



GMP Recombinant Human GM-CSF (carrier-free)

Catalog# / Size 572914 / 25 μg

572916 / 100 µg

Other Names Colony stimulating factor 2, CSF2, Granulocyte/macrophage colony-stimulating

factor, CSF-α, Pluripoietin-α, Eosinophil colony stimulating factor (Eo-CSF), Burst

promoting activity (BPA).

Description GM-CSF plays a key role in signaling emergency hemopoiesis (predominantly

myelopoiesis) in response to infection, including the production of granulocytes and macrophages in the bone marrow and their maintenance, survival, and functional activation at sites of injury or insult. The receptor for GM-CSF is a heterodimer that comprises a major binding subunit (GMR α) and a major signaling subunit (β c). The receptor subunits are always co-expressed on the surface of leukocytes, with β c being expressed at lower levels than GMR α . Certain non-hemopoietic cell types have also been reported to express the GM-CSF receptor and to respond to GM-CSF stimulation *in vitro*, although the *in vivo*

significance of these observations remains uncertain.

Product Details

Source Human GM-CSF, amino acids Ala18- Glu144 (Accession# NM_000758) was

expressed in E. coli.

Molecular Mass The 127 amino acid N-terminal methionylated recombinant protein has a predicted

molecular mass of 14.5 kD. The DTT-reduced protein migrates at approximately 14 kD and the non-reduced protein migrates with slightly greater mobility by SDS-

PAGE.

Purity >95%, as determined by Coomassie stained SDS-PAGE.

Formulation 0.1 μm filtered protein solution is in PBS, pH 7.2.

Endotoxin Level Less than 0.1 EU per µg protein as determined by the LAL method

Concentration 500 μg/mL

Storage & Handling Unopened vial can be stored between 2°C and 8°C for up to 2 weeks, at -20°C for

up to six months, or at -70°C or colder until the expiration date. For maximum results, quick spin vial prior to opening. The protein can be aliquoted and stored at -20°C or colder. Stock solutions can also be prepared at 50 - 100 $\mu g/mL$ in appropriate sterile buffer, carrier protein such as 0.2 - 1% endotoxin-free BSA or HSA can be added when preparing the stock solution. Aliquots can be stored between 2°C and 8°C for up to one week or stored at -20°C or colder for up to 3

months. Avoid repeated freeze/thaw cycles.

Activity ED₅₀ = 0.01 - 0.04 ng/mL as determined by the dose-dependent stimulation of TF-

1 cell proliferation. Deep Blue Cell Viability™ Kit (Cat. No. 424701) is used to

measure the proliferation.

Application Bioassay

Cell Culture

Application Notes BioLegend carrier-free recombinant proteins provided in liquid format are shipped

on blue ice. Our comparison testing data indicates that when handled and stored as recommended, the liquid format has equal or better stability and shelf-life compared to commercially available lyophilized proteins after reconstitution. Our liquid proteins are validated in-house to maintain activity after shipping on blue ice and are backed by our 100% satisfaction guarantee. If you have any concerns,

contact us at tech@biolegend.com.

Disclaimer GMP Recombinant Proteins. BioLegend GMP recombinant proteins are

manufactured in a dedicated GMP facility and compliant with ISO 13485:2016. For research or *ex vivo* cell processing use. Not for use in diagnostic or therapeutic

procedures. Our processes include:

- · Batch-to-batch consistency
- · Material traceability
- · Documented procedures
- Documented employee training
- Equipment maintenance and monitoring records

- · Lot-specific certificates of analysis
- Quality audits per ISO 13485:2016
- QA review of released products

BioLegend GMP recombinant protiens are manufactured and tested in accordance with USP Chapter 1043, Ancillary Materials for Cell, Gene and Tissue-Engineered Products and Ph. Eur. Chapter 5.2.12.

Antigen Details

Structure Cytokine

Distribution T cells, macrophages, fibroblasts, endothelial cells, mast cells.

Function Synergistic with IL-1, IL-3, G-CSF; E21R competitive antagonist for receptor

binding; stored in ECM with heparan sulfate proteoglycans

Interaction Granulocyte/macrophage/erythroid/megakaryocytic progenitors, myeloblasts,

monoblasts, dendritic cells, T cells.

Ligand/Receptor Heterodimer GM-CSFR α subunit (CDw116); β-subunit (CDw131) in common

Bioactivity Measured by its ability to induce proliferation of TF-1 erythroleukemic cells.

Cell Type Embryonic Stem Cells, Hematopoietic stem and progenitors

Biology Area Cell Biology, Stem Cells

Molecular Family Cytokines/Chemokines, Growth Factors

Antigen References

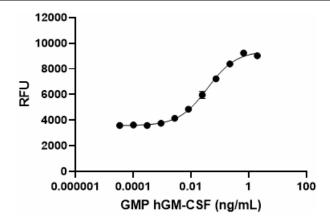
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2. Hayashida K, et al. 1990. Proc Natl Acad Sci USA. 87:9655-9.

3. Walker F & Burgess AW. 1985. EMBO J. 4:933-9.

4. Kitamura T, et al. 1989. J Cell Physiol. 140:323-34.

Gene ID <u>1437</u>



GMP recombinant human GM-CSF induces dose-dependent proliferation of TF-1 erythroleukemic cells. Deep Blue Cell Viability $^{\text{TM}}$ Kit (Cat. No. 424701) is used to measure the proliferation. The ED $_{50}$ range for this effect is 0.01-0.04 ng/mL.

Symbols Glossary*

Symbol	Meaning	Symbol Title	Symbol No.	Symbol	Meaning	Symbol Title	Symbol No.
REF	Catalog number	Catalogue number	5.1.6	(i	Indicates the need for the user to consult the instructions for use.	Consult instructions for use	5.4.3
1	Indicates the temperature limits to which the medical device can be safely exposed.	Temperature limit	5.3.7	淡	Indicates a medical device that needs protection from light sources.	Keep away from sunlight	5.3.2
K	Indicates the upper limit of temperature to which the medical device can be safely exposed.	temperature	5.3.6	Ω	Indicates the date after which the medical device is not to be used.	Use-by date	5.1.4
	Indicates the medical device manufacturer.	Manufacturer	5.1.1	EC REP	Indicates the authorized representative in the European Community.	Authorized representative in the European Community	5.1.2
	Indicates the manufacturer's	Batch code	5.1.5		Indicates a medical device that is	In vitro diagnostic	5.5.1
LOT	batch code so that the batch or lot can be identified.			IVD	intended to be used as an in vitro diagnostic medical device.	medical device	

* Symbol information is from EN ISO 15223-1:2016 Medical devices – Symbols to be used with medical device labels, labelling and information to be supplied – Part 1: General requirements

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