

DESCRIPTION

Source *E. coli*-derived
Glu23-Cys156, with an N-terminal Met
Accession # P01581

N-terminal Sequence Analysis Met

Predicted Molecular Mass 15.5 kDa

SPECIFICATIONS

Activity Measured in an anti-viral assay using L-929 mouse fibroblast cells infected with encephalomyocarditis (EMC) virus. Meager, A. (1987) in *Lymphokines and Interferons, a Practical Approach*. Clemens, M.J. *et al.* (eds): IRL Press. 129.
The ED₅₀ for this effect is 0.1-0.6 ng/mL.

Endotoxin Level <0.01 EU per 1 μ g of the protein by the LAL method.

Purity >97%, by SDS-PAGE under reducing conditions and visualized by silver stain.

Formulation Lyophilized from a 0.2 μ m filtered solution in Tris, NaCl and PEG with BSA as a carrier protein. See Certificate of Analysis for details.

PREPARATION AND STORAGE

Reconstitution Reconstitute at 100 μ g/mL in sterile PBS containing at least 0.1% human or bovine serum albumin.

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

Interferon-gamma (IFN- γ), also known as type II or immune interferon, exerts a wide range of immunoregulatory activities and is considered to be the prototype proinflammatory cytokine (1, 2). Mature rat IFN- γ exists as a noncovalently linked homodimer of 20 - 25 kDa variably glycosylated subunits (3). It shares 86% amino acid sequence identity with mouse IFN- γ and 37%-45% with bovine, canine, cotton rat, equine, feline, human, porcine, and rhesus IFN- γ . IFN- γ dimers bind to IFN- γ RI (alpha subunits) which then interact with IFN- γ RII (beta subunits) to form the functional receptor complex of two α and two β subunits. Inclusion of IFN- γ RII increases the binding affinity for ligand and the efficiency of signal transduction (4, 5). IFN- γ is produced by a variety of immune cells under inflammatory conditions, notably by T cells and NK cells (6). It plays a key role in host defense by promoting the development and activation of Th1 cells, chemoattraction and activation of monocytes and macrophages, upregulation of antigen presentation molecules, and immunoglobulin class switching in B cells. It also exhibits antiviral, antiproliferative, and apoptotic effects (6, 7). In addition, IFN- γ functions as an anti-inflammatory mediator by promoting the development of regulatory T cells and inhibiting Th17 cell differentiation (8, 9). The pleiotropic effects of IFN- γ contribute to the development of multiple aspects of atherosclerosis (7).

References:

1. Billiau, A. and P. Matthys (2009) *Cytokine Growth Factor Rev.* **20**:97.
2. Pestka, S. *et al.* (2004) *Immunol. Rev.* **202**:8.
3. Dijkema, R. *et al.* (1985) *EMBO J.* **4**:761.
4. Marsters, S.A. *et al.* (1995) *Proc. Natl. Acad. Sci.* **92**:5401.
5. Krause, C.D. *et al.* (2000) *J. Biol. Chem.* **275**:22995.
6. Schroder, K. *et al.* (2004) *J. Leukoc. Biol.* **75**:163.
7. McLaren, J.E. and D.P. Ramji (2009) *Cytokine Growth Factor Rev.* **20**:125.
8. Muhl, H. and J. Pfeilschifter (2003) *Int. Immunopharmacol.* **3**:1247.
9. Kelchtermans, H. *et al.* (2008) *Trends Immunol.* **29**:479.