Phospho-Stat6 (Tyr641) (C11A12) Rabbit mAb 7926#



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Applications: W	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 100	Source/Isotype: Rabbit	UniProt ID: #P42226	Entrez-Gene Id: 6778			
Product Usage Information		Application Western Blotting			Dilution 1:1000				
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.							
Specificity/Sen	sitivity	Phospho-Stat6 (Tyr641) (C11A12) Rabbit mAb detects endogenous levels of Stat6 only when phosphorylated at Tyr641. This antibody does not cross-react with the corresponding phospho-tyrosine residues of other Stat proteins. Cross-reactivity has been observed with receptor tyrosine kinases including phosphorylated PDGF receptor that can lead to inapproriate membrane staining.							
Species predict based on 100% homology		Monkey							
Source / Purific	cation	Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Tyr641 of human Stat6.							
Background		Upon activation by Janus kinases, Stat6 translocates to the nucleus where it regulates cytokine-induced gene expression. Stat6 is activated via phosphorylation at Tyr641 and is required for responsiveness to IL-4 and IL-13 (1-4). In addition, Stat6 is activated by IFN-α in B cells, where it forms transcriptionally active complexes with Stat2 and p48 (5,6). Protein phosphatase 2A is also involved in regulation of IL-4-mediated Stat6 signaling (7).							
Background Re	eferences	1. Nelms, K. et al. (1999) <i>Ann. Rev. Immunol.</i> 17, 701-738. 2. Malabarba, M.G. et al. (1996) <i>Biochem. J.</i> 319, 865-872. 3. Hou, J. et al. (1994) <i>Science</i> 265, 1701-1706. 4. Quelle, F.W. et al. (1995) <i>Mol. Cell. Biol.</i> 15, 3336-3343. 5. Takeda, K. et al. (1996) <i>Nature</i> 380, 627-630. 6. Gupta, S. et al. (1999) <i>J. Immunol.</i> 163, 3834-3841. 7. Woetmann, A. et al. (2003) <i>J. Biol. Chem.</i> 278, 2787-2791.							
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							
Cross-Reactivit	ty Key	H: Human							
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