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IL-13 Monoclonal Antibody (eBio13A), PE, eBioscience™

100 µg
Mouse
Mouse, Human
Rat / IgG1, kappa
Rat IgG1 kappa Isotype Control (eBRG1), PE, eBioscience™
Monoclonal
Antibody
eBio13A
PE
Liquid
0.2 mg/mL
Affinity chromatography
PBS, pH 7.2, with 0.1% gelatin
0.09% sodium azide
4° C, store in dark, DO NOT FREEZE!
AB_763559

Applications	Tested Dilution	Publications
Flow Cytometry (Flow)	0.25 μg/test	50 Publications

Product Specific Information

Description: The eBio13A antibody reacts with mouse IL-13. IL-13 is a cytokine produced mainly by Th2 cells, but also by antigenprimed CD8 T cells. IL-13 has a strong involvement in allergic inflammation and parasitic clearing and in cancer models has been shown to have either inhibitory or stimulatory activity depending on the tumor. In humans, IL-13 is found to play a role in isotype switching in B cells. IL-13 is implicating in down modulating macrophage activity, through the reduction of pro-inflammatory cytokines (IL-1, IL-6, IL-8, IL-10, IL-12)

Applications Reported: The eBio13A antibody has been reported useful for ELISA and ELISPOT capture, as well as intracellular staining for flow cytometric analysis.

Applications Tested: This eBio13A antibody is tested intracellular staining of cultured mouse splenocytes. It is offered in 2 formats: - μ g size: has been tested intracellular staining of cultured mouse splenocytes. This can be used at less than or equal to 0.25 μ g per test. A test is defined as the amount (μ g) of antibody that will stain a cell sample in a final volume of 100 μ L. Cell number should be determined empirically but can range from 10^5 to 10^8 cells/test. It is recommended that the antibody be carefully titrated for optimal performance in the assay of interest. - test size: has been pre-titrated and tested intracellular staining of cultured mouse splenocytes. This can be used at 5 μ L (0.125 μ g) per test. A test is defined as the amount (μ g) of antibody that will stain a cell sample in a final volume of 100 μ L. Cell number should be determined empirically but can range from 10^5 to 10^8 cells/test. Excitation: 488-561 nm; Emission: 578 nm; Laser: Blue Laser, Green Laser, Yellow-Green Laser.

Filtration: 0.2 µm post-manufacturing filtered.

O Advanced Verification Data



IL-13 Antibody (12-7133-82)

Fig. 3 cGVHD is associated with expansion of PSGL-1 lo CD4 + T cells. BALB/c recipients were irradiated (850 cGy) and given 2.5 x 10 6 TCD-BM or 2.5 x 10 6 TCD-BM plus 1 x 10 6 splenocytes from C57BL/6 donors. a, b Twenty-one, 30, and 45 days after HCT, spleen and lung were harvested. Splenocytes and mononuclear cells isolated from lung were stained for CD4, CD44, PSGL-1, and CD62L. Gated CD4 + CD44 hi are shown as PSGL-1 versus CD62L. PSGL-1 low and CD62L low cells were gated as extrafollicular CD4 + T cells. Percentages of PSGL-1 lo CD62L lo cells among CD4 + CD44 hi cells are shown as mean +- SE (n = 8). c Twenty-one days after HCT, splenocytes from no-GVHD or cGVHD recipients given wild-type C57BL/6 transplants were harvested and stained for CD4, CD44, PSGL-1, and CD62L. CD44 hi CD62L lo PSGL-1 lo CD4 + T cells were sorted and used for RNA isolation and RNA-Seg microarray analysis. Heat maps of RNA expression of CXCR4, CXCR5, and CCR7 are shown as mean centered log 2 expression, RNA-Seg microarray measurements were performed on duplicate samples from no-GVHD group and cGVHD group. Each sample represents splenocytes from eight recipients. d Twenty-one days after HCT, sorted CD4 + CD44 hi PSGL-1 lo CD62L lo cells were stimulated with PMA and ionomycin for 24 h. Stimulated cells were stained and are shown as CD4 versus IFN-gamma, IL-13, IL-17, or IL-21. Percentages of CD4 + IFNgamma + , CD4 + IL-13 + , CD4 + IL-17 + , or CD4 + IL-21 + cells among CD4 + T cells are shown a Cell treatment validation info.

Product Images For IL-13 Monoclonal Antibody (eBio13A), PE, eBioscience™



IL-13 Antibody (12-7133-82) in Flow

BALB/c splenocytes were stimulated for 3 days with plate-bound Anti-Mouse CD3e Functional Grade Purified (Product # 16-0031-82), soluble Anti-Mouse CD28 Functional Grade Purified (Product # 16-0281-82), Mouse IL-2 Recombinant Protein (Product # 14-8021-64), and Mouse IL-4 Recombinant Protein (Product # 14-8041-80). Cells were then restimulated with Cell Stimulation Cocktail (plus protein transport inhibitors) for 5 hours. Following restimulation, cells were fixed and permeabilized then stained with Anti-Mouse CD4 FITC (Product # 11-0042-82) and 0.125 µg of Rat IgG1 K Isotype Control PE (Product # 12-4301-82) or 0.125 µg of Anti-Mouse IL-13 PE. Viable cells, as determined by Fixable Viability Dye eFluor® 780, were used for analysis.

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□ 50 References

Flow Cytometry (50)

Nature communications Arf1-mediated lipid metabolism sustains cancer cells and its ablation induces anti-tumor immune responses in mice. "Published figure using IL-13 monoclonal antibody (Product # 12-7133-82) in Flow Cytometry" Authors: Wang G,Xu J,Zhao J,Yin W,Liu D,Chen W,Hou SX	Species Not Applicable Dilution Not Cited Year 2020
Frontiers in immunology Nematode-Infected Mice Acquire Resistance to Subsequent Infection With Unrelated Nematode by Inducing Highly Responsive Group 2 Innate Lymphoid Cells in the Lung. "12-7133 was used in Flow cytometry/Cell sorting to demonstrate that Strongyloides venezuelensis-experienced mice become significantly resistant against infection by Nippostrongylus brasiliensis." Authors: Yasuda K,Adachi T,Koida A,Nakanishi K	Species Mouse Dilution Not Cited Year 2019

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