

DESCRIPTION

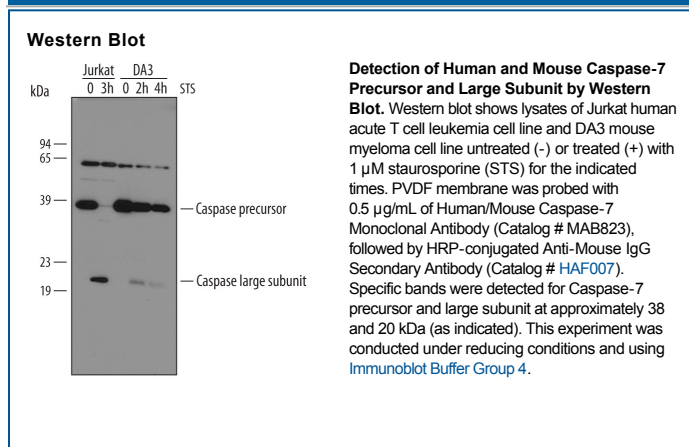
Species Reactivity	Human/Mouse
Specificity	Detects human and mouse precursor Caspase-7 and the large subunit of cleaved Caspase-7.
Source	Monoclonal Mouse IgG ₁ Clone # MCH3101.62
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	<i>E. coli</i> -derived recombinant human Caspase-7 Accession # P55210
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied as a 0.2 µm filtered solution in PBS.

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. *General Protocols* are available in the *Technical Information* section on our website.

	Recommended Concentration	Sample
Western Blot	0.5 µg/mL	See Below

DATA



PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.5 mg/mL in sterile PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. <ul style="list-style-type: none"> ● 12 months from date of receipt, -20 to -70 °C as supplied. ● 1 month, 2 to 8 °C under sterile conditions after reconstitution. ● 6 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

Caspase-7 (Cysteine-aspartic acid protease 7/Casp7; also CMH-1, ICE-LAP3 and Mch3) is a 32 kDa member of the peptidase C14A/IL-1 β -converting family of enzymes (1, 2, 3). It is widely expressed, except in brain, and is best known as an integral component of the apoptotic cascade. Caspase-7 is considered to be an executioner caspase, as a downstream mediator of apoptotic-associated proteolysis (2, 3). Upon activation, Caspase-7 is known to utilize a Cys residue to cleave multiple substrates, including PARP, procaspase 6, Gas2 and calpstatin (1). Human procaspase-7 is a 34 - 36 kDa, 303 amino acid (aa) protein (4, 5, 6). Normally, it is an inactive homodimer (1, 2, 7, 8). But following an upstream signal that activates processing proteases, procaspase-7 undergoes proteolytic cleavage to generate an N-terminal 23 aa propeptide, a 175 aa p20/20 kDa subunit (aa 24 - 198), and a 105 aa C-terminal p12/12 kDa subunit (5). The p20 and p12 subunits noncovalently heterodimerize, and subsequently associate with another p20/p12 heterodimer to form an active antiparallel homodimer. Additional processing of p20 may remove aa 24 - 36 to generate p18, while additional processing of p12 will remove aa 199 - 206 to generate p11 (9, 10). Multiple proteases can use Caspase-7 as a substrate, and include caspase-1, -3, -8, and -10, granzyme B, calpain-1 and Caspase-7 itself (3, 6, 9, 11). Caspase-7 is found in both cytosol and nucleus, and possesses a potential KKKK nuclear localization signal between aa 38 - 41 that likely undergoes sumoylation (9, 12). There are two potential isoform variants, one which shows an alternate start site 33 aa upstream of the standard start site, and a second that shows a 105 aa substitution for aa 149 - 303. Human and mouse Caspase-7 are 82% aa identical at the amino acid level.

References:

1. Chowdhury, I. *et al.* (2008) *Comp. Biochem. Physiol. B* **151**:10.
2. Boatright, K.M. and G.S. Salvesen (2003) *Curr. Opin. Cell Biol.* **15**:725.
3. Launay, S. *et al.* (2005) *Oncogene* **24**:5137.
4. Juan, T. *et al.* (1997) *Genomics* **40**:86.
5. Fernandez-Alnemri, T. *et al.* (1995) *Cancer Res.* **55**:6045.
6. Fernandez-Alnemri, T. *et al.* (1996) *Proc. Natl. Acad. Sci. USA* **93**:7464.
7. Gao, Z. *et al.* (2007) *J. Biol. Chem.* **282**:30718.
8. Riedl, S.J. *et al.* (2001) *Proc. Natl. Acad. Sci. USA* **98**:14790.
9. Gafni, J. *et al.* (2009) *J. Biol. Chem.* July 21 [epub ahead of print].
10. Lippke, J.A. *et al.* (1996) *J. Biol. Chem.* **271**:1825.
11. Lamkanfi, M. *et al.* (2008) *Mol. Cell. Proteomics* **7**:2350.
12. Hayashi, N. *et al.* (2006) *Neurosci. Lett.* **397**:5.