## PAK2 (C17A10) Rabbit mAb





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Applications: W	<b>Reactivity:</b> H M Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 61	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q13177	Entrez-Gene Id: 5062		
Product Usage Information	2	<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/Sensitivity		PAK2 (C17A10) Rabbit mAb detects endogenous levels of total PAK2 protein. This antibody does not cross-react with PAK1, PAK3 or PAK4-6.						
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to the amino-terminal region of human PAK2.						
Background		The p21-activated kinase (PAK) family of serine/threonine kinases is engaged in multiple cellular processes, including cytoskeletal reorganization, MAPK signaling, apoptotic signaling, control of phagocyte NADPH oxidase, and growth factor-induced neurite outgrowth (1,2). Several mechanisms that induce PAK activity have been reported. Binding of Rac/Cdc42 to the CRIB (or PBD) domain near the amino terminus of PAK causes autophosphorylation and conformational changes in PAK (1). Phosphorylation of PAK1 at Thr423 by PDK induces activation of PAK1 (3). Several autophosphorylation sites have been identified, including Ser199 and Ser204 of PAK1, and Ser192 and Ser197 of PAK2 (4,5). Because the autophosphorylation sites are located in the amino-terminal inhibitory domain, it has been hypothesized that modification in this region prevents the kinase from reverting to an inactive conformation (6). Research indicates that phosphorylation at Ser144 of PAK1 or Ser139 of PAK3 (located in the kinase inhibitory domain) affects kinase activity (7). Phosphorylation at Ser21 of PAK1 or Ser20 of PAK2 regulates binding with the adaptor protein Nck (8). PAK4, PAK5/7, and PAK6 have lower sequence similarity with PAK1-3 in the amino-terminal regulatory region (9). Phosphorylation at Ser474 of PAK4, a site analogous to Thr423 of PAK1, may play a pivotal role in regulating the activity and function of PAK4 (10). PAK family members are widely expressed, and often overexpressed in human cancer (11,12).						
Background References		<ol> <li>Knaus, U.G. and Bokoch, G.M. (1998) Int. J. Biochem. Cell Biol. 30, 857-62.</li> <li>Daniels, R.H. et al. (1998) EMBO J. 17, 754-64.</li> <li>King, C.C. et al. (2000) J. Biol. Chem. 275, 41201-9.</li> <li>Manser, E. et al. (1997) Mol. Cell. Biol. 17, 1129-43.</li> <li>Gatti, A. et al. (1999) J. Biol. Chem. 274, 8022-8.</li> <li>Lei, M. et al. (2000) Cell 102, 387-97.</li> <li>Chong, C. et al. (2000) Mol. Cell. Biol. 20, 3906-17.</li> <li>Abo, A. et al. (1998) EMBO J. 17, 6527-40.</li> <li>Qu, J. et al. (2001) Mol. Cell. Biol. 21, 3523-33.</li> <li>Wen, Y.Y. et al. (2004) Expert Opin Ther Targets 18, 807-15.</li> <li>Molli, P.R. et al. (2009) Oncogene 28, 2545-55.</li> </ol>						
Species Reactivity		Species reactivity is determined by testing in at least one approved 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 Mk: Monkey						
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