

Product Data Sheet

Anti-PPP2R5D Antibody

Catalog # Source Reactivity Applications

CPA4167 Rabbit H, M, B, Mk WB, IH

Description Rabbit polyclonal antibody to PPP2R5D

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the C-term

region of human PPP2R5D. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of PPP2R5D 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 PPP2R5D

Alternative Names Serine/threonine-protein phosphatase 2A 56 kDa regulatory subunit delta isoform;

PP2A B subunit isoform B'-delta; PP2A B subunit isoform B56-delta; PP2A B subunit

isoform PR61-delta: PP2A B subunit isoform R5-delta

Entrez Gene 5528 (Human)

SwissProt Q14738 (Human)

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

freeze/thaw cycles.

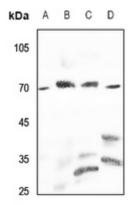
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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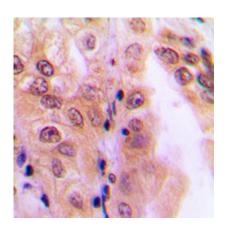
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Western blot analysis of PPP2R5D expression in HEK293 (A), LOVO (B), mouse brain (C), mouse kidney (D) whole cell lysates. (Predicted band size: 69 kD; Observed band size: 70 kD)



Immunohistochemical analysis of PPP2R5D 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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