

Anti-IL-12R β 2 antibodies, mouse

For research use only

9 μ g equal 60 tests, 30 μ g equal 200 tests. One test corresponds to labeling of 10^6 cells.

Product	Content	Order no.
Anti-IL-12R β 2-PE	9 μ g in 300 μ L	130-105-017
Anti-IL-12R β 2-PE	30 μ g in 1 mL	130-104-965
Anti-IL-12R β 2-APC	9 μ g in 300 μ L	130-105-018
Anti-IL-12R β 2-APC	30 μ g in 1 mL	130-104-966
Anti-IL-12R β 2-PE-Vio770	9 μ g in 300 μ L	130-105-020
Anti-IL-12R β 2-PE-Vio770	30 μ g in 1 mL	130-104-968
Anti-IL-12R β 2-APC-Vio770	9 μ g in 300 μ L	130-105-019
Anti-IL-12R β 2-APC-Vio770	30 μ g in 1 mL	130-104-967
Anti-IL-12R β 2-Biotin	9 μ g in 300 μ L	130-105-016
Anti-IL-12R β 2-Biotin	30 μ g in 1 mL	130-104-964

Warnings

Reagents contain sodium azide. Under acidic conditions sodium azide yields hydrazoic acid, which is extremely toxic. Azide compounds should be diluted with running water before discarding. These precautions are recommended to avoid deposits in plumbing where explosive conditions may develop.

Technical data and background information

Antigen	IL-12R β 2
Clone	REA200
Isotype	recombinant human IgG1
Isotype control	REA Control antibodies
Alternative names of antigen	IL12RB2, IL-12RB2, Ifnm
Molecular mass of antigen [kDa]	96
Distribution of antigen	T cells
Product format	Antibodies are supplied in buffer containing stabilizer and 0.05% sodium azide.
Fixation	Cells should be stained prior to fixation, if formaldehyde is used as a fixative.
Storage	Store protected from light at 2–8 °C. Do not freeze.

Clone REA200 recognizes the mouse Interleukin-12 receptor subunit beta-2 (IL-12R β 2) antigen, a Single-pass type I membrane protein which forms the functional high affinity IL-12 receptor together with the IL-12R β 1 subunit. While the IL-12R β 1 subunit is constitutively expressed, the expression of the IL-12R β 2 gene is up-regulated by interferon gamma. IL-12R β 2 is expressed by Th1 cells and IL-12 signaling through the IL-12 receptor leads to the phosphorylation of STAT4 and continued Th1

differentiation. The IL-12R β 2 subunit plays an important role in Th1 cell differentiation, since its absence leads to an abortive Th1 differentiation that has dysfunctional production of Th1 effector molecules. The up-regulation of IL-12R β 2 is found to be associated with a number of infectious diseases, such as Crohn's disease and leprosy, which is thought to contribute to the inflammatory response and host defense.

Additional information: Clone REA200 displays negligible binding to Fc receptors.

Reagent requirements

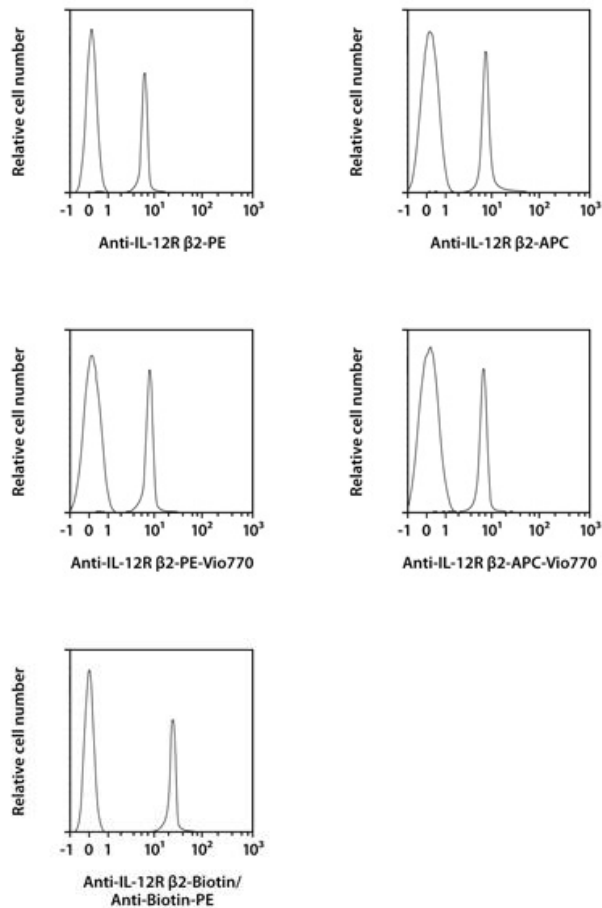
- Buffer: Prepare a solution containing phosphate-buffered saline (PBS), pH 7.2, 0.5% bovine serum albumin (BSA), and 2 mM EDTA by diluting MACS[®] BSA Stock Solution (# 130-091-376) 1:20 with autoMACS[®] Rinsing Solution (# 130-091-222). Keep buffer cold (2–8 °C).
Note: EDTA can be replaced by other supplements such as anticoagulant citrate dextrose formula-A (ACD-A) or citrate phosphate dextrose (CPD). Buffers or media containing Ca^{2+} or Mg^{2+} are not recommended for use.
- (Optional) FcR Blocking Reagent, mouse (# 130-092-575) to avoid Fc receptor-mediated antibody labeling.
- (Optional) Fluorochrome-conjugated anti-biotin antibodies, e.g., Anti-Biotin-PE (# 130-090-756) as secondary antibody reagent in combination with biotinylated antibodies.
- (Optional) Propidium Iodide Solution (# 130-093-233) for flow cytometric exclusion of dead cells without fixation.
- (Optional) Fixation and Dead Cell Discrimination Kit (# 130-091-163) for cell fixation and flow cytometric exclusion of dead cells.

Protocol for cell surface staining

- The recommended antibody dilution for labeling of cells and subsequent analysis by flow cytometry is 1:10 for up to 10^6 cells/50 μL of buffer.
 - Volumes given below are for up to 10^6 nucleated cells. When working with fewer than 10^6 cells, use the same volumes as indicated. When working with higher cell numbers, scale up all reagent volumes and total volumes accordingly (e.g. for 2×10^6 nucleated cells, use twice the volume of all indicated reagent volumes and total volumes).
1. Determine cell number.
 2. Centrifuge cell suspension at $300 \times g$ for 10 minutes. Aspirate supernatant completely.
 3. Resuspend up to 10^6 nucleated cells per 45 μL of buffer.
 4. Add 5 μL of the antibody.
 5. Mix well and incubate for 10 minutes in the dark in the refrigerator (2–8 °C).
Note: Higher temperatures and/or longer incubation times may lead to non-specific cell labeling. Working on ice requires increased incubation times.
 6. Wash cells by adding 1–2 mL of buffer and centrifuge at $300 \times g$ for 10 minutes. Aspirate supernatant completely.
 7. (Optional) If biotinylated antibody was used, resuspend the cell pellet in 100 μL of buffer, add 10 μL of fluorochrome-conjugated anti-biotin antibody, and continue as described in steps 5 and 6.
 8. Resuspend cell pellet in a suitable amount of buffer for analysis by flow cytometry or fluorescence microscopy.

Examples of immunofluorescent staining

Latex beads were coated with recombinant human IL-12 receptor subunit beta-2 protein and then stained with Anti-IL-12R β 2 antibodies or with the corresponding REA control antibodies (left peak). Flow cytometry was performed with the MACSQuant[®] Analyzer.



References

1. **Nishikomori, R. et al.** (2002) Activated STAT4 has an essential role in Th1 differentiation and proliferation that is independent of its role in the maintenance of IL-12R beta 2 chain expression and signaling. *J. Immunol.* 169(8): 4388–4398.
2. **Yin, Z. et al.** (2000) Dominance of IL-12 over IL-4 in gamma delta T cell differentiation leads to default production of IFN-gamma: failure to down-regulate IL-12 receptor beta 2-chain expression. *J. Immunol.* 164(6): 3056–3064.
3. **Presky, D. H. et al.** (1996) A functional interleukin 12 receptor complex is composed of two beta-type cytokine receptor subunits. *Proc. Natl. Acad. Sci. U.S.A.* 93(24): 14002–14007.

Warranty

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