

DESCRIPTION

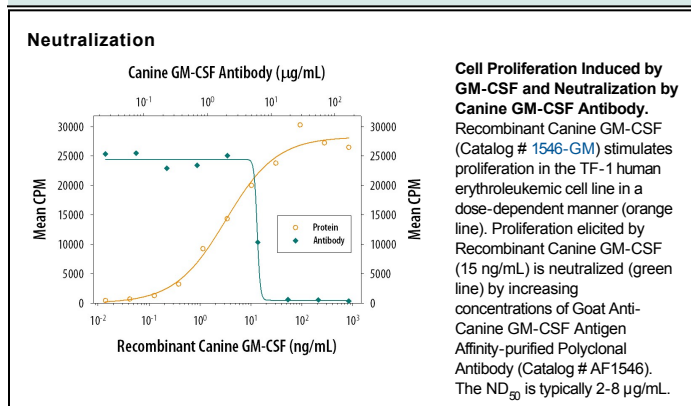
Species Reactivity	Canine
Specificity	Detects canine GM-CSF in direct ELISAs and Western blots. In Western blots, approximately 25% cross-reactivity with recombinant rat GM-CSF is observed, approximately 5% cross-reactivity with recombinant feline GM-CSF and recombinant porcine GM-CSF is observed, and less than 1% cross-reactivity with recombinant human GM-CSF and recombinant mouse GM-CSF is observed.
Source	Polyclonal Goat IgG
Purification	Antigen Affinity-purified
Immunogen	<i>E. coli</i> -derived recombinant canine GM-CSF Ala18-Lys144 Accession # P48749.1
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. *Small pack size (-SP) is supplied as a 0.2 µm filtered solution in PBS.

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 Canine GM-CSF (Catalog # 1546-GM)
Immunocytochemistry	5-15 µg/mL	Immersion fixed canine lymphocytes
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 2-8 µg/mL in the presence of 15 ng/mL Recombinant Canine 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. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C
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 (8-10). Mature canine GM-CSF shares 49-57% amino acid sequence identity with mouse and rat GM-CSF and 69-72% with feline, human, and porcine 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 (11, 12). In addition, GM-CSF binds a naturally occurring soluble form of GM-CSF R α (13). The activity of GM-CSF is species specific between human and mouse, although human GM-CSF is active on canine cells (14, 15).

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. Nash, R.A. *et al.* (1991) *Blood* **78**:930.
11. Onetto-Pothier, N. *et al.* (1990) *Blood* **75**:59.
12. Hayashida, K. *et al.* (1990) *Proc. Natl. Acad. Sci. USA* **87**:9655.
13. Pelley, J.L. *et al.* (2007) *Exp. Hematol.* **35**:1483.
14. Shanafelt, A.B. *et al.* (1991) *J. Biol. Chem.* **266**:13804.
15. Hogge, G.S. *et al.* (1990) *Cancer Gene Ther.* **6**:26.