

Recombinant Human CXCL16 (rHuCXCL16)

PrimeGene Technical Data Sheet

201-16 **Catalog Number:** Source:

E. coli

Molecular Weight:

Approximately 10 kDa, a single non-glycosylated polypeptide chain containing 89 amino acids.

Quantity:

10ug/100µg

AA Sequence:

Asn49-Pro137; Accession # NP_071342

Purity:

> 97 % by SDS-PAGE analyses.

Biological Activity:

Measured by its ability to chemoattract BaF3 mouse pro-B cells transfected with mouse CXCR6. The

ED₅₀ for this effect is 2.5-12 ng/mL.

Physical Appearance:

Sterile Filtered White lyophilized (freeze-dried) powder.

Formulation:

Lyophilized from 0.2 µm filtered concentrated solution in PBS.

Endotoxin:

Less than 1.0 EU/µg of rHuCXCL16 as determined by LAL method.

Reconstitution:

We recommend that this vial be briefly centrifuged prior to opening to bring the contents to the bottom. Reconstitute in sterile PBS to a concentration of 0.1 mg/mL. Further dilutions should be

made in appropriately buffered solutions.

Stability & Storage:

Use a manual defrost freezer and avoid repeated freeze-thaw cycles.

12 months from date of receipt, -20 to -70 °C as supplied.

1 month, 2 to 8 °C under sterile conditions after reconstitution.

3 months, -20 to -70 °C under sterile conditions after reconstitution.

Usage:

This material is offered by Shanghai PrimeGene Bio-Tech for research, laboratory or further

evaluation purposes. NOT FOR HUMAN USE.

Human CXCL16

CXC chemokine ligand 16 (CXCL16) is a type I membrane protein containing a non-ELR motif-containing CXC chemokine domain in its extracellular region. Together with Fractalkine (CX3CL1), CXCL16 constitute the only two transmembrane chemokines within the superfamily. The gene for human CXCL16 predicts a 273 amino acid (aa) residue precursor protein with a putative signal peptide, a CXC chemokine domain, a mucin-like spacer region, a transmembrane domain and a cytoplasmic domain with a potential tyrosine phosphorylation and SH2 protein-binding site. Mouse and human CXCL16 share 70% aa sequence similarity within their chemokine domains and 49% overall aa sequence identity.

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