Semaphorin 3B Antibody



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

Applications: W, IP	Reactivity:	Sensitivity: Endogenous	MW (kDa): 80	Source/Isotype: Rabbit	UniProt ID: #Q13214	Entrez-Gene Id 7869
Product Usage Information		ApplicationDilutionWestern Blotting1:1000Immunoprecipitation1:50				
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Semaphorin 3B Antibody recognizes endogenous levels of total semaphorin 3B protein.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human semaphorin 3B protein. Antibodies are purified by protein A and peptide affinity chromatography.				
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 References		1. Luo, Y. et al. (1993) <i>Cell</i> 75, 217-27. 2. Kruger, R.P. et al. (2005) <i>Nat Rev Mol Cell Biol</i> 6, 789-800. 3. Staton, C.A. (2011) <i>Biochem Soc Trans</i> 39, 1565-70. 4. Ghanem, R.C. et al. (2011) <i>Curr Eye Res</i> 36, 989-96. 5. Nakagawa, Y. et al. (2011) <i>J Immunol</i> 186, 2881-8. 6. Suzuki, K. et al. (2008) <i>Nat Immunol</i> 9, 17-23. 7. He, Z. and Tessier-Lavigne, M. (1997) <i>Cell</i> 90, 739-51. 8. Kolodkin, A.L. et al. (1997) <i>Cell</i> 90, 753-62. 9. Chen, H. et al. (1997) <i>Neuron</i> 19, 547-59. 10. Tomizawa, Y. et al. (2001) <i>Proc Natl Acad Sci U S A</i> 98, 13954-9. 11. Nasarre, P. et al. (2005) <i>Med Sci (Paris)</i> 21, 641-7. 12. Tse, C. et al. (2002) <i>Cancer Res</i> 62, 542-6. 13. Castro-Rivera, E. et al. (2004) <i>Proc Natl Acad Sci U S A</i> 101, 11432-7. 14. Varshavsky, A. et al. (2008) <i>Cancer Res</i> 68, 6922-31.				

Species Reactivity

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

Western Blot 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.

Applications Key W: Western Blotting IP: Immunoprecipitation

Cross-Reactivity Key H: Human

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