



BML473871

Name : Ms. NAREEKA KAINTH

: 07/08/1994

Age / Gender : 30 Y / Female
Referred by : DR ENOMEN

Centre : CITICARE MEDICAL CENTER

Ref No. : 44392

Sample No. : 2410484373

Collected : 05/10/2024 15:31

Registered : 05/10/2024 15:31 **Reported** : 05/10/2024 17:56

BIOCHEMISTRY

Test Result Flag Unit Reference Range Methodology

C-REACTIVE PROTEIN (CRP)

3.6 mg/L < 5.0 Particle-enhanced immunoturbidimetric assay

Source: Roche IFU.

INTERPRETATION NOTES:

DOB

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 M.D (Pathology) Pathologist

This is an electronically authenticated report

P.O Box: 49527

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

Gome V. Shah

Page 1 of 4

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MOHAMMED RASHID CHENANGADATH

Laboratory Technologist
Printed on: 07/10/2024 14:41

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Dubai, UAE





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44392

Laboratory Investigation Report

HENATOLOGY

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DOB 07/08/1994 30 Y / Female Age / Gender Referred by DR ENOMEN

CITICARE MEDICAL CENTER Centre

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: 05/10/2024 16:32 Reported

HEMATOLOGY										
Test	Result	Flag	Unit	Reference Range	Methodology					
COMPLETE BLOOD COUNT (CBC)										
HEMOGLOBIN	12.9		g/dL	12 - 15.5	Photometric					
RBC COUNT	4.4		10^6/μL	3.9 - 5	Electrical Impedance					
HEMATOCRIT	37.6		%	35 - 45	Calculation					
MCV	85		fL	82 - 98	Calculation					
МСН	29.2		pg	27 - 32	Calculation					
мснс	34.3		g/dL	32 - 37	Calculation					
RDW	13.8		%	11.9 - 15.5	Calculation					
RDW-SD	40.7		fL		Calculation					
MPV	7.8		fL	7.6 - 10.8	Calculation					
PLATELET COUNT	252		10^3/uL	150 - 450	Electrical Impedance					
РСТ	0.2		%	0.01 - 9.99	Calculation					
PDW	16.8		Not Applicable	0.1 - 99.9	Calculation					
NUCLEATED RBC (NRBC)^	0.2		/100 WBC		VCS 360 Technology					
ABSOLUTE NRBC COUNT^	0.01		10^3/uL		Calculation					
EARLY GRANULOCYTE COUNT (EGC)^	0.0		%		VCS 360 Technology					
ABSOLUTE EGC^	0.0		10^3/uL		Calculation					
WBC COUNT	5.9		10^3/μL	4 - 11	Electrical Impedance					
DIFFERENTIAL COUNT (DC)										
NEUTROPHILS	73		%	40 - 75	VCS 360 Technology					
LYMPHOCYTES	20 L		%	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					
ABSOLUTE COUNT										
ABSOLUTE NEUTROPHIL COUNT	4.3		10^3/uL	1.6 - 8.25	Calculation					
ABSOLUTE LYMPHOCYTE COUNT	1.3		10^3/uL	1.2 - 6.6	Calculation					
ABSOLUTE MONOCYTE COUNT	0.5		10^3/uL	0.04 - 0.66	Calculation					
ABSOLUTE EOSINOPHIL COUNT	0.0		10^3/uL	0 - 0.66	Calculation					
ABSOLUTE BASOPHIL COUNT	0.0		10^3/uL	0 - 0.11	Calculation					

Comments: Please Correlate Clinically.

Dr. Adley Mark Fernandes

M.D (Pathology)

Gome V. Shah Dr. Vyoma V Shah

M.D (Pathology)

Pathologist Clinical Pathologist This is an electronically authenticated report

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MUBASHER ZAHOOR Laboratory Technologist Printed on: 07/10/2024 14:41

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HEMATOLOGY

End of Report

Test Result Flag Unit Reference Range Methodology

COMPLETE BLOOD COUNT (CBC)

INTERPRETATION NOTES:

Please note update on CBC report format, reference ranges and method(Beckman Coulter).

Sample Type: EDTA Whole Blood

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

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

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IMMUNOLOGY

TestResultFlagUnitReference RangeMethodologyRAPID INFLUENZA TESTRapid Influenza ANegativeNegativeImmunochromatographyRapid Influenza BNegativeNegativeImmunochromatography

Sample Type: Throat / Nasal swab



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

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Gome V. Shah

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Reena Babu Laboratory Technologist Printed on: 07/10/2024 14:41

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IMMUNOLOGY Final Report

FOOD ALLERGEN SCREEN (AlleisaScreen Panel 44 UAE)

All	Conc.	Cl (**)	Allege Breed Street	Conc.	GI (**)
Allergen Description	(*)IU/ml	Class (**)	Allergen Description	(*)IU/ml	Class(**)
CCD1Bromelain	0.00	0	Eggwhite	0.00	0
CCD2HorseradishPeroxidase	0.00	0	Eggyolk	0.00	0
CCD3AscorbatOxidase	0.00	0	Milk	10.4	3
Soybean	0.00	0	Casein	0.15	0
Peanut	0.00	0	Cheese(cow)	1.1	2
Sesameseed	0.00	0	Chicken/Mutton	0.98	2
Hazelnut	0.33	0	Duck	0.00	0
Walnut	0.22	0	Tuna	0.00	0
Cashewnut	0.14	0	Crab/Shrimp	0.00	0
Pistachionut	0.00	0	Codfish/Salmon	0.00	0
Coconut	0.14	0	Apple	0.00	0
Wheatflour	0.00	0	Kiwi	0.00	0
Gluten	0.12	0	Banana	0.04	0
Pea	0.00	0	Mango	0.00	0
BroadBean	0.00	0	Strawberry	0.00	0
GreenBean	0.00	0	Orange	0.00	0
Lentil	0.27	0	Grapes	0.00	0
Carrot	0.10	0	Date	0.00	0
Tomato	0.18	0	Olive	0.00	0
Potato	0.04	0	Cacao	0.00	0
Onion	0.00	0	BellPepper	0.00	0
Riceflour	0.00	0	Sunflowerseed	0.00	0

*CONCENTRATION (IU/ml)	**CLASS	EXPLANATION
0.00-0.34	0	No specific antibody detection.
0.35–0.69	1	Very weak antibody, frequently no clinical evidence in case of an existing sensitization.
0.70–3.49	2	Weak antibody detection, existing sensitization frequently clinical evidence in the upper range of this class.
3.50–17.49	3	Clear antibody detection, clinical evidence is mostly present.
17.5– 49.9	4	Strong antibody detection, nearly always with existing evidence.
50.0–100	5	Very strong antibody detection.
> 100	6	Extremely high antibody titer.

Sample type: Serum Method: Immunoblot

End of Report

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

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Gome V. Sheh

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SUNJO CYRILLA BERKA

Laboratory TechnicianPrinted on: 07/10/2024 14:41

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