

Product Data Sheet

Anti-Caspase 8 (Phospho-S347) Antibody

Catalog # Source Reactivity Applications

CPA1152 Rabbit H, Mk WB, IH

Description Rabbit polyclonal antibody to Caspase 8 (Phospho-S347)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S347 of human Caspase 8 protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of Caspase 8 protein only when phosphorylated at

S347.

Clonality Polyclonal

Conjugation

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)

SwissProt Q14790 (Human)

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

freeze/thaw cycles.

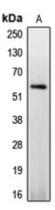
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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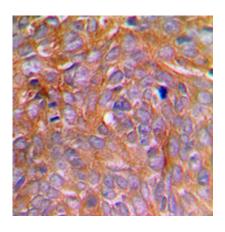
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Western blot analysis of Caspase 8 (Phospho-S347) expression in Jurkat PMA-treated (A) whole cell lysates. (Predicted band size: 55 kD; Observed band size: 57 kD)



Immunohistochemical analysis of Caspase 8 (Phospho-S347) 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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