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anti-Caspase 2 antibody (AA 2-452)

2 Images



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Overview

Quantity:	100 μg
Target:	Caspase 2 (CASP2)
Binding Specificity:	AA 2-452
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This Caspase 2 antibody is un-conjugated
Application:	ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF)

Product Details

Immunogen:	Recombinant Human Caspase-2 protein (2-452AA)
Isotype:	IgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

Target Details

Target:	Caspase 2 (CASP2)
Alternative Name:	CASP2 (CASP2 Products)
Background:	Background: nvolved in the activation cascade of caspases responsible for apoptosis
	execution. Might function by either activating some proteins required for cell death or

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Aliases: CASP 2 antibody, CASP-2 antibody, Casp2 antibody, CASP2_HUMAN antibody, Caspase 2 antibody, Caspase 2 apoptosis related cysteine peptidase antibody, Caspase-2 subunit p12 antibody, Caspase2 antibody, ICH 1 antibody, ICH 1 protease antibody, ICH 1L antibody, ICH1 antibody, ICH1 antibody, NEDD-2 antibody, NEDD-2 antibody, Neural precursor cell expressed developmentally down-regulated protein 2 antibody, PPP1R57 antibody, Protease ICH-1 antibody, Protein phosphatase 1 regulatory subunit 57 antibody

UniProt: P42575

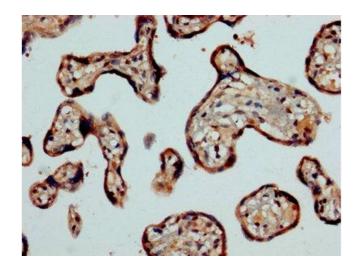
Pathways: Apoptosis, Caspase Cascade in Apoptosis, Neurotrophin Signaling Pathway

Application Details

Application Notes:	Recommended dilution: IHC:1:1000-1:2000, IF:1:200-1:500,
Restrictions:	For Research Use only

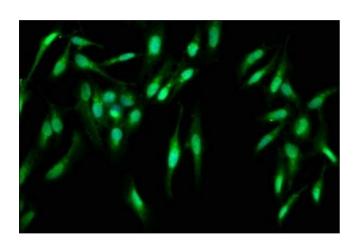
Handling

Format:	Liquid
Buffer:	Preservative: 0.03 % Proclin 300 Constituents: 50 % Glycerol, 0.01M PBS, PH 7.4
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C,-80 °C
Storage Comment:	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.



Immunohistochemistry

Image 1. IHC image of ABIN7146938 diluted at 1:1200 and staining in paraffin-embedded human placenta tissue 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°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



Immunofluorescence

Image 2. Immunofluorescence staining of Hela cells with ABIN7146938 at 1:400, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).