#8041 store at -20C

Met (D1C2) XP[®] Rabbit mAb (Biotinylated)



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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: WB	Reactivity:	Sensitivity: Endogenous	MW (kDa): 140, 170	Source/Isotype: Rabbit IgG	UniProt ID: #P08581	Entrez-Gene Id: 4233	
Product Usage Information	Ар	plication		Dilution			
	We	stern Blotting			1:1000		
Storage		Supplied in 136 mM NaCl, 2.6 mM KCl, 12 mM sodium phosphate (pH 7.4) dibasic, 2 mg/ml BSA, and 50% glycerol. Store at –20°C. Do not aliquot the antibodies.					
Specificity / Sensitiv	/ity Met	Met (D1C2) XP [®] Rabbit mAb (Biotinylated) recognizes endogenous levels of total Met protein.					
Source / Purification	•	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human Met protein.					
Product Description	antil	This Cell Signaling Technology antibody is conjugated to biotin under optimal conditions. The biotinylated antibody is expected to exhibit the same species cross-reactivity as the unconjugated Met (D1C2) XP® Rabbit mAb #8198.					
MW (kDa)		140, 170					

Background

Met, a high affinity tyrosine kinase receptor for hepatocyte growth factor (HGF, also known as scatter factor) is a disulfide-linked heterodimer made of 45 kDa α - and 145 kDa β -subunits (1,2). The α -subunit and the amino-terminal region of the β -subunit form the extracellular domain. The remainder of the β -chain spans the plasma membrane and contains a cytoplasmic region with tyrosine kinase activity. Interaction of Met with HGF results in autophosphorylation at multiple tyrosines, which recruit several downstream signaling components, including Gab1, c-Cbl, and Pl3 kinase (3). These fundamental events are important for all of the biological functions involving Met kinase activity. The addition of a phosphate at cytoplasmic Tyr1003 is essential for Met protein ubiquitination and degradation (4). Phosphorylation at Tyr1234/1235 in the Met kinase domain is critical for kinase activation. Phosphorylation at Tyr1349 in the Met cytoplasmic domain provides a direct binding site for Gab1 (5). Research studies have shown that altered Met levels and/or tyrosine kinase activities are found in several types of tumors, including renal, colon, and breast. Thus, investigators have concluded that Met is an attractive potential cancer therapeutic and diagnostic target (6,7).

Background References

- 1. Cooper, C.S. et al. (1984) Nature 311, 29-33.
- 2. Bottaro, D.P. et al. (1991) Science 251, 802-4.
- 3. Bardelli, A. et al. (1997) *Oncogene* 15, 3103-11.
- 4. Taher, T.E. et al. (2002) J Immunol 169, 3793-800.
- 5. Schaeper, U. et al. (2000) J Cell Biol 149, 1419-32.
- 6. Eder, J.P. et al. (2009) Clin Cancer Res 15, 2207-14.
- 7. Sattler, M. and Salgia, R. (2009) Update Cancer Ther 3, 109-118.

Species Reactivity Species reactivity is determined by testing in at least one approved application (e.g., western blot).

openior reasons, is determined by testing in at reason one approved approaches (e.g., received approaches).

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry

milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key WB: Western Blotting

Cross-Reactivity Key

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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