10320 Camino Santa Fe, Suite G
San Diego, CA 92121
Tel: 858.875.1900 Fax: 858.875.1999

Phospho-mouse CASP3(S26) Antibody Blocking peptide

Synthetic peptide
Catalog # BP3778d

Specification

Phospho-mouse CASP3(S26) Antibody Blocking peptide - Product Information

Primary Accession P70677

Phospho-mouse CASP3(S26) Antibody Blocking peptide - Additional Information

Gene ID 12367

Other Names
Caspase-3, CASP-3, Apopain, Cysteine protease CPP32, CPP-32, LICE, Protein Yama, SREBP cleavage activity 1, SCA-1, Caspase-3 subunit p17, Caspase-3 subunit p12, Casp3, Cpp32

Format
Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

Storage
Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

Precautions
This product is for research use only. Not for use in diagnostic or therapeutic procedures.

Phospho-mouse CASP3(S26) Antibody Blocking peptide - Protein Information

Name Casp3

Synonyms Cpp32 {ECO:0000303|PubMed:8934524}

Function
Thiol protease that acts as a major effector caspase involved in the execution phase of apoptosis (PubMed:8934524, PubMed:16469926). Following cleavage and activation by initiator caspases (CASP8, CASP9 and/or CASP10), mediates execution of apoptosis by catalyzing cleavage of many proteins (PubMed:8934524, PubMed:16469926). At the onset of apoptosis, it proteolytically cleaves poly(ADP-ribose) polymerase PARP1 at a '216-Asp-|-Gly-217' bond. Cleaves and activates sterol regulatory element binding proteins (SREBPs) between the basic helix-loop-helix leucine zipper domain and the membrane attachment domain. Cleaves and activates caspase-6, -7 and -9. Triggers cell adhesion in sympathetic neurons through RET cleavage (By similarity). Cleaves IL-1 beta between an Asp and an Ala, releasing the mature cytokine which is involved in a variety of inflammatory processes (By similarity). Cleaves and inhibits serine/threonine-protein kinase AKT1 in response to oxidative stress (PubMed:12124386). 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 (PubMed:30878284). Also involved in pyroptosis by mediating cleavage and activation of gasdermin-E (GSDME) (By similarity). Cleaves XRCC4 and phospholipid scramblase proteins XKR4, XKR8 and XKR9, leading to promote phosphatidylserine exposure on apoptotic cell surface (PubMed:25231987, PubMed:33725486).

Cellular Location
Cytoplasm {ECO:0000250|UniProtKB:P42574}.

Tissue Location
Highest expression in spleen, lung, liver, kidney and heart (PubMed:9038361). Lower expression in brain, skeletal muscle and testis (PubMed:9038361).

Phospho-mouse CASP3(S26) Antibody Blocking peptide - Protocols

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

- Blocking Peptides

Phospho-mouse CASP3(S26) Antibody Blocking peptide - Images

Phospho-mouse CASP3(S26) Antibody Blocking peptide - Background

This gene encodes a protein which is a member of thecysteine-aspartic acid protease (caspase) family. Sequentialactivation of caspases plays a central role in the execution-phaseof cell apoptosis. Caspases exist as inactive proenzymes whichundergo proteolytic processing at conserved aspartic residues toproduce two subunits, large and small, that dimerize to form theactive enzyme. This protein cleaves and activates caspases 6, 7 and9, and the protein itself is processed by caspases 8, 9 and 10. Itis the predominant caspase involved in the cleavage of amyloid-beta4A precursor protein, which is associated with neuronal death inAlzheimer's disease. Alternative splicing of this gene results intwo transcript variants that encode the same protein. [provided byRefSeq].

Phospho-mouse CASP3(S26) Antibody Blocking peptide - References

Srikanth, C.V., et al. Science 330(6002):390-393(2010)Li, F., et al. Cell Stem Cell 7(4):508-520(2010)Wang, L., et al. J. Neurosci. 30(39):13201-13210(2010)Gascon, E., et al. J. Neurosci. 30(37):12414-12423(2010)Bohsali, A., et al. BMC Microbiol. 10, 237 (2010) :