evision	evision 5						
e at -20C	Caspase-9 Antibody	SE .	Cell Signaling TECHNOLOGY®				
Store		Orde		877-616-CELL (2355) ers@cellsignal.com			
		Supp	ort: 8	77-678-TECH (8324)			
#9502		Web: 3 Trask Lane Danvers		fo@cellsignal.com cellsignal.com setts 01923 USA			
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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W, W-S	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 35, 37, 47	Source/Isotype: Rabbit	UniProt ID: #P55211	Entrez-Gene Id 842		
Product Usage Information		Application Western Blotting Simple Western™			Dilution 1:1000 1:10 - 1:50			
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.						
Specificity/Sen	sitivity	Caspase-9 Antibody detects endogenous levels of full length caspase-9 (47 kDa) and large fragments of caspase-9. The antibody does not recognize other caspases.						
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding aspartic acid 315 of human caspase-9. Antibodies are purified by protein A and peptide affinity chromatography.						
Background		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 Ro	eferences	1. Duan, H. et al. (199 2. Srinivasula, S. M. et 3. Liu, X. et al. (1996) (4. Li, P. et al. (1997) <i>Ce</i> 5. Zou, H. et al. (1999) 6. Srinivasula, S.M. et 7. Deveraux, Q. L. et a 8. Slee, E. A. et al. (199 9. Sun, X.M. et al. (199 10. MacFarlane, M. et	al. (1996) <i>J. Biol. Ch</i> Cell 86, 147-157. ell 91, 479-489. <i>J. Biol. Chem.</i> 274, al. (1998) <i>Mol Cell</i> 1 I. (1998) <i>EMBO J.</i> 17 99) <i>J. Cell Biol.</i> 144, 2 99) <i>J. Biol Chem</i> 274,	nem. 271, 27099-27106. 11549-11556. , 949-57. , 2215-2223. 281-292. 5053-60.				
Species Reacti	vity	Species reactivity is de	etermined by testin	g in at least one approv	ed application (e.g.,	western blot).		
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.						
Applications K	ey	W: Western Blotting W-S: Simple Western™						
Cross-Reactivi	ty Key	H: Human						
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