

Human GM-CSF Rα APC-conjugated Antibody

Monoclonal Mouse IgG₁ Clone # 31916

Catalog Number: FAB706A

100 TESTS

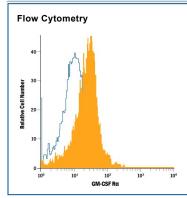
DESCRIPTION			
Species Reactivity	Human		
Specificity	Detects human GM-CSF R α in direct ELISAs and Western blots. In direct ELISAs and Western blots, no cross-reactivity with recombinant human (rh) M-CSF R α 0 or rhGM-CSF R α 1 (α 1) is observed.		
Source	Monoclonal Mouse IgG ₁ Clone # 31916		
Purification	Protein A or G purified from ascites		
Immunogen	S. frugiperda insect ovarian cell line Sf 21-derived recombinant human GM-CSF $Rlpha$ Extracellular domain		
Conjugate	Allophycocyanin Excitation Wavelength: 620-650 nm Emission Wavelength: 660-670 nm		
Formulation	Supplied in a saline solution containing BSA and Sodium Azide. See Certificate of Analysis for details.		
	*Contains <0.1% Sodium Azide, which is not hazardous at this concentration according to GHS classifications. Refer to the Safety Data Shee (SDS) for additional information and handling instructions.		

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Flow Cytometry	10 μL/10 ⁶ cells	See Below

DATA



Detection of GM-CSF Rα in Human Blood Monocytes by Flow Cytometry. Human peripheral blood monocytes were stained with Mouse Anti-Human GM-CSF Rα APC-conjugated Monoclonal Antibody (Catalog # FAB706A, filled histogram) or isotype control antibody (Catalog # IC002A, open histogram). View our protocol for Staining Membraneassociated Proteins.

PREPARATION AND STORAGE

Shipping The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.

Stability & Storage

Protect from light. Do not freeze.

• 12 months from date of receipt, 2 to 8 °C as supplied.





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BACKGROUND

Granulocyte Macrophage Colony Stimulating Factor Receptor alpha (GM-CSF R α), also known as CD116, is a component of the receptor complex that mediates cellular responses to GM-CSF. GM-CSF promotes the differentiation and mobilization of granulocyte-macrophage, erythroid, megakaryocyte, and eosinophil progenitors. It enhances the activation of myeloid cell effector functions and plays a role in the development of Th1 biased immune responses, allergic inflammation, and autoimmunity (1–4). Mature human GM-CSF R α is an 80 kDa type I transmembrane glycoprotein that consists of a 298 amino acid (aa) extracellular domain (ECD) with two fibronectin type III domains and a juxtamembrane WSXWS motif, a 26 aa transmembrane segment, and a 54 aa cytoplasmic domain (5). Within the ECD, human GM-CSF R α shares approximately 33% aa sequence identity with mouse and rat GM-CSF R α . Alternate splicing of human GM-CSF R α generates several additional isoforms that lack the cytoplasmic and/or transmembrane regions. Soluble forms of the receptor retain the ability to bind GM-CSF (6, 7). GM-CSF R α is expressed on hematopoietic stem cells, progenitor and differentiated cells in the myeloid lineage, vascular endothelial cells, placenta, and non-hematopoietic solid tumor cells (8). GM-CSF R α associates with the common beta chain/CD131 (β_c), a 135 kDa transmembrane protein that is also the signal transducing component of the receptors for IL-3 and IL-5 (9, 10). Association with β_c converts GM-CSF R α from a low affinity to a high affinity receptor for GM-CSF (9–11). The shared usage of β_c underlies the synergism between GM-CSF, IL-3, and IL-5 in their effects on myeloid cell differentiation and activation (1, 2).

References:

- 1. Martinez-Moczygemba, M. and D.P. Huston (2003) J. Allergy Clin. Immunol. 112:653.
- 2. Fleetwood, A.J. et al. (2005) Crit. Rev. Immunol. 25:405.
- Eksioglu, E.A. et al. (2007) Exp. Hematol. 35:1163.
- Cao, Y. (2007) J. Clin. Invest. 117:2362.
- 5. Gearing, D.P. et al. (1989) EMBO J. 8:3667
- Pelley, J.L. et al. (2007) Exp. Hematol. 35:1483.
- 7. Raines, M.A. et al. (1991) Proc. Natl. Acad. Sci. 88:8203.
- 8. Chiba, S. et al. (1990) Cell Regul. 1:327.
- 9. Kitamura, T. et al. (1991) Proc. Natl. Acad. Sci. 88:5082.
- 10. Hayashida, K. et al. (1990) Proc. Natl. Acad. Sci. 87:9655.
- 11. Hoang, T. et al. (1993) J. Biol. Chem. 268:11881.

