

# COMPLEMENT C1q DEFICIENT SERUM, Human

Product No. **C8567** Lot 83H9480

### **Precautions and Disclaimer**

For laboratory use only. Not for drug, household or other uses. POTENTIAL BIOHAZARD. Handle as if capable of transmitting infectious agents. Refer to Material Safety Data Sheet.

## Storage/Stability

Store below 0°C. After reconstitution, store at -70°C. Repeated freezing and thawing is **not** recommended.

### **Procedure**

The following procedure is used for the assay of C1q activity.

- Reconstitute the C1q deficient serum with ice cold deionized water in the amount given on the label.
- Add 20 μl of a solution containing 0.3 M CaCl<sub>2</sub> and 1 M MgCl<sub>2</sub> to each ml of C1q deficient serum.
- Prepare nine precooled assay tubes labeled "A" through "I" and two control tubes labeled "100% lysis" and "Spontaneous lysis".
- Dilute purified complement C1q (Product No. C0660) or normal serum to a concentration of 5 μg/ml complement C1q with gelatin veronal buffer (GVB<sup>2+</sup>, Product No. G6514).

# **ProductInformation**

- 5. Prepare a suspension of 1.5 x 10<sup>8</sup> cells/ml of antibody sensitized sheep erythrocytes (Product No. E9383, EA7S) in GVB<sup>2+</sup>.
- 6. Prepare reaction tubes according to Table 1.
- 7. Incubate all tubes in a 37°C water bath with shaking for 3-15 minutes.
- 8. Add 1.0 ml of ice cold GVB<sup>2+</sup> buffer immediately at the end of the incubation time.
- Centrifuge all tubes at 2,000 rpm at 0-4°C for 10 minutes.
- 10. Read the absorbance of each supernatant at 412 nm.
- Subtract the OD<sub>412</sub> of "Spontaneous lysis" tube from the OD<sub>412</sub> of each assay tube (A,B,...,I).
  These values represent OD'<sub>412</sub>. Consider the OD'<sub>412</sub> of tube B as 100% of complement activity.
- 12. The percent of normal complement activity for each tube is calculated as the OD'<sub>412</sub> of each tube divided by the OD'<sub>412</sub> of tube B.

#### **TABLE I**

I -						
Assay Tubes	GVB <sup>2+</sup> (μl)	dH <sub>2</sub> O (μl)	C1q deficient serum (µI)	Normal serum (μΙ)	Purified C1q (μΙ)	EA7S (1.5 x 10 <sup>8</sup> cells/ml) (μl)
A B C D E F G H	300 300 299 295 290 288 286 284 282	     	10  10 10 10 10 10 10	 10    	 1 5 10 12 14 16 18	200 200 200 200 200 200 200 200 200
Control tubes: 100% lysis Spontaneous lysis	310	310				200 200

### **Product Profile**

Protein: Determined by Biuret method.

Form: Lyophilized from amount of serum indicated

on the label.

Source: Prepared from freshly clotted whole blood.

C1q is depleted by the method of Kolb.<sup>1</sup>

Background activity (OD'415): 0.032

Note: Background activity should be determined at the time of assay each time complement C1q deficient serum is used. Recommended volume of C1q deficient serum: 10 µl.

### References

- 1. Kolb, *et al.*, J. Immunol., **122**, 2103 (1979)
- 2. Kabat, E.A. and Mayer, M.M., <u>Experimental</u> <u>Immunochemistry</u>, Springfield, IL, Charles C.

Thomas, 2nd edition, 149 (1961)