

DESCRIPTION

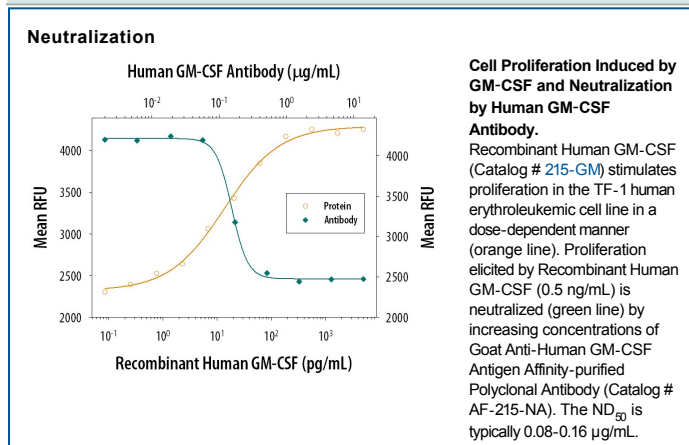
Species Reactivity	Human
Specificity	Detects human GM-CSF in direct ELISAs and Western blots. In direct ELISAs, less than 1% cross-reactivity with recombinant mouse GM-CSF is observed. Neutralizes the biological activity of both recombinant human GM-CSF and natural human GM-CSF.
Source	Polyclonal Goat IgG
Purification	Antigen Affinity-purified
Immunogen	<i>E. coli</i> -derived recombinant human GM-CSF Ala18-Glu144 Accession # P04141
Endotoxin Level	<0.10 EU per 1 µg of the antibody by the LAL method.
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details.

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
Western Blot	0.1 µg/mL	Recombinant Human GM-CSF (Catalog # 215-GM)
Neutralization	Measured by its ability to neutralize GM-CSF-induced proliferation in the TF-1 human erythroleukemic cell line. Kitamura, T. <i>et al.</i> (1989) <i>J. Cell Physiol.</i> 140 :323. The Neutralization Dose (ND ₅₀) is typically 0.08-0.16 µg/mL in the presence of 0.5 ng/mL Recombinant Human GM-CSF.	

DATA



PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.2 mg/mL in sterile PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. <ul style="list-style-type: none"> • 12 months from date of receipt, -20 to -70 °C as supplied. • 1 month, 2 to 8 °C under sterile conditions after reconstitution. • 6 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

GM-CSF was initially characterized as a factor that can support the *in vitro* colony formation of granulocyte-macrophage progenitors. It is also a growth factor for erythroid, megakaryocyte, and eosinophil progenitors. GM-CSF is produced by a number of different cell types (including T cells, B cells, macrophages, mast cells, endothelial cells, fibroblasts, and adipocytes) in response to cytokine or inflammatory stimuli. On mature hematopoietic cells, GM-CSF is a survival factor for and activates the effector functions of granulocytes, monocytes/macrophages, and eosinophils (1, 2). GM-CSF promotes a Th1 biased immune response, angiogenesis, allergic inflammation, and the development of autoimmunity (3-5). It shows clinical effectiveness in ameliorating chemotherapy-induced neutropenia, and GM-CSF transfected tumor cells are utilized as cancer vaccines (6, 7). The 22 kDa glycosylated GM-CSF, similar to IL-3 and IL-5, is a cytokine with a core of four bundled α -helices (8-12). Mature human GM-CSF shares 63%-70% amino acid sequence identity with canine, feline, porcine, and rat GM-CSF and 54% with mouse GM-CSF. GM-CSF exerts its biological effects through a heterodimeric receptor complex composed of GM-CSF R α /CD116 and the signal transducing common β chain (CD131) which is also a component of the high-affinity receptors for IL-3 and IL-5 (13, 14). In addition, GM-CSF binds a naturally occurring soluble form of GM-CSF R α (15). Human GM-CSF is active on canine and feline cells but not on murine cells (16-18).

References:

1. Martinez-Moczygemba, M. and D.P. Huston (2003) J. Allergy Clin. Immunol. **112**:653.
2. Barreda, D.R. *et al.* (2004) Dev. Comp. Immunol. **28**:509.
3. Eksioglu, E.A. *et al.* (2007) Exp. Hematol. **35**:1163.
4. Cao, Y. (2007) J. Clin. Invest. **117**:2362.
5. Fleetwood, A.J. *et al.* (2005) Crit. Rev. Immunol. **25**:405.
6. Heuser, M. *et al.* (2007) Semin. Hematol. **44**:148.
7. Hege, K.M. *et al.* (2006) Int. Rev. Immunol. **25**:321.
8. Kaushansky, K. *et al.* (1992) Biochemistry **31**:1881.
9. Diederichs, K. *et al.* (1991) Science **254**:1779.
10. Cantrell, M.A. *et al.* (1985) Proc. Natl. Acad. Sci. **82**:6250.
11. Lee, F. *et al.* (1985) Proc. Natl. Acad. Sci. **82**:4360.
12. Wong, G.G. *et al.* (1985) Science **228**:810.
13. Onetto-Pothier, N. *et al.* (1990) Blood **75**:59.
14. Hayashida, K. *et al.* (1990) Proc. Natl. Acad. Sci. **87**:9655.
15. Pelley, J.L. *et al.* (2007) Exp. Hematol. **35**:1483.
16. Hogge, G.S. *et al.* (1990) Cancer Gene Ther. **6**:26.
17. Sprague, W.S. *et al.* (2005) J. Comp. Pathol. **133**:136.
18. Shanafelt, A.B. *et al.* (1991) J. Biol. Chem. **266**:13804.