

Screening Libraries

Proteins

Inhibitors

Product Data Sheet



CXCL14/BRAK Protein, Mouse (His)

Cat. No.: HY-P71887A

Synonyms: Cxcl14; Bmac; Kec; Ks1; Mip2g; Scyb14C-X-C motif chemokine 14; B-cell and monocyte-

activating chemokine; Chemokine BRAK; Kidney-expressed chemokine CXC; MIP-2G; Small-

inducible cytokine B14

Species: Mouse Source: E. coli

Q9WUQ5 (S23-E99) Accession:

Gene ID: 57266

Molecular Weight: Approximately 13 kDa

PROPERTIES

	c		
AA	Sec	uen	CE

SKCKCSRKGP KIRYSDVKKL EMKPKYPHCE EKMVIVTTKS MSRYRGQEHC LHPKLQSTKR FIKWYNAWNE KRRVYEE

Biological Activity

The biological activity determined by a chemotaxis bioassay using THP-1 human monocytic leukemia cells. The ED₅₀ for this effect is 1.081 ng/mL, corresponding to a specific activity is 9.25×10⁵ units/mg.

Appearance

Lyophilized powder.

Formulation

Lyophilized from a 0.2 μm filtered solution of 50 mM Tris-HCL, 300 mM NaCl, 300 mM arginine, pH 8.0.

Endotoxin Level

<1 EU/µg, determined by LAL method.

Reconsititution

It is not recommended to reconstitute to a concentration less than 100 μg/mL in ddH₂O. For long term storage it is recommended to add a carrier protein (0.1% BSA, 5% HSA, 10% FBS or 5% Trehalose).

Storage & Stability

Stored at -20°C for 2 years. After reconstitution, it is stable at 4°C for 1 week or -20°C for longer (with carrier protein). It is recommended to freeze aliquots at -20°C or -80°C for extended storage.

Shipping

Room temperature in continental US; may vary elsewhere.

DESCRIPTION

Background

CXCL14/BRAK protein functions as a chemotactic factor specifically attracting CESS B-cells and THP-1 monocytes, with no chemotactic effect on T-cells. This selective chemoattraction underscores its role in mediating the migration of B-cells and monocytes, contributing to immune responses within the microenvironment. The distinct chemotactic properties of CXCL14/BRAK highlight its specificity for certain immune cell types, suggesting a specialized role in regulating the recruitment and activation of B-cells and monocytes while not influencing T-cell migration.

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