

### **Product Data Sheet**

# **Anti-PRKAR1B Antibody**

Catalog # Source Reactivity Applications

CPA1932 Rabbit H, M, R, C, D, Z WB, IH

**Description** Rabbit polyclonal antibody to PRKAR1B

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

region of human PRKAR1B. The exact sequence is proprietary.

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

**Specificity** Recognizes endogenous levels of PRKAR1B 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 PRKAR1B

Alternative Names cAMP-dependent protein kinase type I-beta regulatory subunit

Entrez Gene 5575 (Human); 19085 (Mouse)

SwissProt P31321 (Human); P12849 (Mouse); P81377 (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

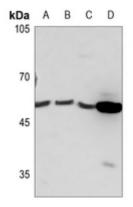
Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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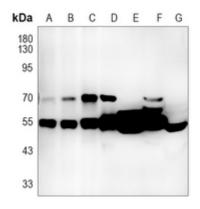
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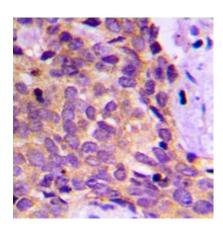
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Western blot analysis of PRKAR1B expression in HEK293T (A), U2OS (B), MCF7 (C), mouse liver (D) whole cell lysates. (Predicted band size: 43 kD; Observed band size: 55 kD)



Western blot analysis of PRKAR1B expression in A2780 (A), MG63 (B), Jurkat (C), mouse brain (D), mouse liver (E), rat brain (F), rat liver (G) whole cell lysates. (Predicted band size: 43 kD; Observed band size: 55 kD)



Immunohistochemical analysis of PRKAR1B 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.

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