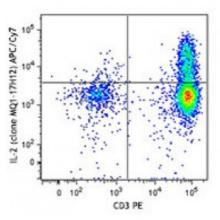
Product Data Sheet

APC/Cy7 anti-human IL-2

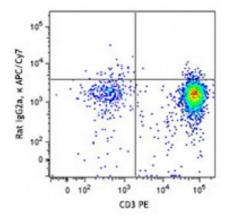
Catalog # / Size:	3101705 / 25 tests 3101710 / 100 tests
Clone:	MQ1-17H12
Isotype:	Rat IgG2a, к
Immunogen:	<i>E. coli</i> - expressed recombinant human IL-2
Reactivity:	Human
Preparation:	The antibody was purified by affinity chromatography and conjugated with APC/Cy7 under optimal conditions. The solution is free of unconjugated APC/Cy7 and unconjugated antibody.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA).
Concentration:	Lot-specific



Human peripheral blood lymphocytes were stimulated with PMA + Ionomycin for 6 hours (in the presence of monensin), surface stained with CD3 PE, fixed, permeabilized, and then stained with IL-2 (clone MQ1-17H12) APC/Cy7 (top) or rat IgG2a, κ APC/Cy7

Applications:

Applications:	Flow Cytometry
Recommended Usage:	Each lot of this antibody is quality control tested by intracellular immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is 5 microL per million cells or 5 microL 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} ,



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	neutralization ¹³ , 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:	 Andersson J, <i>et al.</i> 1994. <i>Immunology</i> 83:16. (IHC) Abrams J, <i>et al.</i> 1992. <i>Immunol. Rev.</i> 127:5. (IP) Abrams JS. 1995. <i>Curr. Prot. Immunol.</i> Unit 6.20. Fernandez V, <i>et al.</i> 1994. <i>Eur. J. Immunol.</i> 24:1808. (IHC) Skansen-Saphir U, <i>et al.</i> 1994. <i>Eur. J. Immunol.</i> 24:916. (IHC) Andersson U, <i>et al.</i> Detection and Quantification of Gene Expression. New York:Springer-Verlag. (IHC) Prussin C, <i>et al.</i> 1995. <i>J. Immunol. Methods.</i> 188:117. Raqib R, <i>et al.</i> 2002. <i>Infect. Immunol.</i> 178:2113. PubMed Colleton BA, <i>et al.</i> 2009. <i>J Virol.</i> 83:6288. PubMed Yoshino N, <i>et al.</i> 2000. <i>Exp. Anim. (Tokyo)</i> 49:97. (FC) Rout N, <i>et al.</i> 2010. <i>PLoS One</i> 5:e9787. (FC)
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.

Antigen1. Fitzgerald K, et al. Eds. 2001. The Cytokine FactsBook. Academic Press, SanReferences:Diego.

- 2. Taniguchi T, et al. 1993. Cell 73:5.
- 3. Nistico G. 1993. Prog. Neurobiol. 40:463.
- 4. Waldmann T, et al.