

ORDERING INFORMATION

Catalog Number: AF2276

Lot Number: XJB01

Size: 100 μg

Formulation: 0.2 µm filtered solution in PBS

with 5% trehalose

Storage: -20° C

Reconstitution: sterile PBS

Specificity: mouse IL-17 RD extracellular

domain

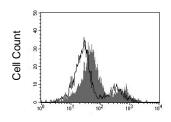
Immunogen: NS0-derived rmIL-17 RD

extracellular domain

Ig Type: goat IgG

Applications: Western blot

Immunohistochemistry Flow cytometry Direct ELISA



IL-17 RD

bEnd.3 cells were stained with anti-IL-17 RD (R&D Systems, Cat. # AF2276, filled histogram) or control antibody (R&D Systems, Cat. # AB-108-C, open histogram) followed by APC-conjugated anti-goat IgG (R&D Systems, Cat. # F0108).

Anti-mouse IL-17 RD/SEF Antibody

Preparation

Produced in goats immunized with purified, NS0-derived, recombinant mouse Interleukin 17 Receptor D (rmIL-17 RD) extracellular domain. Mouse IL-17 RD specific IgG was purified by mouse IL-17 RD affinity chromatography.

Formulation

Lyophilized from a 0.2 μm filtered solution in phosphate-buffered saline (PBS) with 5% trehalose.

Reconstitution

Reconstitute with sterile PBS. If 0.5 mL of PBS is used, the antibody concentration will be 0.2 mg/mL.

Storage

Lyophilized samples are stable for twelve months from date of receipt when stored at -20° C to -70° C. Upon reconstitution, the antibody can be stored at 2° - 8° C for 1 month without detectable loss of activity. Reconstituted antibody can also be aliquotted and stored frozen at -20° C to -70° C in a manual defrost freezer for six months without detectable loss of activity. Avoid repeated freeze-thaw cycles.

Specificity

This antibody has been selected for its ability to recognize mouse IL-17 RD in the applications listed below.

Applications

Western blot - This antibody can be used at 0.1 - 0.2 μ g/mL with the appropriate secondary reagents to detect mouse IL-17 RD. The detection limit for rmIL-17 RD is approximately 5 ng/lane under non-reducing and reducing conditions.

Immunohistochemistry - This antibody will detect IL-17 RD in cells and tissues. The working dilution is 2 - 15 μ g/mL. For chromogenic detection of labeling, use R&D Systems Cell and Tissue Staining Kits (CTS Series).

Flow cytometry - This antibody was tested in flow cytometry using bEnd.3 cells. Dilute this antibody to 25 μ g/mL and add 10 μ L of the diluted solution to 1 - 2.5 x 10 $^{\circ}$ cells in a total reaction volume not exceeding 200 μ L. The binding of unlabled antibodies may be visualized by adding a secondary developing reagent such as anti-goat lgG conjugated to a fluorochrome.

Direct ELISA - This antibody can be used at 0.5 - 1.0 μ g/mL with the appropriate secondary reagents to detect mouse IL-17 RD. The detection limit for rmIL-17 RD is approximately 0.3 ng/well. In this format, this antibody shows approximately 40% cross-reactivity with rhIL-17 RD and less than 1% cross-reactivity with rmIL-17 RC and rmIL-17B R.

Optimal dilutions should be determined by each laboratory for each application.