## **Product Data Sheet**

## FITC anti-mouse GM-CSF

Catalog # / Size:	3127015 / 25 μg 3127020 / 100 μg
Clone:	MP1-22E9
Isotype:	Rat IgG2a, к
Immunogen:	Yeast-expressed, recombinant mouse GM-CSF
<b>Reactivity:</b>	Mouse
Preparation:	The antibody was purified by affinity chromatography, and conjugated with FITC under optimal conditions. The solution is free of unconjugated FITC.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.
<b>Concentration:</b>	0.5

## **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 $\leq 0.25$ microg per $10^6$ cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.
Application Notes:	<ul> <li>ELISA or ELISPOT Capture<sup>1,3-5</sup>: The purified MP1-22E9 antibody is useful as the capture antibody in a sandwich ELISA or ELISPOT assay, when used in conjunction with the biotinylated MP1-31G6 antibody (Cat. No. 505502) as the detecting antibody. The LEAF™ purified antibody is suggested for ELISPOT capture.</li> <li>Flow Cytometry<sup>8</sup>: The fluorochrome-labeled MP1-22E9 antibody is useful for intracellular immunofluorescent staining and flow cytometric analysis to identify GM-CSF -producing cells within mixed cell populations.</li> <li>Neutralization<sup>2-4</sup>: The MP1-22E9 antibody can neutralize the bioactivity of natural or recombinant GM-CSF. The LEAF™ purified antibody (Endotoxin &lt;0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for <i>in vivo</i> and <i>in vitro</i> neutralization (Cat. No. 505408).</li> <li>Additional reported applications (for the relevant formats) include: immunohistochemical staining of paraformaldehyde-fixed, saponintreated frozen tissue sections<sup>1,6,7</sup>, and immunocytochemistry<sup>8</sup>.</li> </ul>
Application References:	<ol> <li>Sander B, et al. 1993. J. Immunol. Methods 166:201.</li> <li>Suda T, et al. 1990. Cell. Immunol. 129:228.</li> <li>Nozaki S, et al. 1991. J. Invest. Dermatol. 97:10.</li> <li>Abrams JS, et al. 1992. Immunol. Rev. 127:5.</li> <li>Abrams JS. 2001. Curr. Protoc. Immunol. Unit 6.20.</li> <li>Sander B, et al. 1991. Immunol. Rev. 119:65.</li> <li>Andersson U, et al. 1999. Detection and quantification of gene expression. New York:Springer-Verlag.</li> <li>Larkin J, et al. 2006. J. Immunol. 177:268.</li> <li>Lee PH, et al. 2014. J. Immunol. 192:178. PubMed</li> </ol>

For research use only. Not for diagnostic use. Not for resale. Sony Biotechnology Inc. will not be held responsible for patent infringement or other violations that may occur with the use of our products. Sony Biotechnology Inc. 1730 North First Street, San Jose, CA 95112 www.sonybiotechnology.com **Description:** GM-CSF is a hematopoietic factor that is produced by T cells, macrophages, fibroblasts and endothelial cells. This multifunctional cytokine stimulates progenitor cells of neutrophils, eosinophils, and macrophages. GM-CSF is also a differentiation and activating factor for granulocytic and monocytic cells.

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

- 2. Demetri, G., et al. 1991. Blood 78:2791.
- 3. Fan, D., *et al.* 1991. *In vivo* 5:571.
- 4. Negrin, R.,