

Porcine IL-12 Antibody

Antigen Affinity-purified Polyclonal Goat IgG Catalog Number: AF912

DESCRIPTION			
Species Reactivity	Porcine		
Specificity	Detects porcine IL-12 in direct ELISAs and Western blots. In direct ELISAs, less than 5% cross-reactivity with recombinant mouse IL-12 is observed.		
Source	Polyclonal Goat IgG		
Purification	Antigen Affinity-purified		
Immunogen	Chinese hamster ovary cell line CHO-derived recombinant porcine IL-12 p35/p40 heterodimer Ile23-Asn324 (p40), Arg26-Ser222 (p35) Accession # Q28938 (p40), Q29053 (p35)		
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 either lyophilized or 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 Porcine IL-12 (Catalog # 912-PL)
Neutralization	Measured by its ability to neutralize IL-12-induced proliferation in PHA-activated human peripheral blood mononuclear cells (PBMC) [Yokota, T. <i>et al.</i> (1986) Proc. Natl. Acad. Sci. USA 83 :5894]. The Neutralization Dose (ND ₅₀) is typically 0.01-0.04 μg/mL in the presence of 0.75 ng/mL Recombinant Porcine IL-12.	

DATA

Neutralization Porcine IL-12 Antibody (ng/mL) 5000 5000 4000 Mean CPM 3000 3000 2000 2000 1000 Recombinant Porcine IL-12 (ng/mL)

Cell Proliferation Induced by IL-12 and Neutralization by Porcine IL-12 Antibody. Recombinant Porcine IL-12 (Catalog # 912-PL) stimulates

proliferation in PHA-activated human peripheral blood mononuclear cells (PBMC) in a dose-dependent manner (orange line). Proliferation elicited by Recombinant Porcine IL-12 (0.75 ng/mL) is neutralized (green line) by increasing concentrations of Goat Anti-Porcine IL-12 Antigen Affinitypurified Polyclonal Antibody (Catalog # AF912). The ND₅₀ is typically 0.01-0.04 µg/mL.

Reconstitution	Reconstitute at 0.2 mg

PREPARATION AND STORAGE

g/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.

- 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.

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BACKGROUND

Interleukin 12, also known as natural killer cell stimulatory factor (NKSF) or cytotoxic lymphocyte maturation factor (CLMF), is a heterodimeric pleiotropic cytokine made up of a 40 kDa (p40) subunit and a 35 kDa (p35) subunit. IL-12 is produced by macrophages and B lymphocytes and has been shown to have multiple effects on T cells and natural killer (NK) cells. Some of these IL-12 activities include the induction of IFN-y and TNF in resting and activated T and NK cells, the enhancement of cytotoxic activity of resting NK and T cells, the stimulation of resting T cell proliferation in the presence of a comitogen, and the enhancement of NK cell proliferation. Current evidence indicates that IL-12 is a key mediator of cellular-immunity and induces the differentiation of Th1 cells from precursor T helper cells. Based on its activities, it has been suggested that IL-12 may have therapeutic potential as a vaccine adjuvant that promotes cellular-immunity and as an anti-tumor and anti-viral agent.

Porcine IL-12 subunits p35 and p40 share about 85% homology to the human subunits, but differ by containing a 3 amino acid addition in the p35 subunit and a 4 amino acid deletion in the p40 subunit. Porcine IL-12 induces proliferation of human lymphoblasts and IFN-γ secretion by human and porcine lymph nodes. Porcine IL-12 has been detected in lymphoid tissues including inguinal and mesenteric lymph nodes, Peyer's patches, spleen, and thymus.

References:

1. Foss, D. et al. (1997) Vet. Immunol. Immunopathol. 57:121.

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