

Tel: 971 4 398 8567 www.biosytech.com

Mr. MUHAMMAD WAQAR

PID NO: 47623

Age: 31 Years Sex: Male DOB: 04-Apr-1994

Reference: Dr. AMAIZAH ISHTIAQ

Referred Client:

CITICARE MEDICAL CENTER

Unit G03, Al Barsha South Bldg, Al Barhsa South

Third, Dubai

VID: 5080104776

Collected on:

Registered on: 16-Aug-2025 03:43 PM

Reported on:

Abnormal Result(s) Summary

Test Name	Result Value	Unit	Reference Range	
MONOCYTES	7	%	1 - 6	
* C-REACTIVE PROTEIN (CRP)	25.9	mg/L	< 5.0 Please note change. Source: Roche IFU.	

Test Remark: Note: Please correlate clinically.

Abnormal Result(s) Summary End

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16-Aug-2025 05:40 PM

Printed on:







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CITICARE MEDICAL CENTER

Reference: Dr. AMAIZAH ISHTIAQ

Unit G03, Al Barsha South Bldg, Al Barhsa South

Third, Dubai

VID: 5080104776

Collected on:

15-Aug-2025 11:10 PM

Registered on:

16-Aug-2025 03:43 PM Reported on:

16-Aug-2025 04:37 PM

<u>Investigation</u>	Observed Value	Flag	<u>Unit</u>	Biological Reference Inte	erval <u>Method</u>
COMPLETE BLOOD COUNT (CBC)					
HEMOGLOBIN	14.7		g/dL	13.5 - 17.5	Photometric
RBC COUNT	5.1		10^6/μL	4.3 - 5.7	Electrical Impedance
HEMATOCRIT	42.9		%	38 - 50	Calculation
MCV	83.4		fL	82 - 98	Calculation
МСН	28.6		pg	27 - 32	Calculation
мснс	34.4		g/dL	32 - 37	Calculation
* RDW	12.7		%	11.8 - 15.6	Calculation
* RDW-SD	37.20		fL		Calculation
MPV	9.7		fL	7.6 - 10.8	Calculation
PLATELET COUNT	227		10^3/uL	150 - 450	Electrical Impedance
* NUCLEATED RBC (NRBC)	0.60		/100 WBC		VCS 360 Technology
* ABSOLUTE NRBC COUNT	0.04		10^3/uL		Calculation
TOTAL & DIFFERENTIAL COUNT (DC)					
WBC COUNT	6.5		10^3/μL	4 - 11	Electrical Impedance
NEUTROPHILS	67		%	40 - 75	VCS 360 Technology
LYMPHOCYTES	25		%	20 - 45	VCS 360 Technology
EOSINOPHILS	0		%	0 - 6	VCS 360 Technology
MONOCYTES	7	Н	%	1 - 6	VCS 360 Technology
BASOPHILS	1		%	0 - 1	VCS 360 Technology
ABSOLUTE COUNT					
ABSOLUTE NEUTROPHIL COUNT	4.36		10^3/uL	1.6 - 8.25	Calculation
ABSOLUTE LYMPHOCYTE COUNT	1.63		10^3/uL	0.8 - 4.95	Calculation
ABSOLUTE MONOCYTE COUNT	0.46		10^3/uL	0.04 - 0.66	Calculation
ABSOLUTE EOSINOPHIL COUNT	0		10^3/uL	0 - 0.66	Calculation
ABSOLUTE BASOPHIL COUNT	0.07		10^3/uL	0 - 0.11	Calculation

DR. ADLEY MARK FERNANDES

Sample Type: EDTA Whole Blood

DR. VYOMA SHAH M.D (Pathology) M.D (Pathology) **Pathologist**

This is an Electronically Authenticated Report.

Clinical Pathologist

Cyona V. Shah

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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 unless specified by (*).





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Investigation Observed Value Biological Reference Interval Flag Unit

25.9

* C-REACTIVE PROTEIN (CRP)

(Serum, Particle-enhanced immunoturbidimetric assay)

Note: Please correlate clinically.

< 5.0 mg/L

> Please note change. Source: Roche IFU.

INTERPRETATION:

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

- C-reactive protein is the classic acute phase protein in inflammatory reactions.
- 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).
- CRP response may be less pronounced in patients suffering from liver disease.
- 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."

----- End Of Report -----

ayana V. Shah

DR. VYOMA SHAH M.D (Pathology) **Clinical Pathologist**

This is an Electronically Authenticated Report.

DR. ADLEY MARK FERNANDES

M.D (Pathology)

Pathologist

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ACCREDITED

192 - LBPMBe 3 of 3



M RASHID CHENANGADATH

Laboratory Technologist