ELISA for Mouse IL-12 (p70)

Product Code: 3456-1H-20

CONTENTS, development kit for 20 plates:

Vial 1 (green top) Monoclonal antibody C17.8 (500 µl) Concentration: 1 mg/ml

Vial 2 (blue top) Biotinylated monoclonal antibody IL12-IV (250 μl) Concentration: 1 mg/ml

Vial 3 (white top) Streptavidin-Horseradish Peroxidase (250 µl)

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.

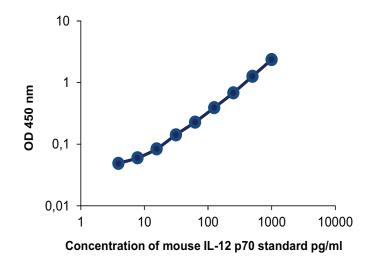
General

Intended use: For quantitative determination of native and recombinant mouse IL-12 (p70) 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-HRP is supplied in PBS with 1% BSA and 0.15% Kathon CG.

Standard range: 10-1000 pg/ml

Standard calibration: No international standard exists for calibration.



Guidelines for Mouse IL-12 (p70) ELISA

- **Day 1** 1. Coat a high protein binding ELISA plate with mAb C17.8, diluted to 1 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 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 $0.5 \mu g/ml$. 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 /well of mAb IL12-IV-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-HRP diluted 1:1000 in incubation buffer. Incubate for 1 hour at room temperature. **Please note that sodium azide used in buffers will inhibit HRP activity.**
 - 11. Wash as in step 4.
 - 12. Add 100 µl/well of appropriate substrate solution.
 - 13. Measure the optical density in an ELISA reader after suitable developing time. If required stop the reaction first.

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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