Anti-Caspase-12 antibody ab18766

Overview

Product name: Anti-Caspase-12 antibody
Description: Rabbit polyclonal to Caspase-12
Host species: Rabbit
Specificity: Recognises full-length and cleaved fragment of Caspase-12
Tested applications: Suitable for: IHC-P, WB, ICC/IF
Species reactivity: Reacts with: Mouse, Rat
Predicted to work with: Human
Immunogen: Synthetic peptide corresponding to Mouse Caspase-12.
Database link: Q6UXS9
(Peptide available as ab38274)
Positive control: Spleen tissue lysate

Properties

Form: Liquid
Storage buffer: Preservative: 0.02% Sodium Azide
Constituents: 50% Glycerol, 1% BSA, PBS, pH 7.2
Purity: Protein A purified
Clonality: Polyclonal
Isotype: IgG

Applications

Our Abpromise guarantee covers the use of ab18766 in the following tested applications.
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.
**Function**
Has no protease activity. May reduce cytokine release in response to bacterial lipopolysaccharide during infections. Reduces activation of NF-kappa-B in response to TNF.

**Tissue specificity**
Detected in heart, kidney, liver, lung, pancreas, small intestine, spleen, stomach, thymus and testis.

**Sequence similarities**
Belongs to the peptidase C14A family. Contains 1 CARD domain.

**Images**
- Western blot - Anti-Caspase-12 antibody (ab18766) + Mouse spleen tissue lysate

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<td>IHC-P</td>
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**Notes**
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- WB:
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- ICC/IF:
  Use a concentration of 1 µg/ml.
ab18766 (1µg/ml) staining caspase-12 in human liver using an automated system (DAKO Autostainer Plus). Using this protocol there is strong staining of the cytoplasm and nuclei in hepatic arteries and plates of hepatic cells.

Sections were rehydrated and antigen retrieved with the Dako 3 in 1 AR buffers EDTA pH 9.0 in a DAKO PT link. Slides were peroxidase blocked in 3% H2O2 in methanol for 10 mins. They were then blocked with Dako Protein block for 10 minutes (containing casein 0.25% in PBS) then incubated with primary antibody for 20 min and detected with Dako envision flex amplification kit for 30 minutes. Colorimetric detection was completed with Diaminobenzidine for 5 minutes. Slides were counterstained with Haematoxylin and coverslipped under DePeX. Please note that, for manual staining, optimization of primary antibody concentration and incubation time is recommended. Signal amplification may be required.

ICC/IF image of ab18766 stained HeLa cells. The cells were 100% methanol fixed (5 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab18766, 1µg/ml) overnight at +4°C. The secondary antibody (green) was Alexa Fluor® 488 goat anti-rabbit IgG (H+L) used at a 1:1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1:200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43µM.
Western blot - Anti-Caspase-12 antibody (ab18766) + Rat kidney tissue lysate

Please note: All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE"

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