

Product Data Sheet

Anti-PINX1 Antibody

Catalog # Source Reactivity Applications

CPA4155 Rabbit H, M, R WB, IH

Description Rabbit polyclonal antibody to PINX1

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

region of human PINX1. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of PINX1 protein.

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 PINX1

Alternative Names LPTL; LPTS; PIN2/TERF1-interacting telomerase inhibitor 1; Liver-related putative

tumor suppressor; Pin2-interacting protein X1; Protein 67-11-3; TRF1-interacting

protein 1

Entrez Gene 54984 (Human); 72400 (Mouse); 305963 (Rat)

SwissProt Q96BK5 (Human); Q9CZX5 (Mouse); A4L691 (Rat)

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

freeze/thaw cycles.

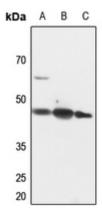
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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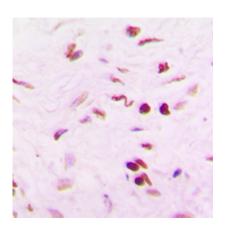
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Western blot analysis of PINX1 expression in rat heart (A), rat muscle (B), rat brain (C) whole cell lysates. (Predicted band size: 37 kD; Observed band size: 45 kD)



Immunohistochemical analysis of PINX1 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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