## **Cleaved Caspase-9 (Asp353) Antibody**



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## For Research Use Only, Not for Use in Diagnostic Procedures

Applications: WB, IF-IC	Reactivity: M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 17, 38	Source: Rabbit	<b>UniProt ID:</b> #Q9JHK1	Entrez-Gene Id: 58918	
Product Usage Information	Ар	plication				Dilution	
	We	stern Blotting				1:1000	
	Imr	nunofluorescence (	Immunocytochemis	stry)		1:200	
Storage	•	Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.					
Specificity / Sensit	with	Cleaved Caspase-9 (Asp353) Antibody detects endogenous levels of the large fragment (17 kDa or 38 kDa with prodomain) of caspase-9 resulting from cleavage at aspartic acid 353. The antibody does not recognize full length caspase-9 or any other cleaved caspases.					
Source / Purification	amii	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to amino-terminal residues surrounding Asp353 of rat caspase-9. Antibodies are purified by protein A and peptide affinity chromatography.					
Background	fami kDa resu proc proc	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 Refere	2. Si 3. Li 4. Li 5. Zi 6. Si 7. D 8. Si	<ol> <li>Duan, H. et al. (1996) J. Biol. Chem. 271, 16720-16724.</li> <li>Srinivasula, S. M. et al. (1996) J. Biol. Chem. 271, 27099-27106.</li> <li>Liu, X. et al. (1996) Cell 86, 147-157.</li> <li>Li, P. et al. (1997) Cell 91, 479-489.</li> <li>Zou, H. et al. (1999) J. Biol. Chem. 274, 11549-11556.</li> <li>Srinivasula, S.M. et al. (1998) Mol Cell 1, 949-57.</li> <li>Deveraux, Q. L. et al. (1998) EMBO J. 17, 2215-2223.</li> <li>Slee, E. A. et al. (1999) J. Cell Biol. 144, 281-292.</li> <li>Sun, X.M. et al. (1999) J Biol Chem 274, 5053-60.</li> </ol>					

**Species Reactivity** 

Species reactivity is determined by testing in at least one approved 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 Key** 

WB: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)

**Cross-Reactivity Key** 

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster

X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

10. MacFarlane, M. et al. (1997) J. Cell Biol. 137, 469-479.

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**Limited Uses** 

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