

DESCRIPTION

Source *E. coli*-derived human IL-21 protein
Gln32-Ser162, with a N-terminal Met
Accession # Q9HBE4.3
Produced in an animal-free laboratory. Manufactured and tested under cGMP guidelines

N-terminal Sequence Analysis Met-Gln-Asp-Arg-His-Met-Ile-Arg-Met-Arg

Predicted Molecular Mass 15 kDa

SPECIFICATIONS

SDS-PAGE 16 kDa, reducing conditions

Activity Measured in a cell proliferation assay using B9 mouse hybridoma cells.
The ED₅₀ for this effect is 5.00-50.0 ng/mL.
The specific activity of recombinant human IL-21 is >1.00 x 10⁶ units/mg, which is calibrated against an internal reference standard for human IL-21.

Measured by its ability to enhance IFN-γ secretion in NK-92 human natural killer lymphoma cells.
The ED₅₀ for this effect is ≤8 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.

Host Cell Protein < 0.1 ng per µg of protein when tested by ELISA.

Mycoplasma Negative for Mycoplasma.

Formulation Lyophilized from a 0.2 µm filtered solution in PBS. See Certificate of Analysis for details.

PREPARATION AND STORAGE

Reconstitution Reconstitute at 100-200 µg/mL in PBS.

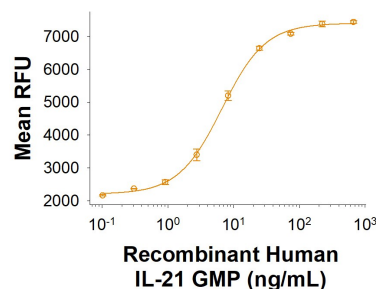
Shipping The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.

Stability & Storage **Use a manual defrost freezer and avoid repeated freeze-thaw cycles.**

- A minimum of 12 months when stored at ≤ -20 °C as supplied. Refer to lot specific COA for the Use by Date.
- 1 month, 2 to 8 °C under sterile conditions after reconstitution.
- 3 months, ≤ -20 °C under sterile conditions after reconstitution.

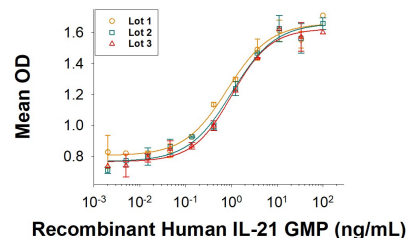
DATA

Bioactivity



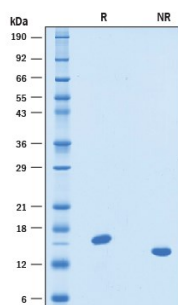
Recombinant Human IL-21 GMP Protein Bioactivity. Recombinant Human IL-21 GMP (Catalog # 8879-GMP) induces B9 mouse hybridoma cell proliferation. The ED₅₀ for this effect is 5.00-50.0 ng/mL.

Bioactivity



Recombinant Human IL-21 GMP Protein Bioactivity GMP-grade Recombinant Human IL-21 (Catalog # 8879-GMP) enhances IFN-gamma secretion in NK-92 human natural killer lymphoma cells. The ED₅₀ for this effect is ≤8 ng/mL. Three independent lots were tested for activity and plotted on the same graph to show lot-to-lot consistency of GMP IL-21.

SDS-PAGE



Recombinant Human IL-21 GMP Protein SDS-PAGE 2 µg/lane of GMP-grade Recombinant Human IL-21 (Catalog # 8879-GMP) was resolved with SDS-PAGE under reducing (R) and non-reducing (NR) conditions and visualized by Coomassie® Blue staining, showing R band at 16.4 kDa and NR band at 14.8 kDa, respectively.

BACKGROUND

Interleukin-21 (IL-21) is an approximately 14 kDa four-helix-bundle member of the family of cytokines that utilize the common gamma chain (γ_c) as a receptor subunit. γ_c is also a subunit of the receptors for IL-2, IL-4, IL-7, IL-9, and IL-15 (1). IL-21 is produced by activated T follicular helper cells (Tfh), Th17 cells, and NKT cells (2-6). It exerts its biological effects through a heterodimeric receptor complex of γ_c and the IL-21-specific IL-21 R (2, 7). Tfh-derived IL-21 plays an important role in the development of humoral immunity through its autocrine effects on the Tfh cell and paracrine effects on immunoglobulin affinity maturation, plasma cell differentiation, and B cell memory responses (4, 8, 9). It is also required for the migration of dendritic cells to draining lymph nodes (10). IL-21 regulates several aspects of T cell function. It co-stimulates the activation, proliferation, and survival of CD8⁺ T cells and NKT cells and promotes Th17 cell polarization (3, 5, 6, 11, 12). It blocks the generation of regulatory T cells and their suppressive effects on CD4⁺ T cells (13, 14). IL-21 R engagement enhances the cytolytic activity and IFN- γ production of activated NK cells but limits the expansion of resting NK cells (15). In addition, IL-21 suppresses cutaneous hypersensitivity reactions by limiting allergen-specific IgE production and mast cell degranulation (16). Dysregulation of the IL-21/IL-21 R system contributes to the development of multiple immunological disorders (1, 17). The 133 amino acid (aa) mature human IL-21 shares 63% and 61% aa sequence identity with mouse and rat IL-21, respectively. Alternative splicing generates an additional isoform with a substitution of the C-terminal 16 amino acids (18).

References:

1. Tangye, S.G. (2015) *Curr. Opin. Immunol.* **34**:107.
2. Parrish-Novak, *et al.* (2000) *Nature* **408**:57.
3. Coquet, J.M. *et al.* (2007) *J. Immunol.* **178**:2827.
4. Vogelzang, A. *et al.* (2008) *Immunity* **29**:127.
5. Korn, T. *et al.* (2007) *Nature* **448**:484.
6. Nurieva, R. *et al.* (2007) *Nature* **448**:480.
7. Asao, H. *et al.* (2001) *J. Immunol.* **167**:1.
8. Zotos, D. *et al.* (2010) *J. Exp. Med.* **207**:365.
9. Rankin, A.L. *et al.* (2011) *J. Immunol.* **186**:667.
10. Jin, H. *et al.* (2009) *J. Clin. Invest.* **119**:47.
11. Frohlich, A. *et al.* (2009) *Science* **324**:1576.
12. Yi, J.S., *et al.* (2009) *Science* **324**:1572.
13. Peluso, I. *et al.* (2007) *J. Immunol.* **178**:732.
14. Bucher, C. *et al.* (2009) *Blood* **114**:5375.
15. Kasaian, M.T. *et al.* (2002) *Immunity* **16**:559.
16. Tamagawa-Mineoka, R. *et al.* (2011) *J. Invest. Dermatol.* **131**:1513.
17. Ma, J. *et al.* (2011) *Cytokine* **56**:133.
18. Rahman, M. *et al.* (2007) *FEBS Lett.* **581**:4001.

MANUFACTURING SPECIFICATIONS

GMP Proteins

R&D Systems, a Bio-Techne Brand's GMP proteins are produced according to relevant sections of the following documents: USP Chapter 1043, Ancillary Materials for Cell, Gene and Tissue-Engineered Products and Eu. Ph. 5.2.12, Raw Materials of Biological Origin for the Production of Cell-based and Gene Therapy Medicinal Products.

R&D Systems' quality focus includes:

- Manufactured and tested under an ISO 9001:2015 and ISO 13485:2016 certified quality system
- Documented processes and QA control of documentation and process changes
- Personnel training programs
- Raw material testing and vendor qualification/monitoring
- Fully validated equipment, processes and test methods
- Equipment calibration schedules using a computerized calibration program
- Facility maintenance, safety programs and pest control
- Material review process for variances
- Monitoring of stability over product shelf-life

R&D Systems strives to provide our customers with the analytical characteristics of each product so that customers may determine whether our products are appropriate for their research. The Certificate of Analysis provided contains the following lot specific information:

- N-terminal amino acid analysis, SDS-PAGE analysis, and endotoxin level (as determined by LAL assay) performed on each bulk QC lot, not on individual bottlings of each QC lot
- Post-bottling lot-specific bioassay results (compliance with an established range) and results of microbial testing according to USP
- Host Cell Protein testing performed by ELISA
- Mycoplasma testing by ribosomal RNA hybridization assay

Additional testing and documentation requested by the customer can be arranged at an additional cost.

Production records and facilities are available for examination by appropriate personnel on-site at R&D Systems in Minneapolis, Minnesota USA.

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Production

- All molecular biology procedures use animal-free media and dedicated labware.
- Dedicated fermentors are utilized in committed animal-free areas.

Purification

- Protein purification columns are animal-free.
- Bulk proteins are filtered using animal-free filters.
- Purified proteins are stored in animal-free containers in a dedicated cold storage room.

Quality Assurance

- Low Endotoxin Level.
- No impairment of biological activity.
- High quality product obtained under stringent conditions.

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