

FITC anti-human IL-2

Catalog # / Size: 3101525 / 50 µg
3101520 / 100 tests

Clone: MQ1-17H12

Isotype: Rat IgG2a, κ

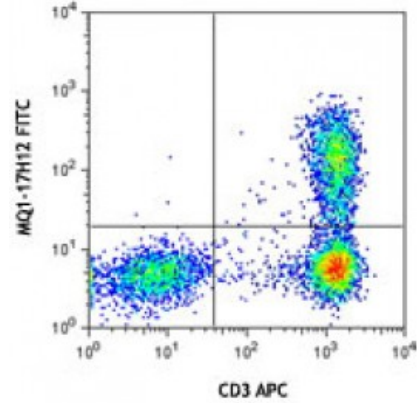
Immunogen: *E. coli*- expressed recombinant human IL-2

Reactivity: Human

Preparation: The antibody was purified by affinity chromatography, and conjugated with FITC under optimal conditions. The solution is free of unconjugated FITC.

Formulation: test size: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA).
microg size: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.

Concentration: microg sizes: 0.5 mg/ml
test sizes: lot-specific

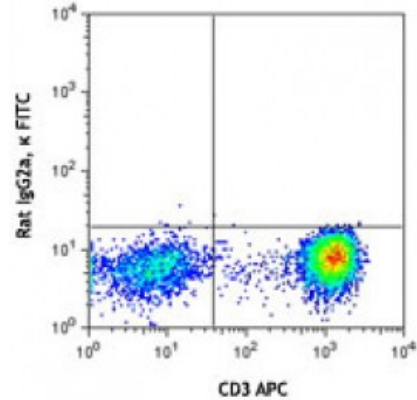


5 hours PMA + ionomycin-stimulated human peripheral blood lymphocytes (in the presence of monensin) were stained with CD3 APC, fixed, permeabilized, and then stained with IL-2 (clone MQ1-17H12) FITC (top) or rat IgG2a, κ FITC isotype control (bottom)

Applications:

Applications: Flow Cytometry

Recommended Usage: Each lot of this antibody is quality control tested by intracellular immunofluorescent staining with flow cytometric analysis. **Test size products are transitioning from 20 microL to 5 microL per test.** Please check your vial or your CoA to find the suggested use of this reagent per million cells in 100 microL staining volume or per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.



Application Notes: **ELISA or ELISPOT Capture^{2,3}:** The purified MQ1-17H12 antibody is useful as the capture antibody in a sandwich ELISA or ELISPOT assay, when used in conjunction with the biotinylated Poly5176 antibody (Cat. No. 517605) as the detecting antibody. The LEAF™ purified antibody is suggested for ELISPOT capture. For ELISPOT capture applications, a concentration range of 4-8 microg/ml is recommended.
Additional reported applications (for the relevant formats) include: immunoprecipitation², immunohistochemical staining of paraformaldehyde-fixed, saponin-

treated frozen tissue sections^{1,4-6,8},
neutralization¹³, and
immunocytochemistry.

Note: For testing human IL-2 in serum
or plasma, BioLegend's LEGEND MAX™
Kits (Cat. No. 431807 & 431808) are
specially developed and recommended.

- Application**
- References:**
1. Andersson J, *et al.* 1994. *Immunology* 83:16. (IHC)
 2. Abrams J, *et al.* 1992. *Immunol. Rev.* 127:5. (IP)
 3. Abrams JS. 1995. *Curr. Prot. Immunol.* Unit 6.20.
 4. Fernandez V, *et al.* 1994. *Eur. J. Immunol.* 24:1808. (IHC)
 5. Skansen-Saphir U, *et al.* 1994. *Eur. J. Immunol.* 24:916. (IHC)
 6. Andersson U, *et al.* *Detection and Quantification of Gene Expression.* New York:Springer-Verlag. (IHC)
 7. Prussin C, *et al.* 1995. *J. Immunol. Methods.* 188:117.
 8. Raqib R, *et al.* 2002. *Infect. Immun.* 70:3199. (IHC)
 9. Dzhagalov I, *et al.* 2007. *J. Immunol.* 178:2113. [PubMed](#)
 10. Colleton BA, *et al.* 2009. *J Virol.* 83:6288. [PubMed](#)
 11. Yoshino N, *et al.* 2000. *Exp. Anim. (Tokyo)* 49:97. (FC)
 12. Rout N, *et al.* 2010. *PLoS One* 5:e9787. (FC)
 13. Wu Z, *et al.* 2015. *J Virol.* 89:6435. [PubMed](#)
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Description: IL-2 is a potent lymphoid cell growth factor which exerts its biological activity primarily on T cells, promoting proliferation and maturation. Additionally, IL-2 has been found to stimulate growth and differentiation of B cells, NK cells, LAK cells, monocytes, and oligodendrocytes.

- Antigen**
- References:**
1. Fitzgerald K, *et al.* Eds. 2001. *The Cytokine FactsBook.* Academic Press, San Diego.
 2. Taniguchi T, *et al.* 1993. *Cell* 73:5.
 3. Nistico G. 1993. *Prog. Neurobiol.* 40:463.
 4. Waldmann T, *et al.*