ELISA for Mouse IL-12/-23 (p40)

Product Code: 3451-1A-20

CONTENTS, development kit for 20 plates:

Vial 1 (red top)

Monoclonal antibody C15.6 (500 µl)

Concentration: 1 mg/ml

Vial 2 (green top)

Biotinylated monoclonal antibody C17.8 (250 μ l)

Concentration: 1 mg/ml

Vial 3 (white top)

Streptavidin-Alkaline Phosphatase (250 μ l)

Vial 4

Recombinant mouse IL-12 p70 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.

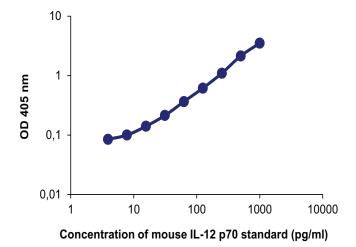
General

Intended use: For quantitative determination of native and recombinant mouse IL-12/-23 (p40) in solution, e.g. cell culture supernatant and serum/plasma samples.

Reagents: Antibodies are supplied in sterile-filtered (0.2 μ m) PBS with sodium azide (0.02%). Streptavidin-ALP is supplied in 0.1 M Tris buffer with 0.15% Kathon CG.

Standard range: 10-1000 pg/ml

Standard calibration: No international standard exists for calibration.



Guidelines for Mouse IL-12/-23 (p40) ELISA

- Day 1 1. Coat a high protein binding ELISA plate with mAb C15.6, 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 μ l/well of PBS with 0.05% Tween 20 containing 0.1% BSA (incubation buffer). Incubate for 1 hour at room temperature.
 - 4. Wash five times with PBS containing 0.05% Tween.
 - 5. Prepare mouse IL-12 p70 standard by reconstituting contents of vial 4 in 1 ml PBS to give a concentration of 0.5 µg/ml. Leave at room temperature for 15 minutes and then vortex the tube and spin down. Use immediately or store in aliquots at -20°C for future use. We recommend the aliquots not to 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 μ l/well of mAb C17.8-biotin at 0.5 μ g/ml in incubation buffer. Incubate for 1 hour at room temperature.
 - 9. Wash as in step 4.
 - 10. Add 100 μl/well of Streptavidin-ALP diluted 1:1000 in incubation buffer. Incubate for 1 hour at room temperature.
 - 11. Wash as in step 4.
 - 12. Add 100 μl/well of appropriate substrate solution e.g. p-nitrophenyl-phosphate (pNPP).
 - 13. Measure the optical density (405 nm for pNPP) in an ELISA reader after suitable developing time.

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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2013-01-24

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





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