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anti-Caspase 8 antibody (N-Term)



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Publications



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Quantity:	100 μg	
Target:	Caspase 8 (CASP8)	
Binding Specificity:	AA 410-449, N-Term	
Reactivity:	Human, Mouse, Rat	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This Caspase 8 antibody is un-conjugated	
Application:	Western Blotting (WB), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p))	
Product Details		
Purpose:	Rabbit IgG polyclonal antibody for Caspase-8(CASP8) detection. Tested with WB, IHC-P in Human,Mouse,Rat.	
Immunogen:	A synthetic peptide corresponding to a sequence at the N-terminus of human Caspase 8 (410-449aa VSYRNPAEGTWYIQSLCQSLRERCPRGDDILTILTEVNYE), different from the related mouse and rat sequences by seven amino acids.	
Sequence:	VSYRNPAEGT WYIQSLCQSL RERCPRGDDI LTILTEVNYE	
Isotype:	IgG	
Cross-Reactivity (Details):	No cross reactivity with other proteins.	
Characteristics:	Rabbit IgG polyclonal antibody for Caspase-8(CASP8) detection. Tested with WB, IHC-P in Human,Mouse,Rat. Gene Name: caspase 8	

Product Details

	Protein Name: Caspase-8	
Purification:	Immunogen affinity purified.	
Target Details		
Target:	Caspase 8 (CASP8)	
Alternative Name:	CASP8 (CASP8 Products)	
Background:	CASP8 is also known as CAP4, MACH or MCH5. This gene encodes 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 composed of a prodomain, a large protease subunit, and a small protease subunit. Activation of caspases requires proteolytic processing at conserved internal aspartic residues to generate a heterodimeric enzyme consisting of the large and small subunits. This protein is involved in the programmed cell death induced by Fas and various apoptotic stimuli. The N-terminal FADD-like death effector domain of this protein suggests that it may interact with Fas-interacting protein FADD. In addition, this protein was detected in the insoluble fraction of the affected brain region from Huntington disease patients but not in those from normal controls, which implicated the role in neurodegenerative diseases. Many alternatively spliced transcript variants encoding different isoforms have been described, although not all variants have had their full-length sequences determined. Synonyms: Apoptotic protease Mch 5 Apoptotic protease Mch5 Apoptotic protease Mch-5 CAP4 CASP8 CASP8 CASP8 CASP8 Caspase8 Caspase	
	8(CASP-8) CED 3 FADD Like ICE FADD-like ICE FLICE MACH MCH 5 MCH5 Q14790	
Gene ID:	841	
UniProt:	Q14790	
Pathways:	Apoptosis, Caspase Cascade in Apoptosis, TLR Signaling, Activation of Innate immune Response, Tube Formation, Positive Regulation of Endopeptidase Activity, Toll-Like Receptors Cascades	
Application Details		
Application Notes:	WB: Concentration: 0.1-0.5 µg/mL, Tested Species: Human, Mouse, Rat IHC-P: Concentration: 0.5-1 µg/mL, Tested Species: Human, Mouse, Rat, Epitope Retrieval Heat: Boiling the paraffin sections in 10 mM citrate buffer, pH 6.0, for 20 mins is required to	

Application Details

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	staining of formalin/paraffin sections.	
	Notes: Tested Species: Species with positive results. Other applications have not been tested.	
	Optimal dilutions should be determined by end users.	
Comment:	Antibody can be supported by chemiluminescence kit ABIN921124 in WB, supported by	
	ABIN921231 in IHC(P).	
Restrictions:	For Research Use only	
Handling		
Format:	Lyophilized	
Reconstitution:	Add 0.2 mL of distilled water will yield a concentration of 500 µg/mL.	
Concentration:	500 μg/mL	
Buffer:	Each vial contains 5 mg BSA, 0.9 mg NaCl, 0.2 mg Na2HPO4, 0.05 mg Sodium azide.	
Preservative:	Sodium azide	
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which	
	should be handled by trained staff only.	
Handling Advice:	Avoid repeated freezing and thawing.	
Storage:	4 °C/-20 °C	
Storage Comment:	At -20°C for one year. After reconstitution, at 4°C for one month.	
	It can also be aliquotted and stored frozen at -20 °C for a longer time. Avoid repeated freezing	
	and thawing.	
Publications		
Product cited in:	Guo, Fu, Wang, Wang, Li, Huang, Gao, Li: "Di-2-pyridylhydrazone Dithiocarbamate Butyric Acid	
	Ester Exerted Its Proliferative Inhibition against Gastric Cell via ROS-Mediated Apoptosis and	
	Autophagy." in: Oxidative medicine and cellular longevity, Vol. 2018, pp. 4950705, (2018) (
	PubMed).	
	Zhan, Hu, Yi, An, Huang: "[Corrigendum] Inhibitory activity of apogossypol in human prostate	
	cancer in vitro and in vivo." in: Molecular medicine reports, Vol. 17, Issue 6, pp. 8010, (2018) (
	PubMed).	

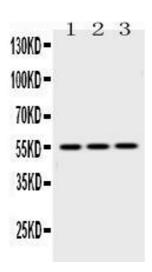
Zhou, He, Wang, Xie, Liu, Liu, Song, Ma: "Intravenous Administration Is an Effective and Safe Route for Cancer Gene Therapy Using the Bifidobacterium-Mediated Recombinant HSV-1 Thymidine Kinase and Ganciclovir." in: **International journal of molecular sciences**, Vol. 17, Issue 6, (2017) (PubMed).

Zhou, Hong, Yu, Nie, Gong, Xiong, Xie: "Exopolysaccharides from Lactobacillus plantarum NCU116 induce c-Jun dependent Fas/Fasl-mediated apoptosis via TLR2 in mouse intestinal epithelial cancer cells." in: **Scientific reports**, Vol. 7, Issue 1, pp. 14247, (2017) (PubMed).

Zhang, Huang, Wang, Luo, Wang: "2-DG-Regulated RIP and c-FLIP Effect on Liver Cancer Cell Apoptosis Induced by TRAIL." in: **Medical science monitor: international medical journal of experimental and clinical research**, Vol. 21, pp. 3442-8, (2016) (PubMed).

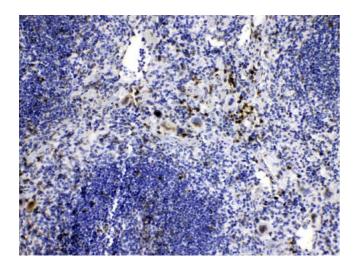
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Images



Western Blotting

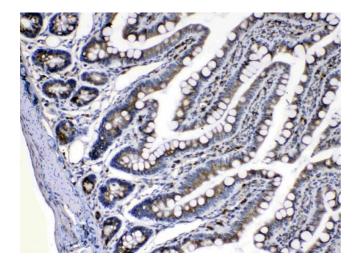
Image 1. Western blot analysis of Caspase8 using anti-Caspase8 antibody . Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel) / 90V (Resolving gel) for 2-3 hours. The sample well of each Lane was loaded with 50ug of sample under reducing conditions. Lane 1: rat liver tissue lysates, Lane 2: mouse liver tissue lysates, Lane 3: HEPG2 whole cell lysates. After Electrophoresis, proteins were transferred to a Nitrocellulose membrane at 150mA for 50-90 minutes. Blocked the membrane with 5% Non-fat Milk/ TBS for 1.5 hour at RT. The membrane was incubated with rabbit anti-Caspase8 antigen affinity purified polyclonal antibody (Catalog #) at 0.5 µg/mL overnight at 4°C, then washed with TBS-0.1%Tween 3 times with 5 minutes each and probed with a goat anti-rabbit IgG-HRP secondary antibody at a dilution of 1:10000 for 1.5 hour at RT. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog # EK1002) with Tanon 5200



system. A specific band was detected for Caspase8 at approximately 55KD. The expected band size for Caspase8 is at 55KD.

Immunohistochemistry

Image 2. IHC analysis of Caspase8 using anti-Caspase8 antibody . Caspase8 was detected in paraffin-embedded section of mouse spleen tissues. Heat mediated antigen retrieval was performed in citrate buffer (pH6, epitope retrieval solution) for 20 mins. The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1µg/ml rabbit anti-Caspase8 Antibody overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Strepavidin-Biotin-Complex (SABC)(Catalog # SA1022) with DAB as the chromogen.



Immunohistochemistry

Image 3. IHC analysis of Caspase8 using anti-Caspase8 antibody . Caspase8 was detected in paraffin-embedded section of rat intestine tissues. Heat mediated antigen retrieval was performed in citrate buffer (pH6, epitope retrieval solution) for 20 mins. The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1µg/ml rabbit anti-Caspase8 Antibody overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Strepavidin-Biotin-Complex (SABC)(Catalog # SA1022) with DAB as the chromogen.

Please check the product details page for more images. Overall 6 images are available for ABIN4886501.