



41308

Laboratory Investigation Report

Name : Mr. VISHAL KUMAR CHAUDHARY

 Age / Gender
 : 30 Y / Male
 Collected
 : 23/07/2024 12:00

 Referred by
 : DR HUMAIRA
 Registered
 : 23/07/2024 14:42

 Centre
 : CITICARE MEDICAL CENTER
 Reported
 : 23/07/2024 18:29

BIOCHEMISTRY

Test Result Flag Unit Reference Range Methodology

C-REACTIVE PROTEIN (CRP) 0,7 mg/L < 5,0 Immunoturbidimetry

Please note change. Source: Roche IFU.

Ref No.

INTERPRETATION NOTES:

1. CRP measurements are used as aid in diagnosis, monitoring, prognosis, and management of suspected inflammatory disorders and associated diseases, acute infections and tissue injury.

- 2. C-reactive protein is the classic acute phase protein in inflammatory reactions.
- 3. CRP is the most sensitive of the acute phase reactants and its concentration increases rapidly during inflammatory processes. The CRP response frequently precedes clinical symptoms, including fever. After onset of an acute phase response, the serum CRP concentration rises rapidly and extensively. The increase begins within 6 to 12 hours and the peak value is reached within 24 to 48 hours. Levels above 100 mg/L are associated with severe stimuli such as major trauma and severe infection (sepsis).
- 4. CRP response may be less pronounced in patients suffering from liver disease.
- 5. CRP assays are used to detect systemic inflammatory processes (apart from certain types of inflammation such as systemic lupus erythematosus (SLE) and Colitis ulcerosa); to assess treatment of bacterial infections with antibiotics; to detect intrauterine infections with concomitant premature amniorrhexis; to differentiate between active and inactive forms of disease with concurrent infection, e.g. in patients suffering from SLE or Colitis ulcerosa; to therapeutically monitor rheumatic disease and assess anti-inflammatory therapy; to determine the presence of post-operative complications at an early stage, such as infected wounds, thrombosis and pneumonia, and to distinguish between infection and bone marrow transplant rejection.

Sample Type : Serum

End of Report

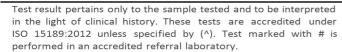
Dr. Adley Mark Fernandes Dr. Vyoma V Shah
M.D (Pathology) M.D (Pathology)
Pathologist Clinical Pathologist

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P.O Box: 49527

NAZAR MOHAMED ALI Laboratory Technologist

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Dubai, UAE



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Laboratory Investigation Report

Name : Mr. VISHAL KUMAR CHAUDHARY

 DOB
 : 06/08/1993

 Age / Gender
 : 30 Y / Male

 Referred by
 : DR HUMAIRA

Centre : CITICARE MEDICAL CENTER

Ref No. : 41308

Sample No. : 2407452476

Collected : 23/07/2024 12:00 **Registered** : 23/07/2024 14:42

Reported : 23/07/2024 16:14

HEMATOLOGY				
Test	Result Flag	Unit	Reference Range	Methodology
COMPLETE BLOOD COUNT (CBC)				
HEMOGLOBIN	15.9	g/dL	13.5 - 17.5	Spectrophotometry (Oxyhemoglobin)
RBC COUNT	5.2	10^6/μL	4.3 - 5.7	Electrical Impedance
HEMATOCRIT	46.7	%	38 - 50	Calculation
MCV	90.4	fL	82 - 98	Calculation
мсн	30.7	pg	27 - 32	Calculation
мснс	34	g/dL	32 - 37	Calculation
RDW	13.8	%	11.8 - 15.6	Calculation
RDW-SD	43.3	fL		Calculation
MPV	9.6	fL	7.6 - 10.8	Calculation
PLATELET COUNT	209	10^3/uL	150 - 450	Electrical Impedance
PCT	0.2	%	0.01 - 9.99	Calculation
PDW	17.9	Not Applicable	0.1 - 99.9	Calculation
NUCLEATED RBC (NRBC)^	0.2	/100 WBC		Flow Cytometry
ABSOLUTE NRBC COUNTA	0.02	10^3/uL		Calculation
EARLY GRANULOCYTE COUNT (EGC)^	0.2	%		Flow Cytometry
ABSOLUTE EGC^	0.0	10^3/uL		Calculation
WBC COUNT	9.7	10^3/μL	4 - 11	Electrical Impedance
DIFFERENTIAL COUNT (DC)				
NEUTROPHILS	55	%	40 - 75	Flow Cytometry
LYMPHOCYTES	38	%	20 - 45	Flow Cytometry
EOSINOPHILS	3	%	0 - 6	Flow Cytometry
MONOCYTES	4	%	1 - 6	Flow Cytometry
BASOPHILS	0	%	0 - 1	Flow Cytometry
ABSOLUTE COUNT				
ABSOLUTE NEUTROPHIL COUNT	5.3	10^3/uL	1.6 - 8.25	Calculation
ABSOLUTE LYMPHOCYTE COUNT	3.7	10^3/uL	0.8 - 4.95	Calculation
ABSOLUTE MONOCYTE COUNT	0.3	10^3/uL	0.04 - 0.66	Calculation
ABSOLUTE EOSINOPHIL COUNT	0.3	10^3/uL	0 - 0.66	Calculation
ABSOLUTE BASOPHIL COUNT	0.0	10^3/uL	0 - 0.11	Calculation

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Test result pertains only to the sample tested and to be interpreted in the light of clinical history. These tests are accredited under ISO 15189:2012 unless specified by $(^{\wedge})$. Test marked with # is performed in an accredited referral laboratory.





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Reported

Laboratory Investigation Report

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DOB 06/08/1993 Sample No. 2407452476 23/07/2024 12:00 Age / Gender 30 Y / Male **Collected** Referred by DR HUMAIRA Registered 23/07/2024 14:42 23/07/2024 16:14

CITICARE MEDICAL CENTER Centre

HEMATOLOGY

Test Result Flag Unit **Reference Range** Methodology

COMPLETE BLOOD COUNT (CBC)

INTERPRETATION NOTES: Please note update on CBC report format and changes in reference ranges.



Dr. Adley Mark Fernandes M.D (Pathology) **Pathologist**

P.O Box: 49527

Dr. Vyoma V Shah M.D (Pathology) **Clinical Pathologist**

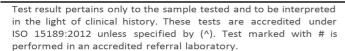
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: 06/08/1993

Age / Gender : 30 Y / Male Referred by : DR HUMAIRA

Centre : CITICARE MEDICAL CENTER

Ref No. : 41308

Sample No. : 2407452476

Collected : 23/07/2024 12:00

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Reported : 23/07/2024 16:34

HAEMATOLOGY

Test Result Flag Unit Reference Range Methodology
ERYTHROCYTE SEDIMENTATION RATE (ESR) 2 mm/hr < 15 Automated

Please note change in reference range and method.

INTERPRETATION NOTES:

DOB

Increased ESR is seen in inflammation, pregnancy, anemia, autoimmune disorders (such as rheumatoid arthritis and lupus), infections, some kidney diseases and some cancers (such as lymphoma and multiple myeloma).

The ESR is decreased in polycythemia, hyperviscosity, sickle cell anemia, leukemia, low plasma protein (due to liver or kidney disease), congestive heart failure, hypofibrinogenemia and leukocytosis.

Sample Type: EDTA Whole Blood

End of Report

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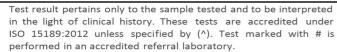
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