PKCδ Antibody



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

Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:	
W, IP	H M R Mk	Endogenous	78	Rabbit	#Q05655	5580	
Product Usage Information	2	Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:25		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.					
Specificity/Sensitivity		PKC δ Antibody detects endogenous levels of total PKC δ protein. The antibody does not cross-react with endogenous levels of other PKC isoforms.					
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to human PKCδ. Antibodies are purified by protein A and peptide affinity chromatography.					
Background		human PKC8. Antibodies are purified by protein A and peptide affinity chromatography. Activation of protein kinase C (PKC) is one of the earliest events in a cascade that controls a variety of cellular responses, including secretion, gene expression, proliferation, and muscle contraction (1,2). PKC isoforms belong to three groups based on calcium dependency and activators. Classical PKCs are calcium-dependent via their C2 domains and are activated by phosphatidylserine (PS), diacylglycerol (DAG), and phorbol esters (TPA, PMA) through their cysteine-rich C1 domains. Both novel and atypical PKCs are calcium-independent, but only novel PKCs are activated by PS, DAG, and phorbol esters (3-5). Members of these three PKC groups contain a pseudo-substrate or autoinhibitory domain that binds to substrate-binding sites in the catalytic domain to prevent activation in the absence of cofactors or activators. Control of PKC activity is regulated through three distinct phosphorylation events. Phosphorylation occurs <i>in vivo</i> at Thr500 in the activation loop, at Thr641 through autophosphorylation, and at the carboxy-terminal hydrophobic site Ser660 (2). Atypical PKC isoforms lack hydrophobic region phosphorylation, which correlates with the presence of glutamic acid rather than the serine or threonine residues found in more typical PKC isoforms. The enzyme PDK1 or a close relative is responsible for PKC activation. A recent addition to the PKC superfamily is PKCµ (PKD), which is regulated by DAG and TPA through its C1 domain. PKD is distinguished by the presence of a PH domain and by its unique substrate recognition and Golgi localization (6). PKC-related kinases (PRK) lack the C1 domain and do not respond to DAG or phorbol esters. Phosphotyldinositol lipids activate PRKs, and small Rho-family GTPases bind to the homology region 1 (HR1) to regulate PRK kinase activity (7). Phosphorylatioin of tyrosine residues in PKCdelta are suggested to play a role in determining its functional properties. Phosphorylated tyrosine residue					
Background References		3. Mellor, H. and Parke 4. Ron, D. and Kazanie 5. Moscat, J. and Diaz-I 6. Baron, C.L. and Mall 7. Flynn, P. et al. (2000) 8. Steinberg, S.F. (2004	(1995) <i>Curr Biol</i> 5, r, P.J. (1998) <i>Bioche</i> tz, M.G. (1999) <i>FAS</i> Meco, M.T. (2000) <i>E</i> notra, V. (2002) <i>Scio</i> J <i>Biol Chem</i> 275, 1 Biochem J 384, 44	95) Curr Biol 5, 1394-403. J. (1998) Biochem J 332 (Pt 2), 281-92. M.G. (1999) FASEB J 13, 1658-76. Proposition of the second of the secon			

10. Konishi, H. et al. (2001) *Proc Natl Acad Sci U S A* 98, 6587-92.

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 IP: Immunoprecipitation

Cross-Reactivity Key H: Human M: Mouse R: Rat Mk: Monkey

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