

## **Product Data Sheet**

## Anti-Presenilin 1 (Phospho-S357) Antibody

Catalog # Source Reactivity Applications

CPA6057 Rabbit H, M, R, C, Mk WB, IH

**Description** Rabbit polyclonal antibody to Presentilin 1 (Phospho-S357)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S357 of human Presenilin 1 protein. The exact sequence is proprietary.

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

**Specificity** Recognizes endogenous levels of Presenilin 1 protein only when phosphorylated at

S357.

**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 (1/50 - 1/200)

Gene Symbol PSEN1

Alternative Names AD3; PS1; PSNL1; Presenilin-1; PS-1; Protein S182

Entrez Gene 5663 (Human); 19164 (Mouse); 29192 (Rat)

SwissProt P49768 (Human); P49769 (Mouse); P97887 (Rat)

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

freeze/thaw cycles.

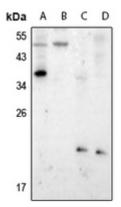
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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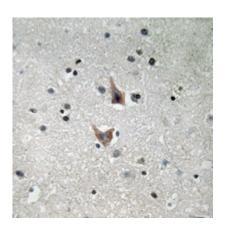
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Western blot analysis of Presenilin 1 (Phospho-S357) expression in HCT116 (A), Beas2B (B), mouse embryo (C), rat brain (D) whole cell lysates. (Predicted band size: 52 kD; Observed band size: 52; 22 kD)



Immunohistochemical analysis of Presenilin 1 (Phospho-S357) staining in human brain 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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