 564, 1st floor, Buddhanagar, Near Sai Baba Temple Peerzadiguda Boduppal Hyderabad, Telangana. ICMR Reg. No. SAPALAPVLHT (Covid -19)

REPORT

Name : Mr. MD UMMED : A0012530

Age/Gender : 53 Years/Male Reg. No : 0312312300039

Referred by : Dr. DURGA PRASAD T SPP Code : SPL-CV-172

Referring Customer : V CARE MEDICAL DIAGNOSTICS Collected On : 30-Dec-2023 01:44 PM
Primary Sample : Whole Blood Received On : 30-Dec-2023 04:17 PM

Sample Tested In : Serum Reported On : 30-Dec-2023 07:58 PM

Client Address : Kimtee colony , Gokul Nagar, Tarnaka Report Status : Final Report

CLINICAL BIOCHEMISTRY

Test Name	Results	Units	Ref. Range	Method

C-Reactive protein-(CRP) 13.15 mg/L Upto:6.0 Immunoturbidimetry

Interpretation:

C-reactive protein (CRP) is produced by the liver. The level of CRP rises when there is inflammation throughout the body. It is one of a group of proteins called acute phase reactants that go up in response to inflammation. The levels of acute phase reactants increase in response to certain inflammatory proteins called cytokines. These proteins are produced by white blood cells during inflammation

A positive test means you have inflammation in the body. This may be due to a variety of conditions, including:

- Connective tissue disease
- Heart attack
- Infection
- Inflammatory bowel disease (IBD)
- Lupus
- Pneumonia
- Rheumatoid arthritis

Result rechecked and verified for abnormal cases

*** End Of Report ***







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Referring Customer : V CARE MEDICAL DIAGNOSTICS Collected On : 30-Dec-2023 01:44 PM

Primary Sample : Whole Blood Received On : 30-Dec-2023 04:17 PM Sample Tested In : Whole Blood EDTA Reported On : 30-Dec-2023 04:59 PM

Client Address : Kimtee colony ,Gokul Nagar,Tarnaka Report Status : Final Report

HAEMATOLOGY					
Test Name	Results	Units	Ref. Range	Method	
Complete Blood Picture(CBP)					
Haemoglobin (Hb)	9.6	g/dL	13-17	Cynmeth Method	
Haematocrit (HCT)	32.1	%	40-50	Calculated	
RBC Count	4.87	10^12/L	4.5-5.5	Cell Impedence	
MCV	66	fl	81-101	Calculated	
MCH	19.7	pg	27-32	Calculated	
MCHC	29.9	g/dL	32.5-34.5	Calculated	
RDW-CV	16.9	%	11.6-14.0	Calculated	
Platelet Count (PLT)	236	10^9/L	150-410	Cell Impedance	
Total WBC Count	5.2	10^9/L	4.0-10.0	Impedance	
Differential Leucocyte Count (DC)					
Neutrophils	76	%	40-70	Cell Impedence	
Lymphocytes	18	%	20-40	Cell Impedence	
Monocytes	03	%	2-10	Microscopy	
Eosinophils	03	%	1-6	Microscopy	
Basophils	0	%	1-2	Microscopy	
Absolute Neutrophils Count	3.95	10^9/L	2.0-7.0	Impedence	
Absolute Lymphocyte Count	0.94	10^9/L	1.0-3.0	Impedence	
Absolute Monocyte Count	0.16	10^9/L	0.2-1.0	Calculated	
Absolute Eosinophils Count	0.16	10^9/L	0.02-0.5	Calculated	
Absolute Basophil ICount	0.00	10^9/L	0.0-0.3	Calculated	
Morphology	Anisocytos Neutrophilia		tic hypochromic anemia and	PAPs Staining	

Result rechecked and verified for abnormal cases

*** End Of Report ***

Laboratory is NABL Accredited







Swarnabala - M DR.SWARNA BALA MD PATHOLOGY





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Primary Sample : Whole Blood EDTA Received On : 30-Dec-2023 04:17 PM
Reported On : 30-Dec-2023 05:16 PM

Client Address : Kimtee colony ,Gokul Nagar,Tarnaka Report Status : Final Report

HAEMATOLOGY

Test Name	Results	Units	Ref. Range	Method

Erythrocyte Sedimentation Rate (ESR) 18 12 or less Westergren method

Result rechecked and verified for abnormal cases

*** End Of Report ***

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Swarnabala - M DR.SWARNA BALA MD PATHOLOGY



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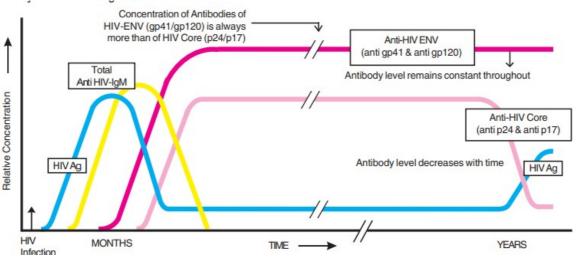
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Primary Sample : Whole Blood Received On : 30-Dec-2023 04:17 PM
Sample Tested In : Serum Reported On : 30-Dec-2023 06:49 PM

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IMMUNOLOGY & SEROLOGY				
Test Name	Results	Units	Ref. Range	Method
HIV (1& 2) Antibody	0.28	S/Co	< 1.00 : Negative > 1.00 : Positive	ELISA

Interpretation

- Non Reactive result implies that antibodies to HIV 1 / 2 have not been detected in the sample. This means the patient has either not been exposed to HIV 1 / 2 infection or the sample has been tested during the "window phase" i.e. before the development of detectable levelsof antibodies. Hence a Non Reactive result does not exclude the possibility of exposure or infection with HIV 1 / 2.
- Pre and Post test counseling to be done by the concerned referring doctor. The sensitivity and specificity of this test has been determined by National HIV Reference Centers of Govt. of India and WHO collaborating Centers, using various other test panels."
- Reactive samples by ELISA Method are confirmed by 2 other supplemental tests for confirm of HIV infection as per NACO guidelines.
- All patients' reports inderminate should be repeated with a second sample taken 14-28 days. In case the serological results continue to be inderminate the sample should be subject to western blot for confirmation.



Correlate Clinically.

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*** End Of Report ***







DR. RUTURAJ MANIKLAL KOLHAPURE MD, MICROBIOLOGIST