

ELISA for Mouse IL-17A

Product Code: 3521-1A-20

CONTENTS, development kit for 20 plates:

Vial 1 (yellow top)

Monoclonal antibody IL17-I (500 μ l)

Concentration: 0.5 mg/ml

Vial 2 (red top)

Biotinylated monoclonal antibody IL17-II (250 μ l)

Concentration: 0.5 mg/ml

Vial 3 (white top)

Streptavidin-Alkaline Phosphatase (250 μ l)

Vial 4

Recombinant mouse IL-17A standard (1 μ g)

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.

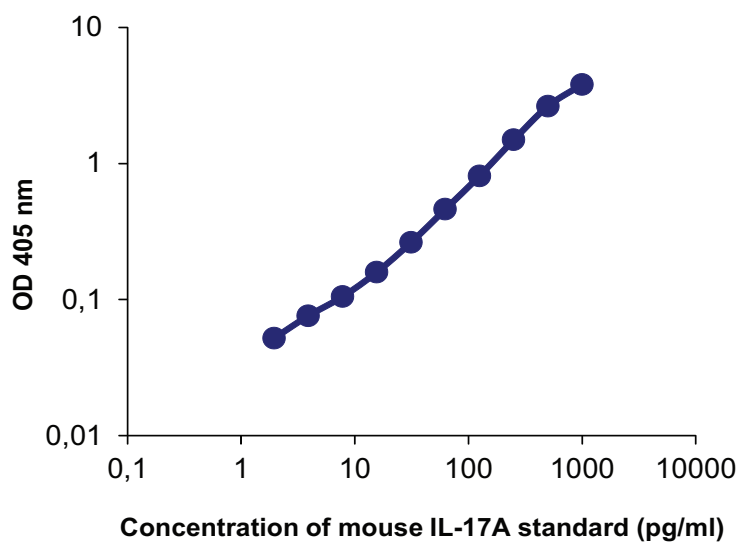
General

Intended use: For quantitative determination of native and recombinant mouse IL-17A in solution, e.g. cell culture supernatant.

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

Standard range: 5-500 pg/ml

Standard calibration: No international standard exists for calibration.



Guidelines for Mouse IL-17A ELISA

- Day 1**
1. Coat a high protein binding ELISA plate with mAb IL17-I, diluted to 1 $\mu\text{g/ml}$ in PBS, pH 7.4, by adding 100 $\mu\text{l/well}$. Incubate overnight at 4-8°C.
- Day 2**
2. Wash twice with PBS (200 $\mu\text{l/well}$).
 3. Block plate by adding 200 $\mu\text{l/well}$ 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-17A standard by reconstituting contents of vial 4 in 1 ml PBS to make a stock solution of 1 $\mu\text{g/ml}$. Allow the standard to dissolve for 5 minutes, mix thoroughly and aliquot. Store at -20°C and avoid repeated freeze-thaw cycles of the standard aliquots. For the test, prepare dilutions of the stock using the standard range as a guideline.
 6. Add 100 $\mu\text{l/well}$ 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 $\mu\text{l/well}$ of mAb IL17-II-biotin at 0.5 $\mu\text{g/ml}$ in incubation buffer or ELISA diluent for serum/plasma samples. Incubate for 1 hour at room temperature.
 9. Wash as in step 4.
 10. Add 100 $\mu\text{l/well}$ 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 $\mu\text{l/well}$ 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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Developed and manufactured by MABTECH AB, Sweden, whose quality management system complies with the following standards:



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