Anti-Human HGFR / c-MET Antibody (PE)

Catalog Number: 10692-R243-P



General Information	
Immunogen:	Recombinant Human HGFR / c-MET protein (Catalog#10692-H08H)
Reagents:	PE-conjugated rabbit monoclonal antibody
Specificity:	Human HGFR / c-MET
Clone ID:	243
Ig Type:	Rabbit IgG
Applications:	Flow Cytometry
Concentration:	5 μl/Test, 0.1 mg/ml
Formulation:	Aqueous solution containing 0.5% BSA and 0.09% sodium azide
Storage:	2 °C - 8 °C in the dark

Preparation

This antibody was obtained from a rabbit immunized with purified, recombinant Human HGFR / c-MET (rh HGFR / c-MET; Catalog#10692-H08H; NP_000236; Met 1-Thr 932) and conjugated with PE under optimum conditions, the unreacted PE was removed.

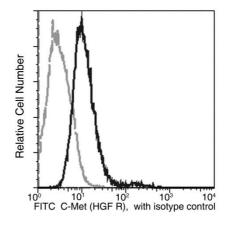
Storage

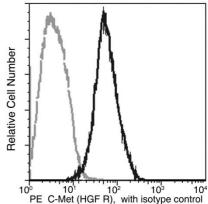
This antibody is stable for 12 months from date of receipt when stored at 2°C - 8°C. Protected from prolonged exposure to light. Do not freeze!

Sodium azide is toxic to cells and should be disposed of properly. Flush with large volumes of water during disposal

Applications

Flow Cytometry -





Profile of anti-human C-Met (HGF R) reactivity on HepG2 cells analyzed by flow cytometry. The Clone R243 and R271 (Cat. No. 10692-R271-F) monoclonal antibodies are not cross-blocking.

Flow cytometry was performed on a BD FACSCalibur flow cytometry system. Please refer to www.sinobiological.com/Flow-Cytometry-FACS-Protocols-a-750.html for technical protocols.

Specificity

Human HGFR / c-MET

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Background

The proto-oncogene MET encoded product MET (mesenchymal-epithelial transition factor), also known as c-Met or hepatocyte growth factor receptor (HGFR), is a multifaceted regulator of growth, motility, and invasion, and is normally expressed by cells of epithelial origin. As the prototypic member of a small subfamily of growth factor receptors, c-Met/HGFR is synthesized as a single chain precursor, and is processed into a mature disulfide-linked heterodimer composed of a extracellular α subunit and a transmembrane β subunit via post-translational cleavage. c-Met/HGFR is identified as a glycosylated receptor tyrosine kinase (RTK), and HGF is the only known ligand. Following ligand binding and autophosphorylation, c-Met/HGFR transmits intercellular signals using a unique multisubstrate docking site which mediates the binding of multiple SH2-containing adapter proteins such as Grb2, SHC, Crk/CRKL, as well as Gab1. Normal c-Met/HGFR signaling is essential for embryonic development, tissue repair or wound healing, whereas aberrantly active c-Met/HGFR has been strongly implicated in tumorigenesis, particularly in the development of invasive and metastatic phenotypes. Two transcript variants encoding different isoforms have been identified.

Reference

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- 3. Weidner, K.M. et al., 1996, Nature. 384: 173-176.
- 4. Pennacchietti, S. et al., 2003, Cancer. Cell. 3: 347-361.
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