PLCγ1 (D9H10) XP[®] Rabbit mAb 0695#



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Applications: W, W-S, IP, IHC-P	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 150	Source/Isotype: Rabbit IgG	UniProt ID: #P19174	Entrez-Gene Id: 5335		
Product Usage Information		Application			Dilutio	n		
information		5	Western Blotting			1:1000 1:10 - 1:50		
		Simple Western™			1:50	.50		
			Immunoprecipitation			1:50 - 1:200		
		Immunohistochemistry (Paraffin)			1.50 - 1.200			
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less tha 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				ol and less than		
		For a carrier free (BSA and azide free) version of this product see product #10607.						
Specificity/Sen	•	PLC γ 1 (D9H10) XP $^{ extsf{B}}$ Rabbit mAb recognizes endogenous levels of total PLC γ 1 protein.						
Source / Purific	cation	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding residues surrounding Leu1220 of human PLCγ1 protein.						
Background		Phosphoinositide-specific phospholipase C (PLC) plays a significant role in transmembrane signaling. In response to extracellular stimuli, such as hormones, growth factors, and neurotransmitters, PLC hydrolyzes phosphatidylinositol 4,5-bisphosphate (PIP ₂) to generate two secondary messengers: inositol 1,4,5-triphosphate (IP ₃) and diacylglycerol (DAG) (1). At least four families of PLCs have been identified: PLC β , PLC γ , PLC δ , and PLC ϵ . Phosphorylation is one of the key mechanisms that regulate the activity of PLC. PLC γ is activated by both receptor and non-receptor tyrosine kinases (2). PLC γ forms a complex with EGF and PDGF receptors, which leads to the phosphorylation of PLC γ at Tyr771, 783, and 1248 (3). Phosphorylation by Syk at Tyr783 activates the enzymatic activity of PLC γ 1 (4). PLC γ 2 is engaged in antigen-dependent signaling in B cells and collagen-dependent signaling in platelets. Phosphorylation by Btk or Lck at Tyr753, 759, 1197, and 1217 is correlated with PLC γ 2 activity (5,6).						
Background Re	eferences	1. Singer, W.D. et al. (1997) <i>Annu Rev Biochem</i> 66, 475-509. 2. Margolis, B. et al. (1989) <i>Cell</i> 57, 1101-7. 3. Kim, H.K. et al. (1991) <i>Cell</i> 65, 435-41. 4. Wang, Z. et al. (1998) <i>Mol Cell Biol</i> 18, 590-7. 5. Watanabe, D. et al. (2001) <i>J Biol Chem</i> 276, 38595-601. 6. Ozdener, F. et al. (2002) <i>Mol Pharmacol</i> 62, 672-9.						
Species Reactiv	vity	Species reactivity is determined by testing in at least one approved application (e.g., western blot).				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 W-S: Simple Western™ IP: Immunoprecipitation IHC-P: Immunohistochemistry (Paraffin)						
Cross-Reactivit	ту Кеу	H: Human M: Mouse R: Rat Mk: Monkey						
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