ELISA for Mouse IL-17A

Product Code: 3521-1H-20

CONTENTS, development kit for 20 plates:

Vial 1 (yellow top) Monoclonal antibody IL17-I (500 μl) Concentration: 0.5 mg/ml

Vial 2 (red top) Biotinylated monoclonal antibody IL17-II (250 µl) Concentration: 0.5 mg/ml

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

Vial 4 Recombinant mouse IL-17A standard (1 μg)

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 at -20°C.

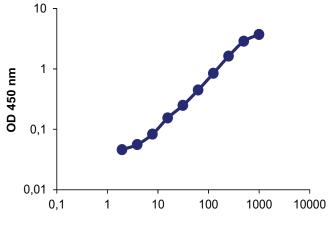
General

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

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: 5-500 pg/ml

Standard calibration: No international standard exists for calibration.



Concentration of mouse IL-17 standard (pg/ml)

Guidelines for Mouse IL-17A ELISA

- **Day 1** 1. Coat a high protein binding ELISA plate with mAb IL17-I, 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-17A standard by reconstituting contents of vial 4 in 1 ml PBS to make a stock solution of 1 μg/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 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 mAb IL17-II-biotin at 0.5 μg/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 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.

MABTECH AB Box 1233 SE-131 28 Nacka Strand Sweden Tel: +46 8 716 27 00 Fax: +46 8 716 27 01 E-mail: mabtech@mabtech.com www.mabtech.com

MABTECH Inc M.E.B. 220 3814 West Street Cincinnati, OH 45227 USA Tel: +1 513 871 4500 Fax: +1 513 871 7353 E-mail: mabtech.usa@mabtech.com

MABTECH AB Büro Deutschland Germany Tel: +49 40 4135 7935 Fax: +49 40 4135 7945 E-mail: mabtech.de@mabtech.com

2013-01-24



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



MABTECH AUSTRALIA Pty Ltd resolvingIMAGES Unit 22, 196 Settlement Road Thomastown Victoria 3074 Australia Tel: +61 3 9466 4007 Fax: +61 3 9466 4003 E-mail: mabtech.au@mabtech.com

MABTECH AB Bureau de liaison France BP 255, 1300 route des Crêtes 06905 Sophia Antipolis France Tel: +33 (0)4 92 38 80 70 Fax:+33 (0)4 92 38 80 71 E-mail: mabtech.fr@mabtech.com