# Mouse IgA ELISA development kit

Product Code: 3865-1AD-6

**CONTENTS:** development kit for 6 plates

Vial 1 (blue top) Monoclonal MT45A antibody (300 µl) Concentration: 0.5 mg/ml

Vial 2 (red top) ALP-conjugated MT39A (80 µl)

**Vial 3** Lyophilised mouse IgA standard

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

#### **STORAGE:**

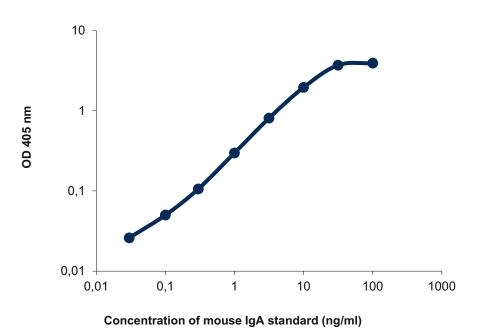
Shipped at ambient temperature. On arrival box 1 should be stored refrigerated at 4-8°C and box 2 should be stored frozen at -20°C.

### General

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

**Reagents:** MT45A is supplied in sterile-filtered (0.2  $\mu$ m) PBS with sodium azide (0.02%). MT39A-ALP is supplied in 0.1 M Tris-buffer with 0.002% Kathon CG.

Recommended standard dilution: 0.1-100 ng/ml



## Guidelines for Mouse IgA ELISA

- Day 1 1. Coat a high protein binding ELISA plate with mAb MT45A, diluted to 2  $\mu$ g/ml in PBS, pH 7.4, by adding 100  $\mu$ l/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.
  - Add 100 μl/well of MT39A-ALP diluted 1:1000 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.



Developed and manufactured by MABTECH AB, Sweden, whose quality management system complies with the standards ISO 9001:2015 & ISO 13485:2016.



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