

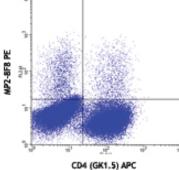
The path to legendary discovery™

## **Product Data Sheet**

## PE anti-mouse IL-3

Catalog # / Size:	503903 / 25 μg 503904 / 100 μg		*
Clone:	MP2-8F8		
Isotype:	Rat IgG1, κ		
Immunogen:	COS-expressed, recombinant mouse IL-3	ш	155
Reactivity:	Mouse	F8 P	
Preparation:	The antibody was purified by affinity chromatography, and conjugated with PE under optimal conditions. The solution is free of unconjugated PE and unconjugated antibody.	MP2-E	-
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.		in.
Concentration:	0.2 mg/ml		-1022
Storage:	The antibody solution should be stored undiluted at 4°C and protected from prolonged exposure to light. <b>Do not freeze</b> .		

## **Applications:**



PMA+lonomycin stimulated Th2-polarized C57BL/6 splenocytes surface stained with CD4 (GK1.5) APC and intracellularly stained with MP2-8F8 PE

Applications:	ICFC - Quality tested	MP2-8F8 PE			
Recommended Usage:	Each lot of this antibody is quality control tested by intracellular immunofluorescent staining with flow cytometric analysis. For immunofluorescent staining, the suggested use of this reagent is $\leq 0.25 \ \mu g$ per 10 <sup>6</sup> cells in 100 $\mu$ l volume. It is recommended that the reagent be titrated for optimal performance for each application.				
Application Notes:	<ul> <li>ELISA or ELISPOT Capture<sup>1-5,8</sup>: The purified MP2-8F8 antibody is useful as the capture antibody in a sandwich ELISA or ELISPOT assay, when used in conjunction with the biotinylated MP2-43D11 antibody (Cat. No. 504002) as the detecting antibody. The LEAF™ purified antibody is suggested for ELISPOT capture.</li> <li>Flow Cytometry<sup>1</sup>: The fluorochrome-labeled MP2-8F8 antibody is useful for intracellular immunofluorescent staining and flow cytometric analysis to identify IL-3 -producing cells within mixed cell populations. For intracellular cytokine staining protocol, please visit www.biolegend.com and click on the support section.</li> <li>Neutralization<sup>1-3,6</sup>: The LEAF™ purified antibody (Endotoxin &lt;0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for neutralization of IL-3 bioactivity <i>in vivo</i> and <i>in vitro</i> (Cat. No. 503906).</li> <li>Additional reported applications (for the relevant formats) include: immunoprecipitation, Western blotting, immunohistochemical staining<sup>4,7</sup> of paraformaldehyde-fixed, saponin-treated frozen tissue sections, and immunocytochemistry.</li> </ul>				
Application References:	<ol> <li>Abrams, J., et al. 1992. Immunol. Rev. 127:5.</li> <li>Abrams, J., et al. 1988. J. Immunol. 140:131.</li> <li>Cockayne, D., et al. 1991. Growth Factors 5:171.</li> <li>Sander, B., et al. 1993. J. Immunol. Meth. 166:201.</li> <li>Abrams, J. 1995. Curr. Prot. Immunol. John Wiley and Sons, New York. Unit 6.20.</li> <li>Finkelman, F., et al. 1993. J. Immunol. 151:1235.</li> <li>Andersson, U., et al. 1999. Detection and quantification of gene expression. New York:Springer-Verlag.</li> <li>Karulin, A., et al. 2000. J. Immunol. 164:1862.</li> </ol>				
Description:	IL-3 is a highly species-specific pleiotropic factor produced primarily by activated T cells though also by mast cells keratinocytes, and astrocytes, which stimulates colony formation of megakaryocytes, neutrophils, and macrophages from bone marrow cultures. The MP2-8F8 antibody reacts with mouse interleukin-3 (IL-3). The MP2-8F8 antibody can neutralize the bioactivity of natural or recombinant IL-3.				
Antigen References:	<ol> <li>Fitzgerald, K., et al. Eds. 2001. The Cytokine FactsBook. Academic Press, San Diego.</li> <li>Frendl, G., 1992. Int. J. Immunopharmacol. 14:421.</li> <li>Ihle, J., 1992. Chem. Immunology 51:65.</li> <li>Valen T, P., et al. 1990. Blut 61:338.</li> </ol>				
Related Products	<b>Product</b> PE Rat IgG1, κ Isotype Ctrl Cell Staining Buffer Fixation Buffer Permeabilization Wash Buffer (10X) Brefeldin A Solution (1,000X) Monensin Solution (1,000X) RBC Lysis Buffer (10X)	<b>Clone</b> RTK2071	Application FC, ICFC FC, ICC, ICFC ICC, ICFC ICC, ICFC, IHC ICFC ICFC FC, ICFC		



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