

Anti-Caspase 2 Antibody

Rabbit polyclonal antibody to Caspase 2 Catalog # AP60235

Product Information

Application WB, IHC
Primary Accession P42575
Reactivity Human, Rat
Host Rabbit
Clonality Polyclonal
Calculated MW 50685

Additional Information

Gene ID 835

Other Names ICH1; NEDD2; Caspase-2; CASP-2; Neural precursor cell expressed

developmentally down-regulated protein 2; NEDD-2; Protease ICH-1

Target/Specificity Recognizes endogenous levels of Caspase 2 protein.

Dilution WB~~WB (1/500 - 1/1000), IHC (1/100 - 1/200) IHC~~WB (1/500 - 1/1000), IHC

(1/100 - 1/200)

Format Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

Protein Information

Name CASP2

Synonyms ICH1, NEDD2

Function Is a regulator of the cascade of caspases responsible for apoptosis execution

(PubMed:<u>11156409</u>, PubMed:<u>15073321</u>, PubMed:<u>8087842</u>). Might function by either activating some proteins required for cell death or inactivating proteins necessary for cell survival (PubMed:<u>15073321</u>). Associates with PIDD1 and CRADD to form the PIDDosome, a complex that activates CASP2 and triggers

apoptosis in response to genotoxic stress (PubMed: 15073321).

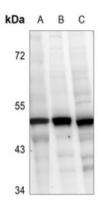
Tissue Location Expressed at higher levels in the embryonic lung, liver and kidney than in the

heart and brain. In adults, higher level expression is seen in the placenta, lung, kidney, and pancreas than in the heart, brain, liver and skeletal muscle

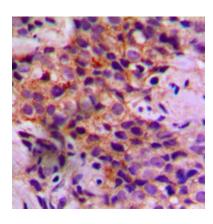
Background

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human Caspase 2. The exact sequence is proprietary.

Images



Western blot analysis of Caspase 2 expression in Jurkat (A), Hela (B), PC12 (C) whole cell lysates.



Immunohistochemical analysis of Caspase 2 staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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