

# ELISA for Mouse IgA

Product Code: 3835-1AD-6

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## **CONTENTS:**

### **Vial 1 (blue top)**

Monoclonal anti-IgA antibody (300 µl)

Concentration: 0.5 mg/ml

### **Vial 2 (red top)**

ALP-conjugated anti-IgA antibody (150 µl)

### **Vial 3**

Lyophilised mouse IgA standard

To ensure total recovery of stated quantity, vials have been overfilled.

## **STORAGE:**

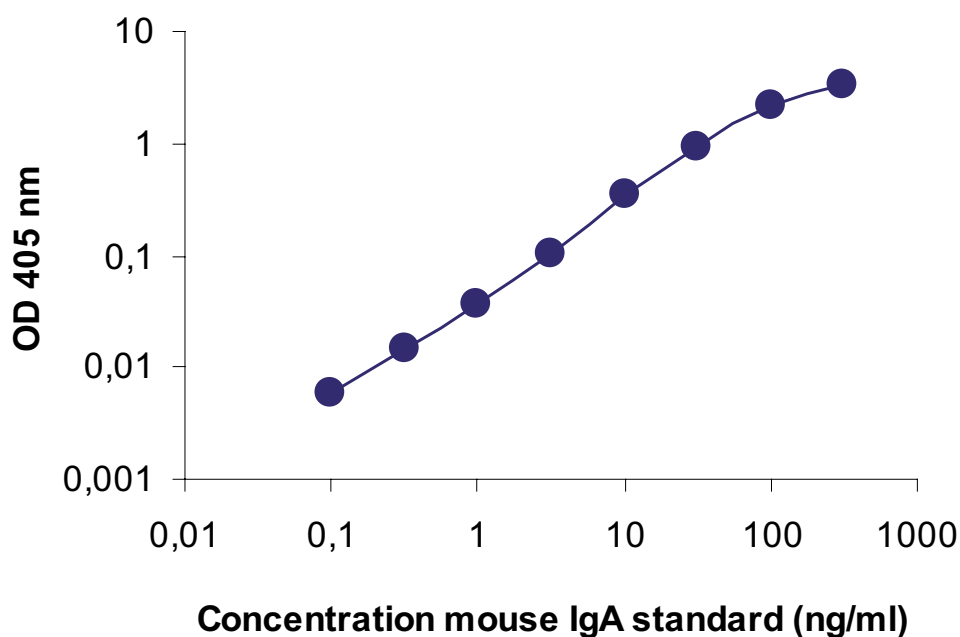
Shipped at ambient temperature. On arrival box 1 should be stored refrigerated at 4-8°C and box 2 should be stored frozen at -20°C.

# General

**Intended use:** For quantitative determination of native mouse IgA in serum and plasma.

**Reagents:** Monoclonal anti-IgA antibody is supplied in sterile-filtered (0.2  $\mu\text{m}$ ) PBS with sodium azide (0.02%). ALP-conjugated anti-IgA antibody is supplied in 0.1 M Tris-buffer with 0.15% Kathon CG.

**Recommended standard dilution:** 0.1-500 ng/ml



# Guidelines for Mouse IgA ELISA

- Day 1** 1. Coat a high protein binding ELISA plate with monoclonal anti-IgA antibody, diluted to 2 µg/ml in PBS, pH 7.4, by adding 100 µl/well. Incubate overnight at 4-8°C.
- Day 2** 2. Wash twice with PBS (200 µl/well).
3. Block plate by adding 200 µl/well of PBS with 0.05% Tween 20 (PBS-Tween) containing 0.1% BSA (incubation buffer\*). Incubate for 1 hour at room temperature.
4. Wash five times with PBS-Tween.
5. Prepare mouse IgA standard by reconstituting contents of vial 3 in 500 µl PBS to make up a stock solution of 50 µg/ml. The stock solution should be used immediately or stored in aliquots at -20°C for future use. We recommend the aliquots not be refrozen after initial use. For the test, prepare dilutions of the stock using the standard range as a guideline.
6. Add 100 µl/well of samples or standards diluted in incubation buffer and incubate for 2 hours at room temperature.
7. Wash as in step 4.
8. Add 100 µl/well of anti-IgA-ALP diluted 1:500 in incubation buffer. Incubate for 1 hour at room temperature.
9. Wash as in step 4.
10. Add 100 µl/well of appropriate substrate solution e.g. p-nitrophenyl-phosphate (pNPP).
11. Measure the optical density (405 nm for pNPP) in an ELISA reader after suitable developing time.

\* The same buffer is used for blocking and for dilution.

**NOTE; for research use only.**

MABTECH shall not be liable for the use or handling of the product or for consequential, special, indirect or incidental damages therefrom.



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Developed and manufactured by MABTECH AB, Sweden, whose quality management system complies with the following standards:



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2012-10-25