

PRODUCT INFORMATION

Croyez GMP® IL-3 (Interleukin-3), Human

v. 230901

Catalog number	C01005-GMP-100 / C01005-GMP-1000
Package	100 μg / 1 mg
Description	Interleukin-3 (IL-3) is an interleukin, a type of biological signal (cytokine) that can improve the body's natural response to disease as part of the immune system. It acts by binding to the IL-3 receptor. IL-3 stimulates the differentiation of multipotent hematopoietic stem cells into myeloid progenitor cells or, with the addition of IL-7, into lymphoid progenitor cells. In addition, IL-3 stimulates proliferation of all cells in the myeloid lineage (granulocytes, monocytes, and dendritic cells), in conjunction with other cytokines, e.g., Erythropoietin (EPO), Granulocyte macrophage colony-stimulating factor (GM-CSF), and IL-6. It is secreted by basophils and activated T cells to support growth and differentiation of T cells from the bone marrow in an immune response.
Expression System	Escherichia coli
Species of Origin	Human
Affinity Tag	His Tag (C-term)
Sequence	Ala20-Phe152
Endotoxin level	<0.05 EU per 1 µg of the protein by the LAL method.
Activity	Measure by its ability to induce TF-1 cells proliferation. The ED $_{50}$ for this effect is <0.15 ng/mL. The specific activity of recombinant human IL-3 is approximately >1.2 x 10^6 IU/mg.
Purity	>98% as determined by SDS-PAGE analysis.
Mycoplasma	Not detected
Form	Lyophilized
Storage Buffer	Lyophilized from a 0.2 µm filtered solution of PBS, pH 8.0.
Reconstitution	It is recommended to reconstitute the lyophilized protein in sterile H_2O to a concentration not less than 0.5 mg/mL and incubate the stock solution for at least 20 min to ensure sufficient re-dissolved .
Stability & Storage	This product is stable after storage at: -20°C for 12 months in lyophilized state from date of receipt.



0°C for 1 month under sterile conditions after reconstitution. d freeze/thaw cycles. recombinant proteins are manufactured in ISO 13485:2016 and facility. The processes include: reagent and laboratory ed and tested under GMP guideline d traceability of raw material the maintenance and equipment calibration
facility. The processes include: reagent and laboratory ed and tested under GMP guideline d traceability of raw material
raining records atch consistency ation of QA control and process changes ed and tested under an ISO 13485:2016 certified quality
nt system onitor of product shelf-life
er G et al. (1990) <i>Biotherapy.</i> 2,4: 337-45. MA. et al. (1998) <i>Stem Cells.</i> 16,5: 301-13. 992) <i>Chem Immunol.</i> 51: 65-106. . et al. (2019) <i>Immunity.</i> 50,4: 796-811.
kDa 75- 60- 45- 35- 25- 17- 11- E analysis of Croyez GMP® IL-3, Human

For Research Use Only. Not for use in diagnostic or therapeutic procedures.