Revision 5		
Cleaved Caspase-9 (Asp315) Antibody	C C	ell Signaling
Store	Orders:	877-616-CELL (2355) orders@cellsignal.com
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Applications: W, IP	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 35	<b>Source/Isotype:</b> Rabbit	<b>UniProt ID:</b> #P55211	Entrez-Gene Id 842		
Product Usage Information	2	<b>Application</b> Western Blotting Immunoprecipitatior	1		<b>Dilution</b> 1:1000 1:100			
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/Sensitivity		Cleaved Caspase-9 (Asp315) Antibody detects endogenous levels of the 35 kDa large fragment of caspase-9 following cleavage at aspartic acid 315. The antibody does not react with full length caspase-9 or any other cleaved caspases.						
Source / Purifi	/ Purification Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding residues surrounding Asp315 of human caspase-9. Antibodies are purified by protein A and peptid 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 R	eferences	<ol> <li>Duan, H. et al. (1996) <i>J. Biol. Chem.</i> 271, 16720-16724.</li> <li>Srinivasula, S. M. et al. (1996) <i>J. Biol. Chem.</i> 271, 27099-27106.</li> <li>Liu, X. et al. (1997) <i>Cell</i> 96, 147-157.</li> <li>Li, P. et al. (1997) <i>Cell</i> 91, 479-489.</li> <li>Zou, H. et al. (1997) <i>J. Biol. Chem.</i> 274, 11549-11556.</li> <li>Srinivasula, S.M. et al. (1998) <i>Mol Cell</i> 1, 949-57.</li> <li>Deveraux, Q. L. et al. (1998) <i>EMBO J.</i> 17, 2215-2223.</li> <li>Slee, E. A. et al. (1999) <i>J. Biol Chem.</i> 274, 5053-60.</li> <li>MacFarlane, M. et al. (1997) <i>J. Cell Biol.</i> 137, 469-479.</li> </ol>						
Species Reacti	vity	Species reactivity is d	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).		
Western Blot E	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 IP: Immunoprecipitation						
Cross-Reactivi	ty Key	H: Human						
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