Revision 1

Store at -20C

#9507

Cleaved Caspase-9 (Asp353) Antibody

Cell Signaling TECHNOLOGY®

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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
W, IF-IC	M R	Endogenous	17, 38	Rabbit	#Q9JHK1	58918

Product Usage Information

Application	Dilution
Western Blotting	1:1000
Immunofluorescence (Immunocytochemistry)	1:200

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 (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

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

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 References

1. Duan, H. et al. (1996) *J. Biol. Chem.* 271, 16720-16724.
2. Srinivasula, S. M. et al. (1996) *J. Biol. Chem.* 271, 27099-27106.
3. Liu, X. et al. (1996) *Cell* 86, 147-157.
4. Li, P. et al. (1997) *Cell* 91, 479-489.
5. Zou, H. et al. (1999) *J. Biol. Chem.* 274, 11549-11556.
6. Srinivasula, S.M. et al. (1998) *Mol Cell* 1, 949-57.
7. Deveraux, Q. L. et al. (1998) *EMBO J.* 17, 2215-2223.
8. Slee, E. A. et al. (1999) *J. Cell Biol.* 144, 281-292.
9. Sun, X.M. et al. (1999) *J Biol Chem* 274, 5053-60.
10. MacFarlane, M. et al. (1997) *J. Cell Biol.* 137, 469-479.

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

W: Western Blotting **IF-IC:** Immunofluorescence (Immunocytochemistry)

Cross-Reactivity Key

M: Mouse **R:** Rat

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Limited Uses

force or effect.

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