

CASP3 Antibody

Purified Mouse Monoclonal Antibody Catalog # AO2019a

Product Information

Application Primary Accession Reactivity Host Clonality Clone Names Isotype Calculated MW Description	 WB, E P42574 Human, Mouse, Rat Mouse Monoclonal 3D4D10 IgG1 31608 This gene encodes a protein which is a member of the cysteine-aspartic acid protease (caspase) family. Sequential activation of caspases plays a central role in the execution-phase of cell apoptosis. Caspases exist as inactive proenzymes which undergo proteolytic processing at conserved aspartic residues to produce two subunits, large and small, that dimerize to form the active enzyme. This protein cleaves and activates caspases 6, 7 and 9, and the protein itself is processed by caspases 8, 9 and 10. It is the predominant caspase involved in the cleavage of amyloid-beta 4A precursor protein, which is associated with neuronal death in Alzheimer's disease. Alternative splicing of this gene results in two transcript variants that encode the same protein.
Immunogen	Purified recombinant fragment of human CASP3 (AA: 29-175) expressed in E. Coli.
Formulation	Purified antibody in PBS with 0.05% sodium azide

Additional Information

Gene ID	836
Other Names	Caspase-3, CASP-3, 3.4.22.56, Apopain, Cysteine protease CPP32, CPP-32, Protein Yama, SREBP cleavage activity 1, SCA-1, Caspase-3 subunit p17, Caspase-3 subunit p12, CASP3, CPP32
Dilution	WB~~1/500 - 1/2000 E~~1/10000
Storage	Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
Precautions	CASP3 Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

Protein Information

Name	CASP3
Synonyms	CPP32 {ECO:0000303 PubMed:7983002}
Function	Thiol protease that acts as a major effector caspase involved in the execution phase of apoptosis (PubMed: <u>18723680</u> , PubMed: <u>20566630</u> , PubMed: <u>23650375</u> , PubMed: <u>35338844</u> , PubMed: <u>35446120</u> , PubMed: <u>7596430</u>). Following cleavage and activation by initiator caspases (CASP8, CASP9 and/or CASP10), mediates execution of apoptosis by catalyzing cleavage of many proteins (PubMed: <u>18723680</u> , PubMed: <u>20566630</u> , PubMed: <u>23650375</u> , PubMed: <u>7596430</u>). At the onset of apoptosis, it proteolytically cleaves poly(ADP-ribose) polymerase PARP1 at a '216-Asp-]-Gly-217' bond (PubMed: <u>10497198</u> , PubMed: <u>16374543</u> , PubMed: <u>7596430</u> , PubMed: <u>7774019</u>). Cleaves and activates sterol regulatory element binding proteins (SREBPs) between the basic helix-loop-helix leucine zipper domain and the membrane attachment domain (By similarity). Cleaves and activates caspase-6, -7 and -9 (CASP6, CASP7 and CASP9, respectively) (PubMed: <u>37993714</u> , PubMed: <u>9334240</u>). Involved in the cleavage of huntingtin (PubMed: <u>8696339</u>). Triggers cell adhesion in sympathetic neurons through RET cleavage (PubMed: <u>21357690</u>). Cleaves and inhibits serine/threonine-protein kinase AKT1 in response to oxidative stress (PubMed: <u>23152800</u>). 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: <u>30878284</u>). Also involved in pyroptosis by mediating cleavage and activation of gasdermin-E (GSDME) (PubMed: <u>35338844</u> , PubMed: <u>35446120</u>). Cleaves XRCC4 and phospholipid scramblase proteins XKR4, XKR8 and XKR9, leading to promote phosphatidylserine exposure on apoptotic cell surface (PubMed: <u>23845944</u> , PubMed: <u>33725486</u>). Cleaves BIRC6 following inhibition of BIRC6-caspase binding by DIABLO/SMAC (PubMed: <u>36758104</u> , PubMed: <u>36758106</u>).
Cellular Location	Cytoplasm.
Tissue Location	Highly expressed in lung, spleen, heart, liver and kidney. Moderate levels in brain and skeletal muscle, and low in testis. Also found in many cell lines, highest expression in cells of the immune system.

References

Cell Death Dis. 2013 Jul 11;4:e725.PLoS One. 2013 May 2;8(5):e62303.

Images

