## RIP3 (D4G2A) Rabbit mAb (PE Conjugate)



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Applications: FC-FP	<b>Reactivity:</b> M	<b>Sensitivity:</b> Endogenous	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #Q9QZL0	Entrez-Gene Id: 56532		
Product Usage Information		<b>Application</b> Flow Cytometry (Fixed/Permeabilized)		Dilution 1:50			
Storage		Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 mg/ml BSA. Store at 4°C. Do not aliquot antibody. Protect from light. Do not freeze.					
Specificity/Sensitivity RIP3 (D4G2A) Rabbit mAb (PE Conjugate) recognizes endogenous levels of total RIP3 promouse.					s of total RIP3 protein from		
Source / Purifica	tion	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val370 of mouse RIP3 protein.					
Description		This Cell Signaling Technology antibody is conjugated to phycoerythrin (PE) and tested in-house for direct flow cytometry and CellSimple™ cell-assay analysis in mouse cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated RIP3 (D4G2A) Rabbit mAb #95702.					
Background		The receptor-interacting protein (RIP) family of serine-threonine kinases (RIP, RIP2, RIP3, and RIP4) are important regulators of cellular stress that trigger pro-survival and inflammatory responses through the activation of NF-kB, as well as pro-apoptotic pathways (1). In addition to the kinase domain, RIP contains a death domain responsible for interaction with the death domain receptor Fas and recruitment to TNF-R1 through interaction with TRADD (2,3). RIP-deficient cells show a failure in TNF- mediated NF-kB activation, making the cells more sensitive to apoptosis (4,5). RIP also interacts with TNF-receptor-associated factors (TRAFs) and can recruit IKKs to the TNF-R1 signaling complex via interaction with NEMO, leading to IkB phosphorylation and degradation (6,7). Overexpression of RIP induces both NF-kB activation and apoptosis (2,3). Caspase-8-dependent cleavage of the RIP death domain can trigger the apoptotic activity of RIP (8).					
Background Ref	erences	<ol> <li>Meylan, E. and Tschopp, J. (2005) <i>Trends Biochem Sci</i> 30, 151-9.</li> <li>Hsu, H. et al. (1996) <i>Immunity</i> 4, 387-96.</li> <li>Stanger, B.Z. et al. (1995) <i>Cell</i> 81, 513-23.</li> <li>Ting, A.T. et al. (1996) <i>EMBO J</i> 15, 6189-96.</li> <li>Kelliher, M.A. et al. (1998) <i>Immunity</i> 8, 297-303.</li> <li>Devin, A. et al. (2000) <i>Immunity</i> 12, 419-29.</li> <li>Zhang, S.Q. et al. (2000) <i>Immunity</i> 12, 301-11.</li> <li>Lin, Y. et al. (1999) <i>Genes Dev</i> 13, 2514-26.</li> </ol>					
Species Reactivi	ty	Species reactivity is determined by testing in at least one approved application (e.g., western blot).					
Applications Key	/	FC-FP: Flow Cytometry (Fixed/Permeabilized)					
Cross-Reactivity	Кеу	M: Mouse					
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