



Overview

Description

SDF-1 beta/CXCL12, Mouse; **Synonyms**

> SDF-1a and SDF-1B, members of the chemokine a subfamily that lack the ELR domain. were initially identified using the signal sequence trap cloning strategy from a mouse bone-marrow stromal cell line. These proteins were subsequently also cloned from a human stromal cell line as cytokines that supported the proliferation of a stromal celldependent pre-B-cell line. SDF-1a and SDF-1ß cDNAs encode precursor proteins of 89 and 93 amino acid residues, respectively. Both SDF-1a and SDF-1ß are encoded by a

single gene and arise by alternative splicing. The two proteins are identical except for the four amino acid residues that are present in the carboxy-terminus of SDF-1ß and absent

from SDF-1^a. SDF-1/PBSF is highly conserved between species, with only one amino acid substitution between the mature human and mouse proteins. SDF-1/PBSF acts via the chemokine receptor CXCR4 and has been shown to be a chemoattractant for Tlymphocytes, monocytes, pro- and pre- B cells, but not neutrophils. Mice lacking SDF-1 or CXCR4 have been found to have impaired B-lymphopoiesis, myelopoiesis, vascular development, cardiogenesis and abnormal neuronal cell migration and patterning in the

central nervous system.

Species Mouse Source E. coli

Fully biologically active when compared to standard. The biological activity determined by

Biological Activity a chemotaxis bioassay using human peripheral blood monocytes is in a concentration

range of 50-100 ng/ml.

KPVSLSYRCP CRFFESHIAR ANVKHLKILN TPNCALQIVA Sequence

RLKNNNRQVC IDPKLKWIQE YLEKALNKRL KM

Properties

Measured Molecular Approximately 8.5 kDa, a single non-glycosylated polypeptide chain containing 72 amino

Weight acids.

> 97 % by SDS-PAGE and HPLC analyses. **Purity**

Lyophilized from a 0.2 µm filtered concentrated solution in 20 mM PB, pH 7.4, 150 mM **Formulation**

NaCl.

We recommend that this vial be briefly centrifuged prior to opening to bring the contents to the bottom. Reconstitute in sterile distilled water or aqueous buffer containing 0.1 % BSA

to a concentration of 0.1-1.0 mg/mL. Stock solutions should be apportioned into working aliquots and stored at d -20 °C. Further dilutions should be made in appropriate buffered

solutions.

Endotoxin Level Physical

Reconstitution

Less than 1 EU/µg of rMuSDF-12/CXCL122 as determined by LAL method.

Appearance

Storage

Sterile Filtered White lyophilized (freeze-dried) powder.

This material is for research, laboratory or further evaluation purposes. NOT FOR Usage

HUMAN USE.

This lyophilized preparation is stable at 2-8 °C, but should be kept at -20 °C for long term storage, preferably desiccated. Upon reconstitution, the preparation is stable for up to one week at 2-8 °C. For maximal stability, apportion the reconstituted preparation into working

aliquots and store at -20 °C to -70 °C. Avoid repeated freeze/thaw cycles.

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