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## Phospho-Zap-70 (Tyr319)/Syk (Tyr352) Antibody



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

Applications: W, W-S, IF-IC	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 70 Zap-70, 72 Syk	<b>Source/Isotype:</b> Rabbit	<b>UniProt ID:</b> #P43403, #P43405	Entrez-Gene Id: 7535, 6850		
Product Usage Information		<b>Application</b> Western Blotting Simple Western™ Immunofluorescence (Immunocytochemistry)		itry)	<b>Dilution</b> 1:1000 1:10 - 1:50 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/Sen	sitivity	Phospho-Zap-70 (Tyr319)/Syk (Tyr352) Antibody detects endogenous levels of Zap-70 only when phosphorylated at phosphorylated at Tyr319. It cross-reacts with endogenous levels of Syk when phosphorylated at Tyr352.						
Species predict based on 100% homology		Mouse, Rat, Hamster, Monkey, Chicken, Bovine, Dog, Pig, Horse						
Source / Purific	cation	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Tyr319 of human Zap-70. Antibodies are purified by protein A and peptide affinity chromatography.						
Background		The Syk family protein tyrosine kinase Zap-70 is expressed in T and NK cells and plays a critical role in mediating T cell activation in response to T cell receptor (TCR) engagement (1). Following TCR engagement, Zap-70 is rapidly phosphorylated on several tyrosine residues through autophosphorylation and transphosphorylation by the Src family tyrosine kinase Lck (2-6). Tyrosine phosphorylation correlates with increased Zap-70 kinase activity and downstream signaling events. Expression of Zap-70 is correlated with disease progression and survival in patients with chronic lymphocytic leukemia (7,8). Phosphorylation of Tyr319 is required for the assembly of a Zap-70-containing signaling complex that leads to the activation of the PLC-gamma1-dependent and Ras-dependent signaling cascades in antigen-stimulated T cells (5,6). The orthologous Tyr352 residue in Syk is also involved in the association with PLC-gamma1 (9).						
Background Re	eferences	<ol> <li>Chu, D.H. et al. (1998) <i>Immunol Rev</i> 165, 167-80.</li> <li>Iwashima, M. et al. (1994) <i>Science</i> 263, 1136-9.</li> <li>Neumeister, E.N. et al. (1995) <i>Mol Cell Biol</i> 15, 3171-8.</li> <li>Chan, A.C. et al. (1995) <i>EMBO J</i> 14, 2499-508.</li> <li>Williams, B.L. et al. (1999) <i>EMBO J</i> 18, 1832-44.</li> <li>Di Bartolo, V. et al. (1999) <i>J Biol Chem</i> 274, 6285-94.</li> <li>Wiestner, A. et al. (2003) <i>Blood</i> 101, 4944-51.</li> <li>Crespo, M. et al. (2003) <i>N Engl J Med</i> 348, 1764-75.</li> <li>Law, C.L. et al. (1996) <i>Mol Cell Biol</i> 16, 1305-15.</li> </ol>						
Species Reactiv	vity	Species reactivity is determined by testing in at least one approved application (e.g., western blot).						
Western Blot B	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 K	ey	W: Western Blotting W-S: Simple Western™ IF-IC: Immunofluorescence (Immunocytochemistry)						
Cross-Reactivit	у Кеу	H: Human						
Trademarks an	d Patents	Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc.						

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