

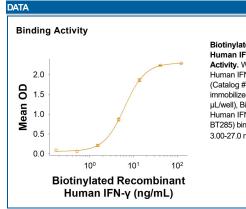
Biotinylated Recombinant Human IFN-y

Catalog Number: BT285

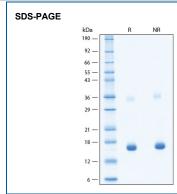
DESCRIPTION	
Source	E. coli-derived human IFN-gamma protein Gln24-Gln166, with a N-terminal Met Accession # CAA31639.1
N-terminal Sequence Analysis	Met
Structure / Form	Biotinylated via Amines
Predicted Molecular	16.9 kDa

SPECIFICATIONS	
SDS-PAGE	14-20 kDa, under reducing conditions.
Activity	Measured by its binding ability in a functional ELISA. When Recombinant Human IFN-γ R1/CD119 (Catalog # 673-IR/CF) is immobilized at 3 μg/mL (100 μL/well), Biotinylated Recombinant Human IFN-γ (Catalog # BT285) binds with an ED ₅₀ of 3.00-27.0 ng/mL.
Endotoxin Level	<0.10 EU per 1 µg of the protein by the LAL method.
Purity	>90%, by SDS-PAGE visualized with Silver Staining and quantitative densitometry by Coomassie® Blue Staining.
Formulation	Supplied as a 0.2 µm filtered solution in Sodium Succinate, Mannitol and Tween®® 80. See Certificate of Analysis for details.

PREPARATION AND STORAGE	
Reconstitution	Reconstitute at 0.2 mg/mL in sterile, deionized water.
Shipping	The product is shipped with dry ice or equivalent. Upon receipt, store it immediately at the temperature recommended below.
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. • 6 months from date of receipt, -70 °C as supplied. • 1 month, 2 to 8 °C under sterile conditions after opening. • 3 months, -20 to -70 °C under sterile conditions after opening.



Biotinylated Recombinant Human IFN-γ Protein Binding Activity. When Recombinant Human IFN-γ R1/CD119 (Catalog # 673-IR/CF) is immobilized at 3 μg/mL (100 μL/well), Biotinylated Recombinant Human IFN-γ Protein (Catalog # BT285) binds with an ED₅₀ of 3.00-27.0 ng/mL.



Biotinylated Recombinant Human IFN-γ Protein SDS-PAGE. 2 μg/lane of Recombinant Human IFN-gamma Biotinylated Protein (Catalog # BT285) was resolved with SDS-PAGE under reducing (R) and non-reducing (NR) conditions and visualized by Coomassie® Blue staining, showing bands at 14-20 kDa.

Rev. 3/7/2023 Page 1 of 2





Biotinylated Recombinant Human IFN-γ

Catalog Number: BT285

BACKGROUND

Interferon-gamma (IFN-gamma), 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 human IFN-gamma exists as a non-covalently linked homodimer of 20-25 kDa variably glycosylated subunits (3). It shares 90% amino acid (aa) sequence identity with rhesus IFN-gamma, 59%-64% with bovine, canine, equine, feline, and porcine IFN-gamma, and 37%-43% with cotton rat, mouse, and rat IFN-gamma. IFN-gamma dimers bind to IFN-gamma RI (alpha subunits) which then interact with IFN-gamma RII (beta subunits) to form the functional receptor complex of two alpha and two beta subunits. Inclusion of IFN-gamma RII increases the binding affinity for ligand and the efficiency of signal transduction (4, 5). IFN-gamma 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, up-regulation of antigen presentation molecules, and immunoglobulin class switching in B cells. It also exhibits antiviral, antiproliferative, and apoptotic effects (6, 7). In addition, IFN-gamma 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-gamma 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. Gray, P.W. and D.V. Goeddel (1982) Nature 298:859.
- 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.