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30-2837: Anti-Human IL-17A MAb (Clone: 9F9)

Clonality :	Monoclonal
Clone Name :	9F9
Application :	IP,ICC,ELISA,FACS
Reactivity :	Human
Gene :	IL17A
Gene ID :	3605
Uniprot ID :	Q16552
Format :	Purified
Alternative Name :	Interleukin 17, Interleukin 17A, IL17A, Anti-Hu IL-17A Purified
Isotype :	Mouse IgG1 kappa
Immunogen Information :	Mammalian-derived human IL-17-IgG fusion protein, boost with recombinant human IL-17A

Description

Specificity: The mouse monoclonal antibody 9F9 recognizes human interleukin 17A (IL-17A; secreted or intracellular). Interleukin 17A (IL-17A) is a proinflammatory cytokine produced by activated T cells. IL-17A-mediated downstream pathways induce the production of inflammatory molecules, chemokines, antimicrobial peptides, and remodeling proteins. It plays an important role in connecting T cell-mediated adaptive immunity and acute inflammatory response to destroy extracellular bacteria and fungi. It is the signature effector cytokine of Th17 cells, and in this role it primarily induces neutrophil activation and recruitment at infection and inflammatory sites. High levels of IL-17A are associated with rheumatoid arthritis, psoriasis, multiple sclerosis, and another inflammatory diseases, including lung injugy during severe COVID 19. This cytokine also contributes to germinal center formation by regulating the chemotactic response of B cells to CXCL12 and CXCL13, enhancing retention of B cells within the germinal centers, B cell somatic hypermutation rate and selection toward plasma cells. It is an effector cytokine for invariant NKT cells (iNKT), and it is involved in epithelial barrier formation upon injury.

Product Info

Amount :	0.1 mg
Purification :	Purified by protein-A affinity chromatography.
Content :	1 mg/ml Storage Buffer: Phosphate buffered saline (PBS), pH 7.4, 15 mM sodium azide
Storage condition :	Store at 2-8°C. Do not freeze.

Application Note

Flow cytometry: Recommended dilution: 0.5-4 µg/ml. Intracellular staining.

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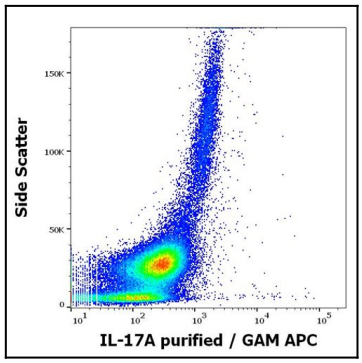

Fig 1: Flow cytometry intracellular staining pattern of human PHA stimulated and Brefeldin A treated peripheral whole blood stained using anti-human IL-17A (9F9) purified antibody (concentration in sample 0,5 Î¼g/ml, GAM APC).

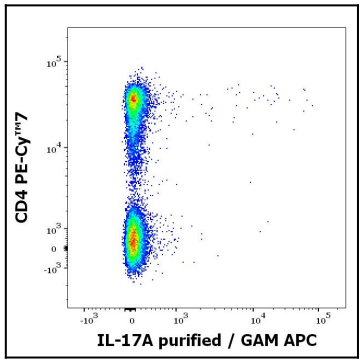

Fig 2: Flow cytometry multicolor intracellular staining of PHA stimulated and Brefeldin A treated peripheral whole blood showing lymphocytes stained using anti-human CD4 (MEM-241) PE-Cy™7 antibody (4 Î¼l reagent / 100 Î¼l of peripheral whole blood) and anti-human IL-17A (9F9) purified antibody (concentration in sample 0,5 Î¼g/ml, GAM APC).

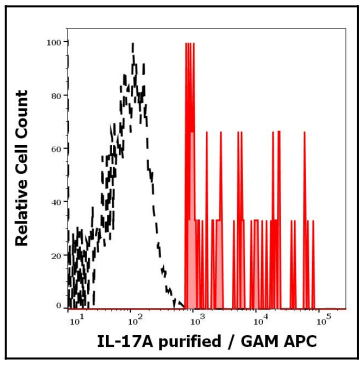

Fig 3: Separation of human CD4 positive IL-17A positive lymphocytes (red-filled) from CD4 negative IL-17A negative lymphocytes (black-dashed) in flow cytometry analysis (intracellular staining) of human PHA stimulated and Brefeldin A treated peripheral whole blood stained using anti-human IL-17A (9F9) purified antibody (concentration in sample 0,5 Î¼g/ml, GAM APC).