



# **Laboratory Investigation Report**

Ms. WASANA DILHANI PANANWELAGE Name

**DOB** 28/07/1990

Age / Gender 34 Y / Female Referred by Dr. AISHA

CITICARE MEDICAL CENTER Centre

Ref No. 39063

Sample No. 2504561353

Collected Registered

10/04/2025 18:46 10/04/2025 23:14

11/04/2025 00:50 Reported

### **BIOCHEMISTRY**

Flag Unit Result Test **Reference Range** Methodology **C-REACTIVE PROTEIN (CRP)** < 5.0 Particle-enhanced 1.8 mg/L Please note change. immunoturbidimetric assay

Source: Roche IFU.

#### **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.
- 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.

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

This is an electronically authenticated report

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MOHAMMED RASHID CHENANGADATH

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Laboratory Technologist Printed on: 11/04/2025 00:52

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





P.O Box: 49527 Tel: +971 4 398 8567 Dubai, UAE reports@biosytech.ae





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15.1

Age / Gender : 34 Y / Female Referred by : Dr. AISHA

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Centre : CITICARE MEDICAL CENTER

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### IMMUNOLOGY

IU/mL

Test Result Flag Unit Reference Range Methodology

Refer to Table below in interpretation notes

ECLIA

### **INTERPRETATION NOTES:**

**IGE TOTAL ANTIBODY** 

#### Age - wise Reference Range:

Age group	IU/mL
Neonates	<1.5
Infants in 1st year of life	<15
Children aged 1 - 5 years	<60
Children aged 6 - 9 years	<90
Children aged 10 - 15 years	<200
Adults	<100
Please note change in reference range (Source: Roche)	

- 1. Immunoglobulin E (IgE) is a type of antibody synthesized by plasma cells
- 2. IgE plays an important role in immunological protection against parasitic infections and in allergy (type 1 hypersensitivity).
- 3. The IgE concentration in serum is normally very low as IgE is the least abundant antibody in serum (0.05 % of the IgG concentration). The IgE concentration is age-dependent, with the lowest values being measured at birth. Its concentration gradually increases and becomes stabilized between the age of 5-7, although the IgE values vary greatly within particular age groups.
- 4. Elevated IgE concentrations are seen in patients with Type 1 hypersensitivity reactions such as Anaphylactic reactions (reaction to drugs, bee stings, latex, vaccines, or antigen preparation used in desensitization immunotherapy), allergic diseases such as hay fever, atopic bronchitis, asthma, food allergies, urticaria and dermatitis.
- 5. Increased IgE concentrations can also occur in non-allergic diseases, e.g. congenital immunodeficiency syndromes, HIV infection, graft-versus- host disease, severe burns, some inflammatory diseases, certain cancers and parasitic diseases.
- 6. Low IgE levels may be seen in auto-immune disorders.

<u>Note:</u> Samples should not be taken from patients receiving therapy with high biotin doses (i.e. > 5 mg/day) until at least 8 hours following the last biotin administration.

#### **References:**

**Pathologist** 

- 1. Kit Insert
- 2. Dati F, Ringel KP. Reference values for serum IgE in healthy non- atopic children and adults. Clin Chem 1982;28(7):1556.
- 3. Gould HJ, Sutton BJ, Beavil AJ, Beavil RL, McCloskey N, Coker HA, et al. (2003). "The biology of IGE and the basis of allergic disease". Annual Review of Immunology. 21: 579–628

Sample Type : Serum

End of Report

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

Gome V. Shah

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P.O Box: 49527

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Dubai, UAE





reports@biosytech.ae www.biosytech.com