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Recombinant Human G-CSFR/CD114 Protein (Fc Tag)

Catalog No. PKSH031749

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description

Synonyms CD114;CSF3R;G-CSF R;GCSFR

Species Human

Expression Host

Sequence

Met 1-Pro 621

Accession

NP_000751.1

Calculated Molecular Weight

Observed molecular weight

Tag

HEK293 Cells

Met 1-Pro 621

NP_000751.1

Calculated Molecular Weight

120-130 kDa

C-hFc

Bioactivity Measured by its ability to inhibit GCSF-induced proliferation of NFS60 mouse

myeloid cells. The ED50 for this ettect is typically 2-20 ng/ml in the presence of

0.125ng/ml of recombinant human GCSF.

Properties

Purity > 95 % as determined by reducing SDS-PAGE.

Endotoxin < 1.0 EU per µg of the protein as determined by the LAL method.

Storage Generally, lyophilized proteins are stable for up to 12 months when stored at -20 to

-80°C. Reconstituted protein solution can be stored at 4-8°C for 2-7 days. Aliquots

of reconstituted samples are stable at < -20°C for 3 months.

Shipping This product is provided as lyophilized powder which is shipped with ice packs.

Formulation Lyophilized from sterile PBS, pH 7.4

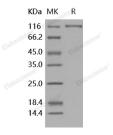
Normally 5 % - 8 % trehalose, mannitol and 0.01% Tween80 are added as

protectants before lyophilization.

Please refer to the specific buffer information in the printed manual.

Reconstitution Please refer to the printed manual for detailed information.

Data



> 95 % as determined by reducing SDS-PAGE.

Background

For Research Use Only

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Granulocyte Colony Stimulating Factor Receptor (G-CSFR), also known as CD114, which belongs to the cytokine receptor superfamily, is a cell surface receptor for colony stimulating factor 3 (CSF3). It is a critical regulator of granulopoiesis. This type I membrane protein has a composite structure consisting of an immunoglobulin(Ig)-like domain, a cytokine receptor-homologous (CRH) domain and three fibronectin type III (FNIII) domains in the extracellular region. Mutations in the G-CSF receptor leading to carboxy-terminal truncation transduce hyperproliferative growth responses, and are implicated in the pathological progression of severe congenital neutropenia (SCN) to acute myelogenous leukemia (AML). Additionally, autocrine/paracrine stimulation of G-CSFR may be important in the biology of solid tumors, including metastasis.

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