



Ms. NAHEEKA KAINTH

PID NO : 44392

Age : 30 Years Sex : Female

DOB: 07-Aug-1994



Reference : Dr. Sandia Bhojwani

Referred Client :

CITICARE MEDICAL CENTER

Unit G03, Al Barsha South Bldg, Al Barsha South
Third, Dubai

VID : 5070102852

Collected on :

08-Jul-2025 01:15 PM

Registered on :

08-Jul-2025 03:26 PM

Reported on :

08-Jul-2025 03:33 PM

<u>Investigation</u>	<u>Observed Value</u>	<u>Flag</u>	<u>Unit</u>	<u>Biological Reference Interval</u>	<u>Method</u>
<u>COMPLETE BLOOD COUNT (CBC)</u>					
HEMOGLOBIN	13.3		g/dL	12 - 15.5	Photometric
RBC COUNT	4.7		10 ⁶ /μL	3.9 - 5	Electrical Impedance
HEMATOCRIT	39.9		%	35 - 45	Calculation
MCV	85.4		fL	82 - 98	Calculation
MCH	28.5		pg	27 - 32	Calculation
MCHC	33.3		g/dL	32 - 37	Calculation
* RDW	13.6		%	11.9 - 15.5	Calculation
* RDW-SD	40.30		fL		Calculation
MPV	7.3	L	fL	7.6 - 10.8	Calculation
PLATELET COUNT	278		10 ³ /uL	150 - 450	Electrical Impedance
* NUCLEATED RBC (NRBC)	0.10		/100 WBC		VCS 360 Technology
* ABSOLUTE NRBC COUNT	0.01		10 ³ /uL		Calculation
<u>TOTAL & DIFFERENTIAL COUNT (DC)</u>					
WBC COUNT	6.6		10 ³ /μL	4 - 11	Electrical Impedance
NEUTROPHILS	56		%	40 - 75	VCS 360 Technology
LYMPHOCYTES	37		%	30 - 60	VCS 360 Technology
EOSINOPHILS	1		%	0 - 6	VCS 360 Technology
MONOCYTES	6		%	1 - 6	VCS 360 Technology
BASOPHILS	0		%	0 - 1	VCS 360 Technology
<u>ABSOLUTE COUNT</u>					
ABSOLUTE NEUTROPHIL COUNT	3.7		10 ³ /uL	1.6 - 8.25	Calculation
ABSOLUTE LYMPHOCYTE COUNT	2.4		10 ³ /uL	1.2 - 6.6	Calculation
ABSOLUTE MONOCYTE COUNT	0.4		10 ³ /uL	0.04 - 0.66	Calculation
ABSOLUTE EOSINOPHIL COUNT	0.1		10 ³ /uL	0 - 0.66	Calculation
ABSOLUTE BASOPHIL COUNT	0		10 ³ /uL	0 - 0.11	Calculation

Sample Type: EDTA Whole Blood

Vyoma V. Shah

DR. ADLEY MARK FERNANDES

M.D (Pathology)
Pathologist

DR. VYOMA SHAH

M.D (Pathology)
Clinical Pathologist

Christeena

CHRISTEENA FRANCIS

Laboratory Technologist

Printed on: 08-Jul-2025 06:02 PM

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





Ms. NAHEEKA KAINTH

PID NO : 44392

Age : 30 Years Sex : Female

DOB: 07-Aug-1994



Reference : Dr. Sandia Bhojwani

Referred Client :

CITICARE MEDICAL CENTER

Unit G03, Al Barsha South Bldg, Al Barsha South
Third, Dubai

VID : 5070102852

Collected on :

08-Jul-2025 01:15 PM

Registered on :

08-Jul-2025 03:26 PM

Reported on :

08-Jul-2025 06:00 PM

Investigation

Observed Value

Flag

Unit

Biological Reference Interval

*** C-REACTIVE PROTEIN (CRP)**

30.9

H

mg/L

< 5.0

(Serum, Particle-enhanced immunoturbidimetric assay)

Please note change.

Source: Roche IFU.

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 -----

Dr. Vyoma Shah

DR. ADLEY MARK FERNANDES

M.D (Pathology)
Pathologist

DR. VYOMA SHAH

M.D (Pathology)
Clinical Pathologist

This is an Electronically Authenticated Report.

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

Dr. Haleem Hakkim

HALEEM HAKKIM

Laboratory Technician

Printed on: 08-Jul-2025 06:02 PM

