Technical Data Sheet

APC Mouse Anti-Human IL-4

Product Information

Material Number: 561233

Alternate Name: Interleukin-4; IL4; BCGF-1; BSF-1; Lymphocyte stimulatory factor 1

Size: 50 test Vol. per Test: 5 μ l Clone: 8D4-8

Immunogen: Recombinant Human IL-4

Isotype:Mouse IgG1, κ Reactivity:Reported: Human

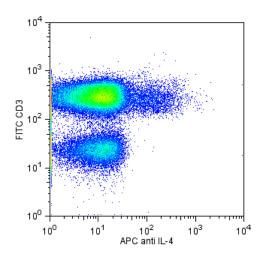
QC Testing: Rhesus or Cynomolgus

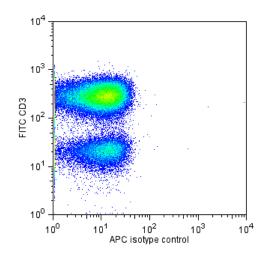
Storage Buffer: Aqueous buffered solution containing BSA and ≤0.09% sodium azide.

Description

The 8D4-8 monoclonal antibody reacts with human interleukin-4 (IL-4). The immunogen used to raise the 8D4-8 hybridoma was recombinant human IL-4. The 8D4-8 antibody binds to an epitope that is different than the epitope recognized by the MP4-25D2 antibody (Cat. No. 554485).

Clone 8D4-8 displays an increased amount of non-specific binding to dead cells when compared to the clone MP4-25D2. It is recommended to use a fixable viability dye in conjunction with this clone.





Expression of IL-4 by stimulated Rhesus macaque peripheral blood lymphocytes. Rhesus macaque peripheral blood mononuclear cells were stimulated for 6 h with Phorbol 12-Myristate 13-Acetate (PMA; Sigma, Cat. No. P-8139) and calcium ionophore A23187 (Sigma, Cat. No. C-9275) in the presence of GolgiStop™ Protein Transport Inhibitor (Cat. No. 554724). The cells were fixed using BD Cytofix™ Fixation Buffer (Cat. No. 554655) and permeabilized with BD Perm/Wash™ Buffer (Cat. No. 554723). The cells were then stained with FITC Mouse Anti-Human CD3 (Cat No. 556611) and APC Mouse anti-Human IL-4 antibody (Cat No. 561233, Left Panel), or APC Mouse IgG1 κ Isotype Control (Cat No.554681, Right Panel) by using BD Biosciences Intracellular Cytokine Staining protocol. Two-color flow cytometric dot plots were derived from gated events with the forward and side light-scatter characteristics of intact lymphocytes. Flow cytometry was performed using a BD™ LSR II Flow Cytometer System.

Preparation and Storage

Store undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

The antibody was conjugated to APC under optimum conditions, and unconjugated antibody and free APC were removed.

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Application Notes

Application

Intracellular staining (flow cytometry)	Routinely Tested	
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Suggested Companion Products

Catalog Number	Name Name	Size	<u>Clone</u>	
554724	Protein Transport Inhibitor (Containing Monensin)	0.7 ml	(none)	
554681	APC Mouse IgG1 κ Isotype Control	0.1 mg	MOPC-21	
554655	Fixation Buffer	100 ml	(none)	
554723	Perm/Wash Buffer	100 ml	(none)	
556611	FITC Mouse Anti-Human CD3ε	50 tests	SP34	

Product Notices

- This reagent has been pre-diluted for use at the recommended Volume per Test. We typically use 1 × 10⁻⁶ cells in a 100-μl experimental sample (a test).
- An isotype control should be used at the same concentration as the antibody of interest.
- Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
- Source of all serum proteins is from USDA inspected abattoirs located in the United States.
- This APC-conjugated reagent can be used in any flow cytometer equipped with a dye, HeNe, or red diode laser.
- For fluorochrome spectra and suitable instrument settings, please refer to our Multicolor Flow Cytometry web page at www.bdbiosciences.com/colors.
- 7. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.

Callard R, Gearing A. Callard R, Gearing A. The Cytokine Facts Book. San Diego: Academic Press; 1994. (Biology) Paul WE. Interleukin-4: a prototypic immunoregulatory lymphokine. Blood. 1991; 77(9):1859-1870. (Biology)

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