## **Syk Antibody**



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

Applications: W	Reactivity: H M	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 72	Source/Isotype: Rabbit	UniProt ID: #P43405	Entrez-Gene Id: 6850	
Product Usage Information		<b>Application</b> Western Blotting			<b>Dilution</b> 1:1000		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.					
Specificity/Sensitivity		Syk Antibody detects endogenous levels of total Syk protein. It does not cross-react with other members of the Syk/Zap-70 tyrosine kinase family.					
Species predicted to react based on 100% sequence homology		Rat					
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to the carboxy-terminal sequence of human Syk. Antibodies are purified by protein A and peptide affinity chromatography.					
Background		Syk is a protein tyrosine kinase that plays an important role in intracellular signal transduction in hematopoietic cells (1-3). Syk interacts with immunoreceptor tyrosine-based activation motifs (ITAMs) located in the cytoplasmic domains of immune receptors (4). It couples the activated immunoreceptors to downstream signaling events that mediate diverse cellular responses, including proliferation, differentiation, and phagocytosis (4). There is also evidence of a role for Syk in nonimmune cells and investigators have indicated that Syk is a potential tumor suppressor in human breast carcinomas (5). Tyr323 is a negative regulatory phosphorylation site within the SH2-kinase linker region in Syk. Phosphorylation at Tyr323 provides a direct binding site for the TKB domain of Cbl (6,7). Tyr352 of Syk is involved in the association of PLCγ1 (8). Tyr525 and Tyr526 are located in the activation loop of the Syk kinase domain; phosphorylation at Tyr525/526 of human Syk (equivalent to Tyr519/520 of mouse Syk) is essential for Syk function (9).					
Background References		2. Kurosaki, T. (1997) ( 3. Chu, D.H. et al. (199 4. Turner, M. et al. (200 5. Coopman, P.J. et al. 6. Deckert, M. et al. (1 7. Rao, N. et al. (2001) 8. Law, C.L. et al. (1996	1. Cheng, A.M. and Chan, A.C. (1997) <i>Curr Opin Immunol</i> 9, 528-33. 2. Kurosaki, T. (1997) <i>Curr Opin Immunol</i> 9, 309-18. 3. Chu, D.H. et al. (1998) <i>Immunol Rev</i> 165, 167-80. 4. Turner, M. et al. (2000) <i>Immunol Today</i> 21, 148-54. 5. Coopman, P.J. et al. (2000) <i>Nature</i> 406, 742-7. 6. Deckert, M. et al. (1998) <i>J Biol Chem</i> 273, 8867-74. 7. Rao, N. et al. (2001) <i>EMBO J</i> 20, 7085-95. 8. Law, C.L. et al. (1996) <i>Mol Cell Biol</i> 16, 1305-15. 9. Zhang, J. et al. (2000) <i>J Biol Chem</i> 275, 35442-7.				
Species Reactiv	rity	Species reactivity is de	etermined by testin	g in at least one approve	ed 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

Cross-Reactivity Key H: Human M: Mouse

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