Cleaved Caspase-6 (Asp162) Antibody





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

Applications: W	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 18	Source/Isotype: Rabbit	UniProt ID: #P55212	Entrez-Gene Id: 839		
Product Usage Information	9	Application Western Blotting			Dilution 1:1000			
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 aliguot the antibody.						
Specificity/Ser	nsitivity	Cleaved Caspase-6 (Asp162) Antibody detects endogenous levels of the large subunit of active caspase- 6 resulting from cleavage at Asp179 in human or Asp162 in mouse and rat (18 kDa). This antibody does not recognize other cleaved caspases.						
Source / Purifi	ication	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to the carboxy-terminal sequence of p18, the large subunit of rat caspase-6. Antibodies are purified by protein A and peptide affinity chromatography.						
Background		Caspase-6 (Mch2) is one of the major executioner caspases functioning in cellular apoptotic processes (1,2). Upon apoptotic stimulation, initiator caspases such as caspase-9 are cleaved and activated (3). The activated upstream caspases further process downstream executioner caspases, such as caspase-3 and caspase-6, by cleaving them into large and small subunits, thereby initiating a caspase cascade leading to apoptosis (4,5). One of the major targets for caspase-6 is the membrane associated protein lamin A (6). The cleavage of this protein causes cell membrane malfunction, membrane blebbing, and eventual cell death.						
Background R	eferences	1. Cohen, G.M. (1997) <i>Biochem J</i> 326 (Pt 1), 1-16. 2. Faleiro, L. et al. (1997) <i>EMBO J</i> 16, 2271-81. 3. Li, P. et al. (1997) <i>Cell</i> 91, 479-89. 4. Slee, E.A. et al. (1999) <i>J Cell Biol</i> 144, 281-92. 5. MacFarlane, M. et al. (1997) <i>J Cell Biol</i> 137, 469-79. 6. Orth, K. et al. (1996) <i>J Biol Chem</i> 271, 16443-6.						
Species Reacti	ivity	Species reactivity is determined by testing in at least one approved application (e.g., western blot).						
Western Blot I	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						
Cross-Reactivi	ty Key	H: Human M: Mouse R: Rat						
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