

Store at
-20C
#52873**Cleaved Caspase-9 (Asp330) (E5Z7N)
Rabbit mAb****Orders:** 877-616-CELL (2355)
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Applications: W, IF-IC	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 37	Source/Isotype: Rabbit IgG	UniProt ID: #P55211	Entrez-Gene Id: 842
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**Product Usage
Information****Application**Western Blotting
Immunofluorescence (Immunocytochemistry)**Dilution**1:1000
1:50**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.

Specificity/Sensitivity

Cleaved Caspase-9 (Asp330) (E5Z7N) Rabbit mAb recognizes endogenous levels of caspase-9 protein only when cleaved at Asp330.

Source / Purification

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Asp330 of human caspase-9 protein.

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

- 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 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****H:** Human**Trademarks and Patents**

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