



GMP Recombinant Human SCF (carrier-free)

Catalog# / Size 573914 / 25 μg

573916 / 100 µg

Other Names Stem cell factor, KIT-ligand, Kitl, KL-1, mast cell grow factor (MGF), steel factor

(SF), FPH2, SHEP7

Description Stem Cell Factor (SCF) is initially synthesized as membrane-bound forms of 248

or 220 amino acids, depending on alternative splicing of exon 6. The 248 amino acid form contains a proteolytic cleavage site encoded by exon 6, and it is cleaved from the cell to release an active soluble protein of 165 amino acid residues. Soluble SCF is glycosylated at both N-linked and O-linked sites. MMP-9 plays a physiological role in SCF release from the membrane, and this action plays a significant role in differentiating and mobilizing stem and progenitors cells from the bone marrow. SCF increases the proliferation of myeloid and lymphoid hematopoietic progenitors in bone marrow cultures. SCF/c-kit interaction in mast cells results in mast cell degranulation with release of mediators, such as histamine and inflammatory cytokines and chemokines. Also, activation of c-kit in dendritic cells regulates T helper cell differentiation and allergic asthma. In addition, SCF plays an important role in revascularization of ischemic limbs. Ischemia induces plasma elevation of SCF and thrombopoietin (TPO) and, in lower levels, GM-CSF and erythropoietin (EPO). SCF and TPO induce the release of CXCL12 from platelets, thereby increasing CXCL12 levels in plasma. This results in an extensive mobilization of CXCR4+VEGFR1+ cells

(hemangiocytes), accelerating revascularization of the ischemic limbs. SCF binds

to its receptor kit that belongs to the type III tyrosine kinase family, whose

members include receptors for M-CSF and PDGF.

Product Details

Source Human SCF, amino acids Glu26-Ala189 (Accession# NM 000899.4) was

expressed in E. coli.

Molecular Mass The 165 amino acid recombinant protein has a predicted molecular mass of

approximately 18.6 kD. The DTT-reduced and non-reduced protein migrate at approximately 20 kD and 18 kD respectively by SDS-PAGE. The N-terminal amino

acid is Met.

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

Formulation 0.1 μm filtered protein solution is in PBS.

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 µg/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 = 3 - 12 ng/mL as determined by its ability to induce proliferation of TF-1

human erythroleukemic cells. Deep Blue Cell Viability™ Kit (Cat. No. 424701) is

used to measure the proliferation.

Application Biossay

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 proteins 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 Homodimer

Distribution Embryonic Stem Cells, Hematopoietic stem and progenitors

Function Plays an essential role in the regulation of cell survival and proliferation,

hematopoiesis, stem cell maintenance, gametogenesis, mast cell development,

migration and function, and in melanogenesis.

Interaction Mast cells, natural killer cells, dendritic cells, eosinophils, epithelial cells,

endothelial cells, melanocytes, germ cells, cholangiocytes, platelets, myeloid

leukemia cells, and intestinal cells of Cajal

Ligand/Receptor c-kit (CD117)

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

Cell Type Embryonic Stem Cells, Hematopoietic stem and progenitors

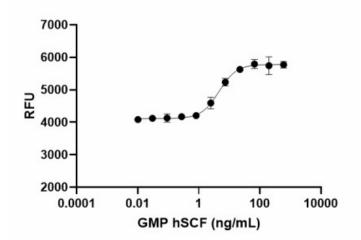
Biology Area Angiogenesis, Cell Biology, Immunology, Signal Transduction, Stem Cells

Molecular Family Cytokines/Chemokines, Growth Factors

Antigen References

- 1. Lu HS, et al. 1996. J. Biol. Chem. 271:11309.
- 2. Heissig B, et al. 2002. Cell 109:625.
- 3. Jin DK, et al. 2006. Nat. Med. 12:557.
- Krishnamoorthy N, et al. 2008. Nat. Med. 14:565.
 Ray P, et al. 2010. Ann. N. Y. Acad. Sci. 1183:104.

Gene ID <u>4254</u>



GMP recombinant human SCF induces proliferation of TF-1 human erythroleukemic cells in a dose-dependent manner with an ED₅₀ range of 3 - 12 ng/mL.

Symbols Glossary*

Symbol	Meaning	Symbol Title	Symbol No.	Symbol	Meaning	Symbol Title	Symbol No.
REF	Catalog number	Catalogue number	5.1.6	<u> </u>	Indicates the need for the user to consult the instructions for use.	Consult instructions for use	5.4.3
X	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
X	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
1	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
LOT	Indicates the manufacturer's batch code so that the batch or lot can be identified.	Batch code	5.1.5	IVD	Indicates a medical device that is intended to be used as an in vitro diagnostic medical device.	In vitro diagnostic medical device	5.5.1

^{*} 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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