

20750

Cleaved Caspase-9 (Asp315) (D8I9E) Rabbit



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Applications:ReactW, IP, IF-IC, FC-FPH			Source/Isotype: Rabbit IgG	UniProt ID: #P55211	Entrez-Gene Id: 842	
Product Usage	Application	Application			Dilution	
Information	Western Blotting	Western Blotting			1:1000	
	Immunoprecipit	ation		1:50		
	Immunofluoresc	Immunofluorescence (Immunocytochemistry)			1:400 - 1:1600	
	Flow Cytometry	Flow Cytometry (Fixed/Permeabilized)			1:100 - 1:400	
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.				
	For a carrier free	(BSA and azide free) vei	rsion of this product see	product #24731.		
Specificity/Sensitivity	only when cleave	Cleaved-Caspase-9 (Asp315) (D8I9E) Rabbit mAb recognizes endogenous levels of caspase-9 protein only when cleaved at Asp315. Non-specific proteins that are induced by apoptosis under certain conditions may be detected.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Asp315 of human caspase-9 protein.				
Background	family (1,2). Upor 47 kDa procaspa processing, resu Asp330, producir caspase-9 furthe	Caspase-9 (ICE-LAP6, Mch6) is an important member of the cysteine aspartic acid protease (caspase) family (1,2). Upon apoptotic stimulation, cytochrome c released from mitochondria associates with the 47 kDa procaspase-9/Apaf-1. Apaf-1 mediated activation of caspase-9 involves intrinsic proteolytic processing, resulting in cleavage at Asp315 and producing a p35 subunit. Another cleavage occurs at Asp330, producing a p37 subunit that can serve to amplify the apoptotic response (3-6). Cleaved caspase-9 further processes other caspase members, including caspase-3 and caspase-7, to initiate a caspase cascade, which leads to apoptosis (7-10).				
Background Reference	2. Srinivasula, S. 3. Liu, X. et al. (19 4. Li, P. et al. (199 5. Zou, H. et al. (1 6. Srinivasula, S.I 7. Deveraux, Q. L 8. Slee, E. A. et al 9. Sun, X.M. et al	 Duan, H. et al. (1996) J. Biol. Chem. 271, 16720-16724. Srinivasula, S. M. et al. (1996) J. Biol. Chem. 271, 27099-27106. Liu, X. et al. (1996) Cell 86, 147-157. Li, P. et al. (1997) Cell 91, 479-489. Zou, H. et al. (1999) J. Biol. Chem. 274, 11549-11556. Srinivasula, S.M. et al. (1998) Mol Cell 1, 949-57. Deveraux, Q. L. et al. (1998) EMBO J. 17, 2215-2223. Slee, E. A. et al. (1999) J. Cell Biol. 144, 281-292. Sun, X.M. et al. (1999) J Biol Chem 274, 5053-60. MacFarlane, M. et al. (1997) J. Cell Biol. 137, 469-479. 				
Species Reactivity	Species reactivity	is determined by testin	g in at least one approve	ed application (e.g.,	western blot).	
		IMPORTANT: For western blots, incubate membrane with diluted				

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X

Applications Key W: Western Blotting IP: Immunoprecipitation IF-IC: Immunofluorescence (Immunocytochemistry) FC-

FP: Flow Cytometry (Fixed/Permeabilized)

Cross-Reactivity Key H: Human

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