

Product datasheet

Recombinant human Met (c-Met) (mutated M1250T) protein ab119729

2 Images

Description	
Product name	Recombinant human Met (c-Met) (mutated M1250T) protein
Biological activity	Specific activity of ab119729 was determined to be 185 nmol/min/mg as per activity assay protocol.
Purity	> 85 % Proprietary Purification. determined by densitometry. Affinity purified.
Expression system	Baculovirus infected Sf9 cells
Accession	<u>P08581</u>
Protein length	Protein fragment
Animal free	No
Nature	Recombinant
Species	Human
Predicted molecular weight	81 kDa including tags
Amino acids	956 to 1390

Specifications

Our Abpromise guarantee covers the use of ab119729 in the following tested applications.

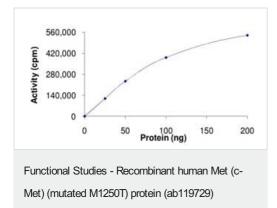
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications	Western blot
	Functional Studies
	SDS-PAGE
Form	Liquid
Preparation and Storage	
Stability and Storage	Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.
	pH: 7.50
	Constituents: 0.79% Tris buffered saline, 0.002% PMSF, 0.003% DTT, 0.003% EDTA, 25% Glycerol (glycerin, glycerine), 0.88% Sodium chloride

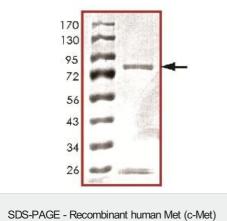
General Info

Function	Receptor for hepatocyte growth factor and scatter factor. Has a tyrosine-protein kinase activity.
	Functions in cell proliferation, scattering, morphogenesis and survival.
Involvement in disease	Note=Activation of MET after rearrangement with the TPR gene produces an oncogenic protein.
	Note=Defects in MET may be associated with gastric cancer.
	Defects in MET are a cause of hepatocellular carcinoma (HCC) [MIM:114550].
	Defects in MET are a cause of renal cell carcinoma papillary (RCCP) [MIM:605074]. It is a
	subtype of renal cell carcinoma tending to show a tubulo-papillary architecture formed by
	numerous, irregular, finger-like projections of connective tissue. Renal cell carcinoma is a
	heterogeneous group of sporadic or hereditary carcinoma derived from cells of the proximal renal
	tubular epithelium. It is subclassified into common renal cell carcinoma (clear cell, non-papillary
	carcinoma), papillary renal cell carcinoma, chromophobe renal cell carcinoma, collecting duct
	carcinoma with medullary carcinoma of the kidney, and unclassified renal cell carcinoma.
	Note=A common allele in the promoter region of the MET shows genetic association with
	susceptibility to autism in some families. Functional assays indicate a decrease in MET promoter
	activity and altered binding of specific transcription factor complexes.
	Note=MET activating mutations may be involved in the development of a highly malignant,
	metastatic syndrome known as cancer of unknown primary origin (CUP) or primary occult
	malignancy. Systemic neoplastic spread is generally a late event in cancer progression. However,
	in some instances, distant dissemination arises at a very early stage, so that metastases reach
	clinical relevance before primary lesions. Sometimes, the primary lesions cannot be identified in
	spite of the progresses in the diagnosis of malignancies.
Sequence similarities	Belongs to the protein kinase superfamily. Tyr protein kinase family.
	Contains 3 IPT/TIG domains.
	Contains 1 protein kinase domain.
	Contains 1 Sema domain.
Domain	The kinase domain is involved in SPSB1 binding.
Post-translational modifications	Dephosphorylated by PTPRJ at Tyr-1349 and Tyr-1365.
Cellular localization	Membrane.

Images



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The purity of ab119729 was determined to be >85% by densitometry, approx. MW 81 kDa.

SDS-PAGE - Recombinant human Met (c-Met) (mutated M1250T) protein (ab119729)

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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