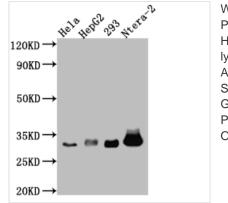
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CASP3 Recombinant Monoclonal Antibody

Product Code	CSB-RA286668A0HU
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Uniprot No.	P42574
Immunogen	A synthesized peptide derived from human pro Caspase 3
Species Reactivity	Human
Tested Applications	ELISA, WB, IHC, IF; Recommended dilution: WB:1:500-1:5000, IHC:1:50-1:200, IF:1:20-1:200
Relevance	Involved in the activation cascade of caspases responsible for apoptosis execution. At the onset of apoptosis it proteolytically cleaves poly(ADP-ribose) polymerase (PARP) 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. Involved in the cleavage of huntingtin. Triggers cell adhesion in sympathetic neurons through RET cleavage.
Form	Liquid
Form Conjugate	Liquid Non-conjugated
	•
Conjugate	Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium
Conjugate Storage Buffer	Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.
Conjugate Storage Buffer Purification Method	Non-conjugatedRabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.Affinity-chromatography
Conjugate Storage Buffer Purification Method Isotype	Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Affinity-chromatography Rabbit IgG
Conjugate Storage Buffer Purification Method Isotype Clonality	Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Affinity-chromatography Rabbit IgG Monoclonal
Conjugate Storage Buffer Purification Method Isotype Clonality Product Type	Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Affinity-chromatography Rabbit IgG Monoclonal Recombinant Antibody
Conjugate Storage Buffer Purification Method Isotype Clonality Product Type Immunogen Species	Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Affinity-chromatography Rabbit IgG Monoclonal Recombinant Antibody Homo sapiens (Human)
Conjugate Storage Buffer Purification Method Isotype Clonality Product Type Immunogen Species Research Area	Non-conjugatedRabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.Affinity-chromatographyRabbit IgGMonoclonalRecombinant AntibodyHomo sapiens (Human)Cancer; Cell biology; Metabolism

Image

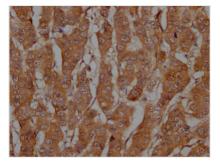


Western Blot Positive WB detected in: Hela whole cell lysate, HepG2 whole cell lysate, HEK293 whole cell lysate, Ntera-2 whole cell lysate All lanes: pro Caspase 3 antibody at 1:1000 Secondary Goat polyclonal to rabbit IgG at 1/50000 dilution Predicted band size: 32 kDa Observed band size: 32 kDa

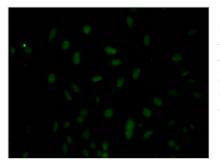
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IHC image of CSB-RA286668A0HU diluted at 1:100 and staining in paraffin-embedded human breast cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4? overnight. The primary is detected by a Goat anti-rabbit IgG polymer labeled by HRP and visualized using 0.05% DAB.



Immunofluorescence staining of Hela Cells with CSB-RA286668A0HU at 1:50, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeated by 0.2% TritonX-100, and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4?. Nuclear DNA was labeled in blue with DAPI. The secondary antibody was FITC-conjugated AffiniPure Goat Anti-Rabbit IgG (H+L).

Description

B cells were extracted from the animal immunized with a synthetic peptide derived from human CASP3, followed by fusion with myeloma cells to generate hybridomas. The variable light and variable heavy domains of CASP3 antibodyproducing hybridomas were sequenced to construct a vector for a recombinant generation. The CASP3 monoclonal antibody gene-containing vector was then transfected into cells for cultivation, and the CASP3 recombinant monoclonal antibody was isolated and purified from the cell culture supernatant using affinity chromatography. This antibody was tested for the detection of human CASP3 protein in ELISA, WB, IHC, and IF applications.

CASP3 is a protease enzyme that plays a crucial role in apoptosis. Once activated, CASP3 cleaves various cellular substrates, including structural proteins, signaling molecules, and DNA repair enzymes, resulting in the dismantling of the cell and the removal of apoptotic cells by phagocytosis. CASP3 has been shown to be involved in a wide range of physiological and pathological processes, including development, tissue homeostasis, and various diseases, such as cancer and neurodegeneration.