Phospho-Numb (Ser276) (D5C2) Rabbit mAb



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8286 # For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W, IF-IC	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 78	Source/Isotype: Rabbit IgG	UniProt ID: #P49757	Entrez-Gene Id: 8650		
Product Usage Information		Application Western Blotting Immunofluorescence	(Immunocytochem	iistry)		Dilution 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 than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.						
Specificity/Ser	sitivity	Phospho-Numb (Ser276) (D5C2) Rabbit mAb recognizes endogenous levels of numb protein only when phosphorylated at Ser276.						
Species predic based on 100% homology	ted to react sequence	Mouse, Rat, Chicken, Xenopus, Zebrafish, Bovine, Horse						
Source / Purifi	cation	Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser276 of human numb protein.						
Background	eferences	 Numb contains an amino-terminal phosphotyrosine binding (PTB) domain and carboxy-terminal endocytic binding motifs for α-adaptin and EH (Eps15 homology) domain-containing proteins, indicating a role in endocytosis (1,2). There are four mammalian Numb splicing isoforms that are differentially expressed and may have distinct functions (3-5). Numb acts as a negative regulator of Notch signaling by promoting ubiquitination and degradation of Notch (6). The protein is asymmetrically segregated into one daughter cell during cell division, producing two daughter cells with different responses to Notch signaling and different cell fates (7,8). The localization of Numb can also be regulated by G protein-coupled receptor (GPCR) and protein kinase C (PKC) signaling (9). Numb can be phosphorylated at several sites including Ser7, Ser276, and Ser295. Phosphorylation at these sites regulates asymmetric membrane localization of Numb and integrin endocytosis (10-12). 1. Berdnik, D. et al. (2002) <i>Dev. Cell</i> 3, 221-231. 2. Santolini, E. et al. (2000) <i>J. Cell Biol.</i> 151, 1345-1352. 3. Dho, S.E. et al. (1999) <i>J. Biol. Chem.</i> 274, 33097-33104. 4. Verdi, J.M. et al. (1999) <i>Proc. Natl. Acad. Sci. USA</i> 96, 10472-10476. 5. Toriya, M. et al. (2006) <i>Dev. Neurosci.</i> 28, 142-155. 6. McGill, M.A. and McGlade, C.J. (2003) <i>J. Biol. Chem.</i> 278, 23196-23203. 7. Verdi, J.M. et al. (1996) <i>Curr. Biol.</i> 6, 1134-1145. 8. Reugels, A.M. et al. (2006) <i>Dev. Dyn.</i> 235, 934-948. 9. Dho S.E. et al. (2006) <i>Mol. Biol. Cell.</i> 17, 4142-4155. 						
		11. Smith, C.A. et al. (2 12. Wirtz-Peitz, F. et al	2007) <i>EMBO J</i> 26, 46 . (2008) <i>Cell</i> 135, 16	8-80. 1-73.				
Species Reacti	vity	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).		
Western Blot E	Buffer	IMPORTANT: For west dry milk, 1X TBS, 0.1%	≀TANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat lk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.					
Applications K	ey	W: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)						
Cross-Reactivi	ty Key	H: Human						
Trademarks ar	nd Patents	Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc.						

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