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## Phospho-RCC1 (Ser11) (D18B5) Rabbit mAb



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Applications: W	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 45	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P18754	Entrez-Gene Id: 1104		
Product Usage Information		Application Western Blotting		Dilution 1:1000				
			n 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than ium azide. Store at –20°C. Do not aliquot the antibody.					
Specificity/Sensitivity		Phospho-RCC1 (Ser11) (D18B5) Rabbit mAb recognizes endogenous levels of RCC1 protein only when phosphorylated at Ser11.						
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser11 of human RCC1 protein.						
Background		The Ras family small GTPase Ran is involved in nuclear envelope formation, assembly of the mitotic spindle, and nuclear transport (1,2). Like other small GTPases, Ran is active in its GTP-bound form and inactive in its GDP-bound form. Nuclear RanGTP concentration is maintained through nuclear localization of guanine nucleotide exchange factor (GEF) activity, which catalyzes the exchange of bound GDP for GTP. Regulator of chromatin condensation 1 (RCC1) is the only known RanGEF (3). RCC1 is dynamically chromatin-bound throughout the cell cycle, and this localization is required for mitosis to proceed normally (4,5). Appropriate association of RCC1 with chromatin is regulated through aminoterminal phosphorylation (5,6) and methylation (7). RCC1 regulation of RanGTP levels in response to histone modifications regulates nuclear import during apoptosis (8). In mitosis RCC1 is phosphorylated at Ser11, possibly by cyclin B/cdc2 (9-11). This phosphorylation may play a role in RCC1 interaction with chromatin and RCC1 RanGEF activity (6).						
Background R	eferences	<ol> <li>Quimby, B.B. and Dasso, M. (2003) <i>Curr Opin Cell Biol</i> 15, 338-44.</li> <li>Hetzer, M. et al. (2002) <i>Nat Cell Biol</i> 4, E177-84.</li> <li>Moore, W. et al. (2002) <i>Curr Biol</i> 12, 1442-7.</li> <li>Ohtsubo, M. et al. (1989) <i>J Cell Biol</i> 109, 1389-97.</li> <li>Li, H.Y. and Zheng, Y. (2004) <i>Genes Dev</i> 18, 512-27.</li> <li>Hutchins, J.R. et al. (2004) <i>Curr Biol</i> 14, 1099-104.</li> <li>Chen, T. et al. (2007) <i>Nat Cell Biol</i> 9, 596-603.</li> <li>Wong, C.H. et al. (2009) <i>Nat Cell Biol</i> 11, 36-45.</li> <li>Horiike, Y. et al. (2009) <i>Mol Biol Rep</i> 36, 717-23.</li> <li>Dephoure, N. et al. (2008) <i>Proc Natl Acad Sci U S A</i> 105, 10762-7.</li> <li>Hood, F.E. and Clarke, P.R. (2007) <i>J Cell Sci</i> 120, 3436-45.</li> </ol>						
Species Reacti	vity	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).		
Western Blot E	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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