

## **SH2D1A Antibody**



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

<b>Applications:</b> W, IP	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 14	<b>Source/Isotype:</b> Rabbit	UniProt ID: #O60880	Entrez-Gene Id 4068
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 and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		SH2D1A Antibody detects endogenous levels of total SH2D1A protein.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues around Gly112 of human SH2D1A. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		SH2D1A and SH2D1B are small, adaptor proteins with a single SH2-domain that play important signal transduction roles mediated by the signaling lymphocytic activation molecule (SLAM) family receptors (1). SH2D1A (also called SAP or SLAM-associated protein) is frequently mutated in patients with X-linked lymphoproliferative disease (Duncan's disease), which is characterized by extreme susceptibility to Epstein-Barr virus; approximately 50 different SH2D1A mutations have been reported to date (2-4). The single SH2D1B gene in humans (also called EAT-2 or Ewing's sarcoma's/FLI1-activated transcript 2) is present as a pair of duplicated EAT-2A and EAT-2B genes with identical genomic organization in mouse and rat (5,6).				
Background References		1. Latour, S. and Veillette, A. (2004) <i>Semin. Immunol.</i> 16, 409-419. 2. Nichols, K.E. et al. (2005) <i>Immunol. Rev.</i> 203, 180-199. 3. Engel, P. et al. (2003) <i>Nat. Rev. Immunol.</i> 3, 813-821. 4. Sumegi, J. et al. (2002) <i>Leuk. Lymphoma</i> 43, 1189-1201. 5. Roncagalli, R. et al. (2005) <i>Nat. Immunol.</i> 6, 1002-1010. 6. Calpe, S. et al. (2006) <i>Immunogenetics</i> 58, 15-25.				

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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