## V07-TerrorgAcetyl-β-Catenin (Lys49) (D7C2) Rabbit<br/>mAb0206#



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Applications: W, IP	<b>Reactivity:</b> H M Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 92	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P35222	Entrez-Gene Id: 1499		
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 1:1000 1:200			
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less thar 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				ol and less than		
Specificity/Sen	sitivity	Acetyl-β-Catenin (Lys49) (D7C2) Rabbit mAb recognizes endogenous levels of β-catenin protein only when acetylated at Lys49.						
Source / Purific	cation	Monoclonal antibody is produced by immunizing animals with a synthetic acetylated peptide corresponding to residues surrounding Lys49 of human $\beta$ -catenin protein.						
Background		$\beta$ -catenin is a key downstream effector in the Wnt signaling pathway (1). It is implicated in two major biological processes in vertebrates: early embryonic development (2) and tumorigenesis (3). CK1 phosphorylates $\beta$ -catenin at Ser45. This phosphorylation event primes $\beta$ -catenin for subsequent phosphorylation by GSK-3 $\beta$ (4-6). GSK-3 $\beta$ destabilizes $\beta$ -catenin by phosphorylating it at Ser33, Ser37, and Thr41 (7). Mutations at these sites result in the stabilization of $\beta$ -catenin protein levels and have been found in many tumor cell lines (8).						
		Lys49 lies in a region that contains several Ser/Thr residues whose phosphorylation status regulates the stability of β-catenin. Research studies have shown that Lys49 is one of few residues frequently mutated in thyroid anaplastic carcinoma (9). CREB-binding protein (CBP) binds and acetylates β-catenin at Lys49 (10,11).						
Background Re	eferences	<ol> <li>Cadigan, K.M. and Nusse, R. (1997) <i>Genes Dev</i> 11, 3286-3305.</li> <li>Wodarz, A. and Nusse, R. (1998) <i>Annu Rev Cell Dev Biol</i> 14, 59-88.</li> <li>Polakis, P. (1999) <i>Curr Opin Genet Dev</i> 9, 15-21.</li> <li>Amit, S. et al. (2002) <i>Genes Dev</i> 16, 1066-76.</li> <li>Liu, C. et al. (2002) <i>Cell</i> 108, 837-47.</li> <li>Yanagawa, S. et al. (2002) <i>EMBO J</i> 21, 1733-42.</li> <li>Yost, C. et al. (1996) <i>Genes Dev</i> 10, 1443-54.</li> <li>Morin, P.J. et al. (1997) <i>Science</i> 275, 1787-90.</li> <li>Polakis, P. (2000) <i>Genes Dev</i> 14, 1837-51.</li> <li>Takemaru, K.I. and Moon, R.T. (2000) <i>J Cell Biol</i> 149, 249-54.</li> <li>Wolf, D. et al. (2002) <i>J Biol Chem</i> 277, 25562-7.</li> </ol>						
Species Reactiv	vity	Species reactivity is det	ermined by testin	g in at least one approve	ed application (e.g.,	western blot).		
Western Blot B	Buffer	IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				ו 5% w/v BSA, 1X		
Applications K	ey	W: Western Blotting IP: Immunoprecipitation						
Cross-Reactivit	ty Key	H: Human M: Mouse Mk: Monkey						
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