

Catalog Number: PPK-002-GMPIU

		RI		

Source E. coli-derived human IL-2 protein

Ala21-Thr153 (Cys145Ser), with and without an N-terminal Met

Accession # P60568.1

Produced using non-animal reagents in an animal-free laboratory.

Manufactured and tested under cGMP guidlelines.

N-terminal Sequence Met-Ala21-Pro-Thr-Ser-Ser-Thr-Lys-Lys & Ala21-Pro-Thr-Ser-Ser-Thr-Lys-Lys-Thr

Analysis

Predicted Molecular 15.5 kDa

Mass

SPECIFICATIONS					
SDS-PAGE	13 kDa, under reducing conditions.				
Activity	Measured in a cell proliferation assay using CTLL-2 mouse cytotoxic T cells. Gearing, A.J.H. and C.B. Bird (1987) in Lymphokines and Interferons, A Practical Approach. Clemens, M.J. <i>et al.</i> (eds): IRL Press. 295. The potency of recombinant human IL-2 is 75-125% of the IU pack size.				
	International units are calibrated from an internal reference standard that is value assigned against the human IL-2 WHO International Standard (NIBSC code: 86/500).				
Endotoxin Level	<5.0 EU/mL by the LAL method.				
Purity	>97%, by SDS-PAGE with quantitative densitometry by Coomassie® Blue Staining.				
Mass Spectrometry	The molecular weight by mass spectrometry is 15521 ± 5 Da, and a second 15390 ± 5 Da product may be present.				
Host Cell Protein	<0.500 ng per μg of protein when tested by ELISA.				
Mycoplasma	Negative for Mycoplasma.				
Host Cell DNA	t Cell DNA <0.00150 ng per μg of protein when tested by PCR.				

Supplied as a 0.2 µm filtered solution in PBS, recombinant HSA, and Trehalose. See Certificate of Analysis for details.

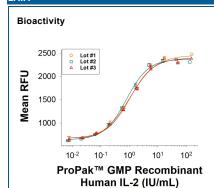
PREPARATION AND STORAGE

Shipping This product is shipped on dry ice. Upon receipt, store immediately at the temperature recommended below.

A minimum of 6 months when stored between -14 °C and -40 °C. Can be stored up to 2 weeks at 2-8 °C. Refer to lot specific COA for the Use by Date.

DATA

Formulation



Lot to Lot Consistency
ProPak™ GMP-grade
Recombinant Human IL-2
International Unit Based Filling
(Catalog # PPK-002-GMPIU)
stimulates proliferation of CTLL-2
mouse cytotoxic T cells. Three
independent lots were tested for
activity and plotted on the same
graph to show lot-to-lot
consistency of ProPak™ GMP IL2.

Rev. 6/5/2025 Page 1 of 4



Catalog Number: PPK-002-GMPIU

BACKGROUND

Recombinant Interleukin-2 (IL-2) is expressed in E. coli and has been engineered to contain the serine for cysteine substitution found in Proleukin® (aldesleukin). Recombinant IL-2 is widely used in cell culture for the expansion of T cells. IL-2 is expressed by CD4+ and CD8+ T cells, γδ T cells, B cells, dendritic cells, and eosinophils (1-3). Mature human IL-2 shares 56% and 66% amino acid (aa) sequence identity with mouse and rat IL-2, respectively. Human and mouse IL-2 exhibit cross-species activity (4). The receptor for IL-2 consists of three subunits that are present on the cell surface in varying preformed complexes (5-7). The 55 kDa IL-2 Rα is specific for IL-2 and binds with low affinity. The 75 kDa IL-2 Rβ, which is also a component of the IL-15 receptor, binds IL-2 with intermediate affinity. The 64 kDa common gamma chain yc/lL-2 Ry, which is shared with the receptors for IL-4, -7, -9, -15, and -21, does not independently interact with IL-2. Upon ligand binding, signal transduction is performed by both IL-2 Rβ and γc. IL-2 is best known for its autocrine and paracrine activity on T cells. It drives resting T cells to proliferate and induces IL-2 and IL-2 Rα synthesis (1, 2). It contributes to T cell homeostasis by promoting the Fas-induced death of naïve CD4+ T cells but not activated CD4+ memory lymphocytes (8). IL-2 plays a central role in the expansion and maintenance of regulatory T cells, although it inhibits the development of Th17 polarized cells (9-11). Thus, IL-2 may be a key cytokine in the natural suppression of autoimmunity (12, 13). IL-2 expression and concentration can have either immunostimulatory effects at high doses or immunosuppressive effects at low doses due to its preferential binding to different receptor subunits expressed by various immune cell types. This has led to the generation of recombinant IL-2 variants aimed at modifying IL-2 receptor binding for increased antitumor efficacy (14, 15). These variants are typically used in combination with immune checkpoint inhibitors instead of as a monotherapy (14). IL-2 can be genetically engineered to express in NK cells for CAR T cell therapies, and in combination with other cytokines like IL-15, can increase cell viability and proliferation (16). In addition to adoptive cell transfer and checkpoint blockade inhibitors, cancer vaccines that boost immune responses have been combined with IL-2 treatment with promising results in recent studies (15). In cell culture, IL-2 is a frequently used cytokine for the proliferation, differentiation, and increased antibody secretion of B cells as they transform into plasma cells in vitro (17). IL-2 is also a classically used cytokine for the expansion of NK cells, early differentiated T cells and effector memory Treg cells for adoptive cell transfer cancer immunotherapy (16, 18). GMP IL-2 is a commonly used supplement for the expansion of these cell types for cellular therapies.

References:

- 1. Ma, A. et al. (2006) Annu. Rev. Immunol. 24:657.
- 2. Gaffen, S.L. and K.D. Liu (2004) Cytokine 28:109.
- 3. Taniguchi, T. et al. (1983) Nature 302:305.
- 4. Mosmann, T.R. et al. (1987) J. Immunol. 138:1813.
- 5. Liparoto, S.F. et al. (2002) Biochemistry 41:2543.
- 6. Wang, X. et al. (2005) Science 310:1159.
- 7. Bodnar, A. et al. (2008) Immunol. Lett. 116:117.
- 8. Jaleco, S. et al. (2003) J. Immunol. 171:61.
- 9. Malek, T.R. (2003) J. Leukoc. Biol. 74:961.
- 10. Laurence, A. et al. (2007) Immunity 26:371.
- 11. Kryczek, I. et al. (2007) J. Immunol. 178:6730.
- 12. Afzali, B. *et al.* (2007) Clin. Exp. Immunol. **148**:32.
- 13. Fehervari, Z. et al. (2006) Trends Immunol. 27:109.
- 14. Xue, D. et al. (2021) Antib Ther. 4:123.
- 15. Wolfarth, A.A. et al. (2022) Immune Netw. 22:e5.
- 16. Koehl, U. et al. (2015) Oncoimmunology. 5:e1115178.
- 17. Marsman, C. et al. (2022) Front. Immunol. 13:815449.
- 18. Chamucero-Millares, J.A.et al. (2021) Cell Immunol. 360:104257



Catalog Number: PPK-002-GMPIU

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, Mass Spectrometry, Host Cell Protein, Host Cell DNA, Mycoplasma testing performed on each bulk QC lot, not on individual finshed product lots
- Finished product testing includes bioassay (compliance with an established range), endotoxin level (as determined by LAL assay), and microbial testing
 according to USP <71>

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.

R&D Systems sells GMP grade products for preclinical or clinical ex vivo cell therapy applications. They are not for in vivo use. Please read the following End User Terms prior to using this product.

Animal-Free Manufacturing Conditions

Our dedicated controlled-access animal-free laboratories ensure that at no point in production are the products exposed to potential contamination by animal components or byproducts. Every stage of manufacturing is conducted in compliance with R&D Systems' stringent Standard Operating Procedures (SOPs). Production and purification procedures use equipment and media that are confirmed animal-free.

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.

China | info.cn@bio-techne.com TEL: 400.821.3475

Quality Assurance

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

Please read our complete Animal-Free Statement.

Rev. 6/5/2025 Page 3 of 4



Catalog Number: PPK-002-GMPIU

PRODUCT SPECIFIC NOTICES

END USER TERMS OF USE OF PRODUCT

The following terms are offered to you upon your acceptance of these End User Terms of Use of Product. By using this product, you indicate your acknowledgment and agreement to these End User Terms of Use of Product. If you do not agree to be bound by and comply with all of the provisions of these End User Terms of Use of Product, you should contact your supplier of the product and make arrangements to return the product.

We suggest you print and retain a copy of these End User Terms of Use of Product for your records.

The End User is aware that R&D Systems, Inc. sells GMP products for preclinical or clinical ex vivo use and not for in vivo use. The End User further agrees, as a condition of the sale of R&D Systems' GMP products that: a) the End User will not use this GMP Product in any procedure wherein the product may be directly or indirectly administered to humans, unless the End User has obtained, or prior to their use will have obtained, an Investigational New Drug (IND) exemption from the FDA and will use the product only in accordance with the protocols of such IND and of the Institutional Review Board overseeing the proposed research, or b) the End User will use the products outside of the United States in accordance with the protocols of research approved by the Institutional Review Board or authorized ethics committee and regulatory agencies to which the End User is subject to in their territory.

R&D Systems, Inc. has the right, at its sole discretion, to modify, add or remove any terms or conditions of these End User Terms of Use without notice or liability to you. Any changes to these End User Terms of Use are effective immediately following the printing of such changes on this product insert. The most recent version of these End User Terms of Use of Product may be found at: RnDSystems.com/Legal.

You agree to review these End User Terms of Use of Product to ensure any subsequent use by you of R&D Systems' GMP Products following changes to these End User Terms of Use of Product constitutes your acceptance of all such changes.

TERMS AND CONDITIONS

The following limitation applies to R&D Systems' warranty and liability for damages: All products are warranted to meet R&D Systems' published specifications when used under normal laboratory conditions.

R&D SYSTEMS DOES NOT MAKE ANY OTHER WARRANTY OR REPRESENTATION WHATSOEVER, WHETHER EXPRESS OR IMPLIED, WITH RESPECT TO ITS PRODUCTS. IN PARTICULAR, R&D SYSTEMS DOES NOT MAKE ANY WARRANTY OF SUITABILITY, NONINFRINGEMENT, MERCHANTABILITY OR FITNESS FOR ANY PARTICULAR PURPOSE.

NOTWITHSTANDING ANY OTHER PROVISIONS OF THESE TERMS AND/OR ANY OTHER AGREEMENT BETWEEN R&D SYSTEMS AND PURCHASER FOR THE PURCHASE OF THE PRODUCTS, R&D SYSTEMS' TOTAL LIABILITY TO PURCHASER ARISING FROM OR IN RELATION TO THESE TERMS, AN AGREEMENT BETWEEN THE PARTIES OR THE PRODUCTS, WHETHER ARISING IN CONTRACT, TORT OR OTHERWISE SHALL BE LIMITED TO THE TOTAL AMOUNT PAID BY PURCHASER TO R&D SYSTEMS FOR THE APPLICABLE PRODUCTS. IN NO EVENT WILL R&D SYSTEMS BE LIABLE FOR THE COST OF PROCUREMENT OF SUBSTITUTE GOODS.

Full details of R&D Systems' Terms and Conditions of Sale can be found online at: RnDSystems.com/Legal.

For preclinical, or clinical ex vivo use Not for in vivo use

Rev. 6/5/2025 Page 4 of 4