

**DESCRIPTION**

<b>Source</b>	<i>Spodoptera frugiperda</i> , Sf 21 (baculovirus)-derived	
	Human IL-12 p40 (Ile23-Ser328) Accession # P29460	
	Human IL-12 p35 (Arg23-Ser219) Accession # P29459	
	N-terminus	C-terminus

**N-terminal Sequence Analysis** Ile23 (p40) & Arg23 (p35)

**Structure / Form** Disulfide-linked heterodimer

**Predicted Molecular Mass** 34.7 kDa (p40) & 22.5 kDa (p35)

**SPECIFICATIONS**

**SDS-PAGE** 41 kDa (p40) & 29 kDa (p35), reducing conditions

**Activity** Measured in a cell proliferation assay using PHA-stimulated human T lymphoblasts. Symons, J.A. *et al.* (1987) in *Lymphokines and Interferons, a Practical Approach*. Clemens, M.J. *et al.* (eds): IRL Press. 272.  
The ED<sub>50</sub> for this effect is typically 0.01-0.05 ng/mL.  
The specific activity of Recombinant Human IL-12 is approximately 1.1 x 10<sup>4</sup> units/μg, which is calibrated against recombinant human IL-12 WHO Standard (NIBSC code: 95/544).

**Endotoxin Level** <1.0 EU per 1 μg of the protein by the LAL method.

**Purity** >97%, 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**

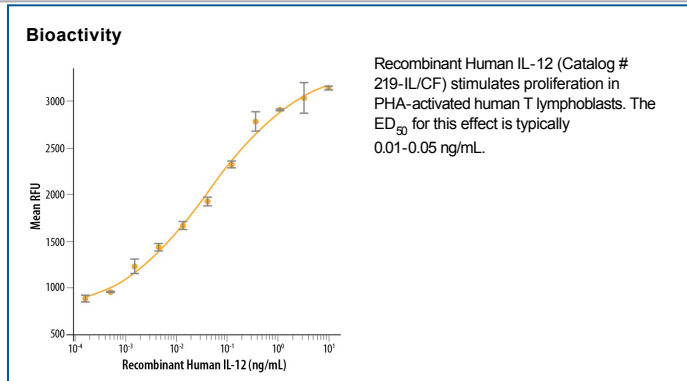
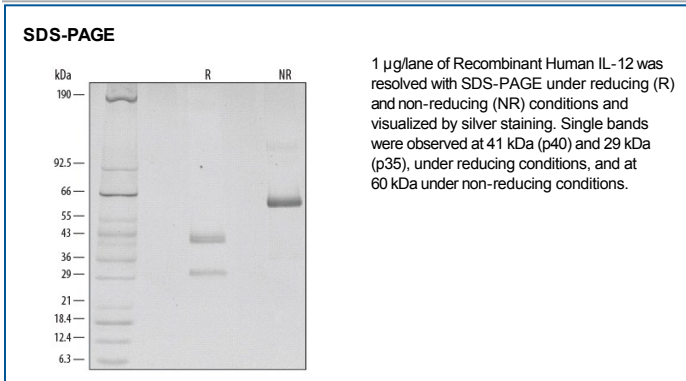
**Reconstitution** Reconstitute 5 μg vials at 50 μg/mL in sterile PBS. Reconstitute 25 μg or larger vials at 100 μg/mL in sterile PBS.

**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.

- 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.

**DATA**



#### BACKGROUND

Interleukin 12, also known as natural killer cell stimulatory factor (NKSF) or cytotoxic lymphocyte maturation factor (CLMF), is a pleiotropic cytokine originally identified in the medium of activated human B lymphoblastoid cell lines. The p40 subunit of IL-12 has been shown to have extensive amino acid sequence homology to the extracellular domain of the human IL-6 receptor while the p35 subunit shows distant but significant sequence similarity to IL-6, G-CSF, and chicken MGF. These observations have led to the suggestion that IL-12 might have evolved from a cytokine/soluble receptor complex. Human and murine IL-12 share 70% and 60% amino acid sequence homology in their p40 and p35 subunits, respectively. IL-12 apparently shows species specificity with human IL-12 reportedly showing minimal activity in the murine system.

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. These effects include inducing production of IFN- $\gamma$  and TNF by resting and activated T and NK cells, synergizing with other IFN- $\gamma$  inducers at both the transcriptional and post-transcriptional levels. This interaction induces IFN- $\gamma$  gene expression, enhancing the cytotoxic activity of resting NK and T cells, inducing and synergizing with IL-2 in the generation of lymphokine-activated killer (LAK) cells, acting as a co-mitogen to stimulate proliferation of resting T cells, and inducing proliferation of activated T and NK cells. Current evidence indicates that IL-12, produced by macrophages in response to infectious agents, is a central mediator of the cell-mediated immune response by its actions on the development, proliferation, and activities of TH1 cells. In its role as the initiator of cell-mediated immunity, it has been suggested that IL-12 has therapeutic potential as a stimulator of cell-mediated immune responses to microbial pathogens, metastatic cancers, and viral infections such as AIDS.