

Caspase-7 Antibody

Catalog # ASC10299

Specification

Caspase-7 Antibody - Product Information

Application
Primary Accession
Other Accession
Reactivity
Host
Clonality
Isotype

Host Rabbit
Clonality Polyclonal
Isotype IgG
Application Notes Casp-7 ant

Casp-7 antibody can be used for the detection of Caspase-7 by Western blot at 0.5 to 1 μ g/mL. Antibody can also be used for immunohistochemistry starting at 2 μ g/mL.

Caspase-7 Antibody - Additional Information

Gene ID 840

Other Names

Caspase-7 Antibody: MCH3, CMH-1, LICE2, CASP-7, ICE-LAP3, MCH3, Caspase-7, Apoptotic protease Mch-3, caspase 7, apoptosis-related cysteine peptidase

Target/Specificity

CASP7; Depending on cell lines or tissues used, other cleavage products may be observed.

Reconstitution & Storage

Caspase-7 antibody can be stored at 4°C for three months and -20°C, stable for up to one year. As with all antibodies care should be taken to avoid repeated freeze thaw cycles. Antibodies should not be exposed to prolonged high temperatures.

WB, IHC P55210

P55210, 1730092

Human, Mouse, Rat

Precautions

Caspase-7 Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

Caspase-7 Antibody - Protein Information

Name CASP7 {ECO:0000303|PubMed:9070923, ECO:0000312|HGNC:HGNC:1508}

Function

Thiol protease involved in different programmed cell death processes, such as apoptosis, pyroptosis or granzyme-mediated programmed cell death, by proteolytically cleaving target proteins (PubMed:8521391, PubMed:8567622, PubMed:8576161, PubMed:<a href="http://www.uniprot.org/citations/9070923"



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target=" blank">9070923</a>, PubMed:<a href="http://www.uniprot.org/citations/16916640"
target="blank">16916640</a>, PubMed:<a href="http://www.uniprot.org/citations/17646170"
target="blank">17646170</a>, PubMed:<a href="http://www.uniprot.org/citations/18723680"
target="_blank">18723680</a>, PubMed:<a href="http://www.uniprot.org/citations/19581639"
target=" blank">19581639</a>, PubMed:<a href="http://www.uniprot.org/citations/11257230"
target=" blank">11257230</a>, PubMed:<a href="http://www.uniprot.org/citations/11257231"
target=" blank">11257231</a>, PubMed:<a href="http://www.uniprot.org/citations/11701129"
target="blank">11701129</a>, PubMed:<a href="http://www.uniprot.org/citations/15314233"
target="blank">15314233</a>). Has a marked preference for Asp-Glu-Val-Asp (DEVD)
consensus sequences, with some plasticity for alternate non-canonical sequences (PubMed: <a
href="http://www.uniprot.org/citations/12824163" target=" blank">12824163</a>, PubMed:<a
href="http://www.uniprot.org/citations/19581639" target="blank">19581639</a>, PubMed:<a
href="http://www.uniprot.org/citations/20566630" target="blank">20566630</a>, PubMed:<a
href="http://www.uniprot.org/citations/15314233" target=" blank">15314233</a>, PubMed:<a
href="http://www.uniprot.org/citations/17697120" target="blank">17697120</a>, PubMed:<a
href="http://www.uniprot.org/citations/23897474" target="blank">23897474</a>, PubMed:<a
href="http://www.uniprot.org/citations/23650375" target="_blank">23650375</a>, PubMed:<a
href="http://www.uniprot.org/citations/27032039" target="blank">27032039</a>). Its
involvement in the different programmed cell death processes is probably determined by
upstream proteases that activate CASP7 (By similarity). Acts as an effector caspase involved in the
execution phase of apoptosis: following cleavage and activation by initiator caspases (CASP8,
CASP9 and/or CASP10), mediates execution of apoptosis by catalyzing cleavage of proteins, such
as CLSPN, PARP1, PTGES3 and YY1 (PubMed: <a href="http://www.uniprot.org/citations/10497198"
target=" blank">10497198</a>, PubMed:<a href="http://www.uniprot.org/citations/16123041"
target="blank">16123041</a>, PubMed:<a href="http://www.uniprot.org/citations/16374543"
target="blank">16374543</a>, PubMed:<a href="http://www.uniprot.org/citations/16916640"
target="blank">16916640</a>, PubMed:<a href="http://www.uniprot.org/citations/18723680"
target=" blank">18723680</a>, PubMed:<a href="http://www.uniprot.org/citations/20566630"
target=" blank">20566630</a>, PubMed:<a href="http://www.uniprot.org/citations/21555521"
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target="blank">28863261</a>, PubMed:<a href="http://www.uniprot.org/citations/31586028"
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target="blank">34156061</a>, PubMed:<a href="http://www.uniprot.org/citations/27889207"
target="blank">27889207</a>, PubMed:<a href="http://www.uniprot.org/citations/35338844"
target="blank">35338844</a>, PubMed:<a href="http://www.uniprot.org/citations/35446120"
target="blank">35446120</a>). Compared to CASP3, acts as a minor executioner caspase and
cleaves a limited set of target proteins (PubMed:<a
href="http://www.uniprot.org/citations/18723680" target="_blank">18723680</a>). Acts as a key
regulator of the inflammatory response in response to bacterial infection by catalyzing cleavage
and activation of the sphingomyelin phosphodiesterase SMPD1 in the extracellular milieu, thereby
promoting membrane repair (PubMed: <a href="http://www.uniprot.org/citations/21157428"
target=" blank">21157428</a>). Regulates pyroptosis in intestinal epithelial cells: cleaved and
activated by CASP1 in response to S.typhimurium infection, promoting its secretion to the
extracellular milieu, where it catalyzes activation of SMPD1, generating ceramides that repair
membranes and counteract the action of gasdermin-D (GSDMD) pores (By similarity). Regulates
granzyme-mediated programmed cell death in hepatocytes: cleaved and activated by granzyme B
(GZMB) in response to bacterial infection, promoting its secretion to the extracellular milieu, where
it catalyzes activation of SMPD1, generating ceramides that repair membranes and counteract the
action of perforin (PRF1) pores (By similarity). Following cleavage by CASP1 in response to
inflammasome activation, catalyzes processing and inactivation of PARP1, alleviating the
transcription repressor activity of PARP1 (PubMed:<a
href="http://www.uniprot.org/citations/22464733" target=" blank">22464733</a>). Acts as an
inhibitor of type I interferon production during virus-induced apoptosis by mediating cleavage of
antiviral proteins CGAS, IRF3 and MAVS, thereby preventing cytokine overproduction (By
similarity). Cleaves and activates sterol regulatory element binding proteins (SREBPs) (PubMed: <a
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href="http://www.uniprot.org/citations/8643593" target=" blank">8643593). Cleaves phospholipid scramblase proteins XKR4, XKR8 and XKR9 (By similarity). In case of infection, catalyzes cleavage of Kaposi sarcoma-associated herpesvirus protein ORF57, thereby preventing expression of viral lytic genes (PubMed: 20159985).

Cellular Location

Cytoplasm, cytosol. Nucleus. Secreted, extracellular space {ECO:0000250|UniProtKB:P97864}. Note=Following cleavage and activation by CASP1 or granzyme B (GZMB), secreted into the extracellular milieu by passing through the gasdermin-D (GSDMD) pores or perforin (PRF1) pore, respectively {ECO:0000250|UniProtKB:P97864}

Tissue Location

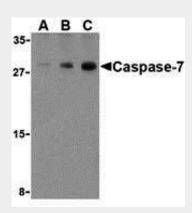
Highly expressed in lung, skeletal muscle, liver, kidney, spleen and heart, and moderately in testis. No expression in the brain.

Caspase-7 Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

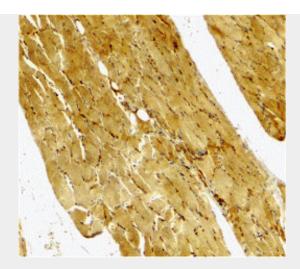
- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

Caspase-7 Antibody - Images



Western blot analysis of Caspase-7 in human skeletal muscle cell lysate with Caspase-7 antibody at (A) 0.5, (B) 1, and (C) 2 μg/mL.





Immunohistochemical staining of human skeletal muscle using Caspase-7 antibody at 2 μg/mL.

Caspase-7 Antibody - Background

Caspase-7 Antibody: Caspases are a family of cysteine proteases that can be divided into the apoptotic and inflammatory caspase subfamilies. Unlike the apoptotic caspases, members of the inflammatory subfamily are generally not involved in cell death but are associated with the immune response to microbial pathogens. The apoptotic subfamily can be further divided into initiator caspases, which are activated in response to death signals, and executioner caspases, which are activated by the initiator caspases and are responsible for cleavage of cellular substrates that ultimately lead to cell death. Caspase-7 is an executioner caspase that was identified based on its homology with caspases 1 and 3, as well as the C. elegans cell death protein CED-3. Alternative splicing of Caspase-7 mRNA results in the production of 3 distinct isoforms. Caspase-7 activity can be directly inhibited by XIAP expression.

Caspase-7 Antibody - References

Martinon F and Tschopp J. Inflammatory caspases: linking an intracellular innate immune system to autoinflammatory diseases. Cell 2004; 117:561-74.

Zhivotovsky B and Orrenius S. Caspase-2 function in response to DNA damage. Biochim. Biophys. Res. Comm. 2005; 331:859-67.

Wolf BB and Green DR. Suicidal tendencies: apoptotic cell death by caspase family proteinases. J. Biol. Chem. 1999; 274:20049-52.

Juan TSC, McNiece IK, Argento JM, et al. Identification and mapping of Casp7, a cysteine protease resembling CPP32b, Interleukin-1b converting enzyme, and CED-3. Genomics 1997; 40:86-93.