





Mr. ANSARI MD TABREZ

PID NO: 47503 Age: 25 Years Sex: Male

DOB: 28-Sep-1999

Reference: Dr. FARHAN

**Referred Client:** 

CITICARE MEDICAL CENTER

Unit G03, Al Barsha South Bldg, Al Barhsa South

Third, Dubai

VID: 5070110556

Collected on:

Registered on: 31-Jul-2025 03:24 PM

Reported on:

## **Abnormal Result(s) Summary**

		` '	
Test Name	Result Value	Unit	Reference Range
* C-REACTIVE PROTEIN (CRP)	20.4	mg/L	< 5.0 Please note change. Source: Roche IFU.

Test Remark: Note: Please correlate clinically.

Abnormal Result(s) Summary End

This is an Electronically Authenticated Report.



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Page 1 of 3

Test marked with # is performed in an accredited referral laboratory.

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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VID: 5070110556

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30-Jul-2025 10:00 PM

Registered on:

31-Jul-2025 03:24 PM

Reported on:

31-Jul-2025 09:40 PM

<u>Investigation</u>	Observed Value Fla	g <u>Unit</u>	Biological Reference In	terval <u>Method</u>
COMPLETE BLOOD COUNT (CBC)				
HEMOGLOBIN	15.8	g/dL	13.5 - 17.5	Photometric
RBC COUNT	5.4	10^6/μL	4.3 - 5.7	Electrical Impedance
HEMATOCRIT	45.2	%	38 - 50	Calculation
MCV	83.5	fL	82 - 98	Calculation
мсн	29.2	pg	27 - 32	Calculation
мснс	34.9	g/dL	32 - 37	Calculation
* RDW	12.4	%	11.8 - 15.6	Calculation
* RDW-SD	36.80	fL		Calculation
MPV	8.1	fL	7.6 - 10.8	Calculation
PLATELET COUNT	287	10^3/uL	150 - 450	Electrical Impedance
* NUCLEATED RBC (NRBC)	0.50	/100 WBC		VCS 360 Technology
* ABSOLUTE NRBC COUNT	0.06	10^3/uL		Calculation
<b>TOTAL &amp; DIFFERENTIAL COUNT (DC)</b>				
WBC COUNT	10.7	10^3/μL	4 - 11	Electrical Impedance
NEUTROPHILS	68	%	40 - 75	VCS 360 Technology
LYMPHOCYTES	25	%	20 - 45	VCS 360 Technology
EOSINOPHILS	4	%	0 - 6	VCS 360 Technology
MONOCYTES	3	%	1 - 6	VCS 360 Technology
BASOPHILS	0	%	0 - 1	VCS 360 Technology
ABSOLUTE COUNT				
ABSOLUTE NEUTROPHIL COUNT	7.28	10^3/uL	1.6 - 8.25	Calculation
ABSOLUTE LYMPHOCYTE COUNT	2.67	10^3/uL	0.8 - 4.95	Calculation
ABSOLUTE MONOCYTE COUNT	0.32	10^3/uL	0.04 - 0.66	Calculation
ABSOLUTE EOSINOPHIL COUNT	0.43	10^3/uL	0 - 0.66	Calculation
ABSOLUTE BASOPHIL COUNT	0	10^3/uL	0 - 0.11	Calculation

Sample Type: EDTA Whole Blood

**DR. ADLEY MARK FERNANDES** 

DR. VYOMA SHAH

ayana V. Shah

M.D (Pathology) M.D (Pathology) **Pathologist Clinical Pathologist** 

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JALADINI DULANKA

**Laboratory Technologist** 









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31-Jul-2025 05:09 PM

**Investigation** 

**Observed Value** 

20.4

**Flag** <u>Unit</u> **Biological Reference Interval** 

< 5.0 Н mg/L

> Please note change. Source: Roche IFU.

\* C-REACTIVE PROTEIN (CRP)

(Serum, Particle-enhanced immunoturbidimetric assay)

Note: Please correlate clinically.

## 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 -----

ayona V. Shah

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

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M RASHID CHENANGADATH

Laboratory Technologist

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