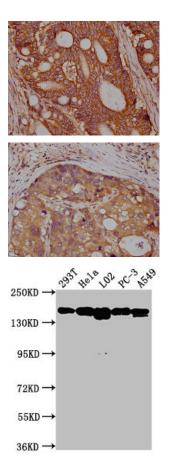
## **MET Recombinant Antibody**

## RAC00222



Product Information	
Size:	Protein Background:
50ul	<ul> <li>Receptor tyrosine kinase that transduces signals from the extracellular matrix into the cytoplasm by binding to hepatocyte growth factor/HGF ligand. Regulates many physiological processes including proliferation, scattering, morphogenesis and survival. Ligand binding at the cell surface induces autophosphorylation of MET on its intracellular domain that provides docking sites for downstream signaling molecules. Following activation by ligand, interacts with the PI3-kinase subunit PIK3R1, PLCG1, SRC, GRB2, STAT3 or the adapter GAB1.</li> <li>MET</li> <li>Diprot</li> <li>P08581</li> <li>Synonyms:</li> </ul>
Reactivity:	
Human	
Source:	
Homo sapiens (Human)	
lsotype:	
Rabbit IgG	
Applications:	
ELISA, WB, IHC, IF, FC	
Recommended dilutions:	
	Hepatocyte growth factor receptor (HGF receptor) (EC 2.7.10.1) (HGF/SF receptor) (Proto-oncogene c-Met) (Scatter factor receptor) (SF receptor) (Tyrosine-protein kinase Met), MET
	Immunogen:
	A synthesized peptide derived from human Met (c-Met).
	Storage:
	Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and

50% glycerol.



IHC image of RACO0222 diluted at 1:100 and staining in paraffinembedded human colon cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a Goat antirabbit IgG polymer labeled by HRP and visualized using 0.05% DAB.

IHC image of RACO0222 diluted at 1:100 and staining in paraffinembedded human liver cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a Goat antirabbit IgG polymer labeled by HRP and visualized using 0.05% DAB.

## Western Blot

Observed band size: 156 kDa

Positive WB detected in( 293T whole cell lysate) Hela whole cell lysate) L02 whole cell lysate) PC-3 whole cell lysate) A549 whole cell lysate) All lanes: MET antibody at 1:1500 Secondary Goat polyclonal to rabbit IgG at 1:50000 dilution Predicted band size: 156, 158, 86 kDa