

Recombinant Human c-MET/HGFR Protein (aa 956-1390, His & GST Tag)

Catalog No. PKSH030396

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

Description

Synonyms AUTS9;c-Met;DFNB97;HGFR;RCCP2

Species Human

Expression Host Baculovirus-Insect Cells

Sequence Lys956-Ser1390

AccessionP08581-1Calculated Molecular Weight76.8 kDaObserved molecular weight68 kDaTagN-His-GST

Bioactivity 1. The specific activity was determined to be 10 nmol/min/mg using MBP as

substrate.

2. Immobilized human HGFR (aa 956-1390) at 10 µg/ml (100 µl/well) can bind

biotinylated human HGF-his with a linear range of 15.6-125 ng/ml.

Properties

Purity > 90 % as determined by reducing SDS-PAGE.

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

Storage Storage Store at < -20°C, stable for 6 months. Please minimize freeze-thaw cycles.

Shipping This product is provided as liquid. It is shipped at frozen temperature with blue

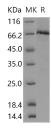
ice/gel packs. Upon receipt, store it immediately at < - 20°C.

Formulation Supplied as sterile solution of 20mM Tris, 500mM NaCl, pH 7.4, 10% glycerol,

3mM DTT

Reconstitution Not Applicable

Data



> 90 % as determined by reducing SDS-PAGE.

Background

Hepatocyte growth factor receptor (HGFR), also known as c-Met or mesenchymal-epithelial transition factor (MET), is a receptor tyrosine kinase (RTK) that has been shown to be overexpressed and/or mutated in a variety of malignancies.

For Research Use Only

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HGFR protein is produced as a single-chain precursor, and HGF is the only known ligand. Normal HGF/HGFR signaling is essential for embryonic development, tissue repair or wound healing, whereas aberrantly active HGFR has been strongly implicated in tumorigenesis, particularly in the development of invasive and metastatic phenotypes. HGFR protein is a multifaceted regulator of growth, motility, and invasion, and is normally expressed by cells of epithelial origin. Preclinical studies suggest that targeting aberrant HGFR signaling could be an attractive therapy in cancer.

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