

## **Product Data Sheet**

## **Anti-Caspase 8 Antibody**

Catalog # Source Reactivity Applications

CPA1154 Rabbit H, M, R WB, IH

**Description** Rabbit polyclonal antibody to Caspase 8

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human Caspase 8. The exact sequence is proprietary.

**Purification** The antibody was purified by immunogen affinity chromatography.

**Specificity** Recognizes endogenous levels of Caspase 8 protein.

**Clonality** Polyclonal

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

and 0.01% sodium azide.

**Dilution** WB (1/500 - 1/1000), IH (1/100 - 1/200)

Gene Symbol CASP8

Alternative Names MCH5; Caspase-8; CASP-8; Apoptotic cysteine protease; Apoptotic protease Mch-5;

CAP4; FADD-homologous ICE/ced-3-like protease; FADD-like ICE; FLICE; ICE-like

apoptotic protease 5; MORT1-associated ced-3 homolog; MACH

Entrez Gene 841 (Human); 12370 (Mouse)

SwissProt Q14790 (Human); O89110 (Mouse)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

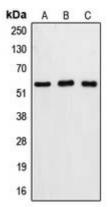
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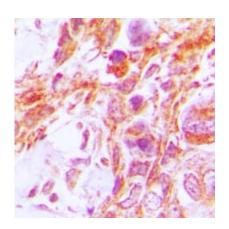




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Western blot analysis of Caspase 8 expression in DLD (A), mouse liver (B), Raw264.7 (C) whole cell lysates.



Immunohistochemical analysis of Caspase 8 staining in human lung 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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