## Semaphorin 3B (D5A2P) Rabbit mAb





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Applications: W, IP	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 80	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q13214	Entrez-Gene Id: 7869	
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 1:1000 1:50		
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.					
Semaphorin 3B (D5A2P) Rabbit mAb recogn				gnizes endogenous leve	l of total semaphor	in 3B protein.	
Source / Purific	cation	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gln717 of human semaphorin 3B protein.					
Background		Semaphorins are a family of cell surface and secreted proteins initially recognized as axon guidance factors that control the development of the central nervous system (1). They are involved in cell migration, angiogenesis, and immune responses (2-6). Based on protein structure, there are eight classes of semaphorins. Class 3-7 semaphorins are expressed in vertebrates. Semaphorin 3 subfamily members are the only secreted semaphorins in vertebrates. There are seven semaphorin 3 proteins and their receptors include neuropilins and the type A/D family plexins (7-9).					
		Semaphorin 3B functions as a tumor suppressor, as research studies have shown that it is deleted or inactivated in lung and breast cancer (10,11). Overexpression of semaphorin 3B inhibits tumor cell proliferation and causes apoptosis (12,13). Semaphorin 3B also inhibits angiogenesis (14). Semaphorin 3B loses its activity upon cleavage by furin-like pro-protein convertases (14).					
Background Re	eferences	<ol> <li>Luo, Y. et al. (1993) <i>Cell</i> 75, 217-27.</li> <li>Kruger, R.P. et al. (2005) <i>Nat Rev Mol Cell Biol</i> 6, 789-800.</li> <li>Staton, C.A. (2011) <i>Biochem Soc Trans</i> 39, 1565-70.</li> <li>Ghanem, R.C. et al. (2011) <i>Curr Eye Res</i> 36, 989-96.</li> <li>Nakagawa, Y. et al. (2011) <i>J Immunol</i> 186, 2881-8.</li> <li>Suzuki, K. et al. (2008) <i>Nat Immunol</i> 9, 17-23.</li> <li>He, Z. and Tessier-Lavigne, M. (1997) <i>Cell</i> 90, 739-51.</li> <li>Kolodkin, A.L. et al. (1997) <i>Cell</i> 90, 753-62.</li> <li>Chen, H. et al. (2001) <i>Proc Natl Acad Sci U S A</i> 98, 13954-9.</li> <li>Nasarre, P. et al. (2005) <i>Med Sci (Paris)</i> 21, 641-7.</li> <li>Tse, C. et al. (2002) <i>Cancer Res</i> 62, 542-6.</li> <li>Castro-Rivera, E. et al. (2008) <i>Cancer Res</i> 68, 6922-31.</li> </ol>					
Species Reactiv	vity	Species reactivity is det	ermined by testing	j in at least one approve	d application (e.g.,	western blot).	
Western Blot B	Suffer	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	су Кеу	H: Human					
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