

Product Data Sheet

Anti-SHPTP1 (Phospho-Y536) Antibody

Catalog # Source Reactivity Applications

CPA7005 Rabbit H, M, R WB, IH, IF/IC

Description Rabbit polyclonal antibody to SHPTP1 (Phospho-Y536)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

Y536 of human SHPTP1 protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of SHPTP1 protein only when phosphorylated at

Y536.

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), IF/IC (1/100 - 1/500)

Gene Symbol PTPN6

Alternative Names HCP; PTP1C; Tyrosine-protein phosphatase non-receptor type 6; Hematopoietic cell

protein-tyrosine phosphatase; Protein-tyrosine phosphatase 1C; PTP-1C;

Protein-tyrosine phosphatase SHP-1; SH-PTP1

Entrez Gene 5777 (Human); 15170 (Mouse); 116689 (Rat)

SwissProt P29350 (Human); P29351 (Mouse); P81718 (Rat)

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

freeze/thaw cycles.

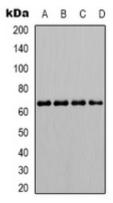
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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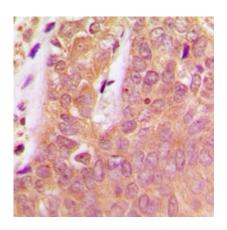
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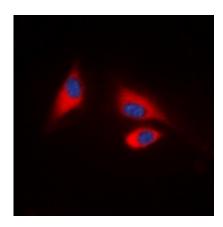
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Western blot analysis of SHPTP1 (Phospho-Y536) expression in K562 (A), Jurkat (B), NIH3T3 (C), Raw264.7 (D) whole cell lysates. (Predicted band size: 67 kD; Observed band size: 68 kD)



Immunohistochemical analysis of SHPTP1 (Phospho-Y536) 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.



Immunofluorescent analysis of SHPTP1 (Phospho-Y536) staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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