

# **Product Data Sheet**

# **Anti-Caspase 3 (Phospho-S150) Antibody**

Catalog # Source Reactivity Applications

CPA4589 Rabbit H, M, R, B, D, Mk, Rb WB, IH

**Description** Rabbit polyclonal antibody to Caspase 3 (Phospho-S150)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S150 of human Caspase 3 protein. The exact sequence is proprietary.

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

**Specificity** Recognizes endogenous levels of Caspase 3 protein only when phosphorylated at

S150.

**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 CASP3

Alternative Names CPP32; Caspase-3; CASP-3; Apopain; Cysteine protease CPP32; CPP-32; Protein

Yama; SREBP cleavage activity 1; SCA-1

**Entrez Gene** 836 (Human); 12367 (Mouse); 25402 (Rat)

SwissProt P42574 (Human); P70677 (Mouse); P55213 (Rat)

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

freeze/thaw cycles.

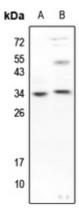
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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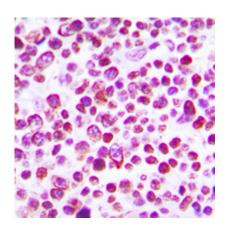
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Western blot analysis of Caspase 3 (Phospho-S150) expression in A549 (A), LO2 (B) whole cell lysates. (Predicted band size: 31 kD; Observed band size: 35 kD)



Immunohistochemical analysis of Caspase 3 (Phospho-S150) staining in human tonsil 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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