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CDK9 (C12F7) Rabbit mAb (PE Conjugate)



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Applications: FC-FP	Reactivity: H M R Hm Mk B Dg	Sensitivity: Endogenous	Source/Isotype: Rabbit	UniProt ID: #P50750	Entrez-Gene Id: 1025		
Product Usage Information		Application Flow Cytometry (Fixed/Po	ermeabilized)		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 the antibody. Protect from light. Do not freeze.					
Specificity/Sensitivity		CDK9 (C12F7) Rabbit mAb (PE Conjugate) detects endogenous levels of total CDK9 protein, both 42 kDa and 55 kDa isoforms.					
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human CDK9.					
Description		This Cell Signaling Technology antibody is conjugated to phycoerythrin (PE) and tested in-house for direct flow cytometry analysis in human cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated CDK9 (C12F7) Rabbit mAb #2316.					
Background		P-TEFb is a general transcription factor that regulates transcription elongation through phosphorylation of the C-terminal tail domain (CTD) of RNA polymerase II (RNAP II). The P-TEFb complex is composed of a catalytic subunit, CDK9, and its regulatory cyclin partner, which can be cyclin T1, T2a, T2b or K (reviewed in 1,2). P-TEFb is recruited by the HIV Tat protein to allow transcriptional elongation, and subsequent replication of the viral genome. Inhibition of P-TEFb function therefore has potential for HIV therapy. CDK9 exists as two isoforms, an abundant 42 kDa isoform, and a less abundant 55 kDa isoform, which contains an amino-terminal extension (3). The two forms likely have distinct purposes based on differential expression during lymphocyte activation (4,5) and on their localization within the nucleus (5). Cyclin dependent kinases (CDKs) are activated in part by cyclin binding and by phosphorylation of a conserved threonine in the T-loop domain. Phosphorylation of CDK9 at the T-loop Thr186 by an unidentified nuclear kinase may be important in P-TEFb activation (6) and regulation of HIV transcription (7). Acetylation of CDK9 at Lys44 affects its ability to phosphorylate the RNAPII CTD (8).					
Background References		 Rice, A.P. and Herrmann, C.H. (2003) <i>Curr HIV Res</i> 1, 395-404. De Falco, G. and Giordano, A. (2002) <i>Cancer Biol Ther</i> 1, 342-7. Shore, S.M. et al. (2003) <i>Gene</i> 307, 175-82. Shore, S.M. et al. (2005) <i>Gene</i> 350, 51-8. Liu, H. and Herrmann, C.H. (2005) <i>J Cell Physiol</i> 203, 251-60. Chen, R. et al. (2004) <i>J Biol Chem</i> 279, 4153-60. Chen, R. et al. (2005) <i>Retrovirology</i> 2, 47. Fu, J. et al. (2007) <i>Mol Cell Biol</i> 27, 4641-51. De Falco, G. and Giordano, A. <i>Cancer Biol Ther</i> 1, 342-7. 					
Species Reacti	vity	Species reactivity is deter	rmined by testing in at lea	ast one approved ap	plication (e.g., western blot).		
Applications Key FC-FP: Flow Cytome		-C-FP: Flow Cytometry (F	/ (Fixed/Permeabilized)				
Cross-Reactivity Key H: Human M: Mouse R: Rat Hm: Hamster Mk: Monkey B: Bovine Dg: Dog				oog			
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