

Recombinant Cynomolgus Monkey IL-12 (linked heterodimer)

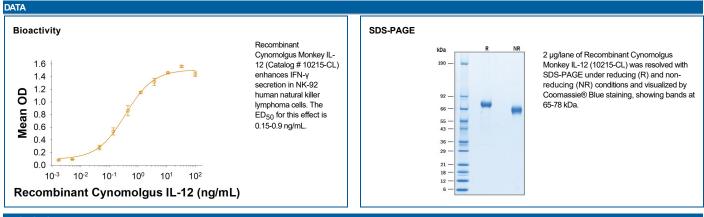
Catalog Number: 10215-CL

DESCRIPTION			
Source	Human embryonic kidney cell, HEK293-derived cynomolgus monkey IL-12 protein		
	Cynomolgus Monkey IL-12p40 (Ile23-Ser328) Accession # NP_001274204	GGGSGGGSGGGS	Cynomolgus Monkey IL-12p35 (Arg57-Ser253) Accession # XP_005546300
	N-terminus		C-terminus

N-terminal Sequence Analysis	N-terminal Sequence IIe23 Analysis		
Structure / Form	Disulfide-linked heterodimer		
Predicted Molecular	58 kDa		

SPECIFICATIONS		
SDS-PAGE	65-78 kDa, under reducing conditions	
Activity	Measured by its ability to enhance IFN-γ secretion in NK-92 human natural killer lymphoma cells. The ED ₅₀ for this effect is 0.15-0.9 ng/mL.	
Endotoxin Level	<0.10 EU per 1 µg of the protein by the LAL method.	
Purity	>95%, by SDS-PAGE visualized with Silver Staining and quantitative densitometry by Coomassie® Blue Staining.	
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS. See Certificate of Analysis for details.	

PREPARATION AND STORAGE			
Reconstitution	Reconstitute at 100 μg/mL in 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. 12 months from date of receipt, -20 to -70 °C as supplied. 1 month, 2 to 8 °C under sterile conditions after reconstitution. 3 months, -20 to -70 °C under sterile conditions after reconstitution.		



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 (1, 2). The cynomolgus monkey p35 subunit shares 93.4% sequence identity with the human homolog, and the p40 subunit shares 96.4% identity with its human homolog. 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-gamma 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 (2, 3). 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 (1-3).

References:

- 1. Hasegawa, H. et al. (2016) Front. Immunol. 7:479.
- 2. Schurich, A. et al. (2017) Rheumatology. 57:246.
- 3. Agarwal, P. (2009). J Immunol. 183:1695.

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