## PP2A A Subunit (6G3) Rat mAb





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Applications: W	<b>Reactivity:</b> H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 62	<b>Source/Isotype:</b> Rat IgG2a	<b>UniProt ID:</b> #P30153	Entrez-Gene Id: 5518	
Product Usage Information		<b>Application</b> 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	PP2A A Subunit (6G3) Antibody detects endogenous levels of PP2A A subunit, alpha isoform. The antibody may also detect the beta isoform. This antibody does not cross-react with other PP2A subunits.					
Species predict based on 100% homology	ted to react sequence	Xenopus, Bovine					
Source / Purific	cation	Monoclonal antibody is produced by immunizing animals with the full-length PP2A A alpha protein.					
Background		Protein phosphatase type 2A (PP2A) is an essential protein serine/threonine phosphatase that is conserved in all eukaryotes. PP2A is a key enzyme within various signal transduction pathways as it regulates fundamental cellular activities such as DNA replication, transcription, translation, metabolism, cell cycle progression, cell division, apoptosis and development (1-3). The core enzyme consists of catalytic C and regulatory A (or PR65) subunits, with each subunit represented by $\alpha$ and $\beta$ isoforms (1). Additional regulatory subunits belong to four different families of unrelated proteins. Both the B (or PR55) and B' regulatory protein families contain $\alpha$ , $\beta$ , $\gamma$ and $\delta$ isoforms, while striatin (PR10) and SG2NA (PR93) are both members of the B''' regulatory protein family. These B subunits competitively bind to a shared binding site on the core A subunit (1). This variable array of holoenzyme components, particularly regulatory B subunits, allows PP2A to act in a diverse set of functions. PP2A function is regulated by expression, localization, holoenzyme composition and post-translational modification. Phosphorylation of PP2A at Tyr307 by Src occurs in response to EGF or insulin and results in a substantial reduction of PP2A activity (4). Reversible methylation on the carboxyl group of Leu309 of PP2A has been observed (5,6). Methylation alters the conformation of PP2A, as well as its localization and association with B regulatory subunits (6-8).					
Background Re	eferences	<ol> <li>Janssens, V. and Goris, J. (2001) <i>Biochem J</i> 353, 417-39.</li> <li>Zolnierowicz, S. (2000) <i>Biochem Pharmacol</i> 60, 1225-35.</li> <li>Millward, T.A. et al. (1999) <i>Trends Biochem Sci</i> 24, 186-91.</li> <li>Chen, J. et al. (1992) <i>Science</i> 257, 1261-4.</li> <li>Turowski, P. et al. (1995) <i>J Cell Biol</i> 129, 397-410.</li> <li>Lee, J. et al. (1996) <i>Proc Natl Acad Sci U S A</i> 93, 6043-7.</li> <li>Tolstykh, T. et al. (2001) <i>Mol Biol Cell</i> 12, 185-99.</li> <li>Kremmer, E. et al. (1997) <i>Mol Cell Biol</i> 17, 1692-701.</li> </ol>					
Species Reactiv	vitv	Species reactivity is de	etermined by testin	g in at least one approve	ed 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 M: Mouse R: Rat Mk: Monkey					
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