

Product Data Sheet

Anti-Adenylate Cyclase 5/6 Antibody

Catalog # Source Reactivity Applications

CPA1019 Rabbit H, M, R, D, Mk, Rb, Z WB, IF/IC

Description Rabbit polyclonal antibody to Adenylate Cyclase 5/6

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

region of human Adenylate Cyclase 5/6. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of Adenylate Cyclase 5/6 protein.

Clonality Polyclonal

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

Gene Symbol ADCY5; ADCY6

Alternative Names ADCY5; Adenylate cyclase type 5; ATP pyrophosphate-lyase 5; Adenylate cyclase type

V; Adenylyl cyclase 5; ADCY6; KIAA0422; Adenylate cyclase type 6; ATP

pyrophosphate-lyase 6; Adenylate cyclase type VI; Adenylyl cyclase 6;

Ca(2+)-inhibitable adenylyl cyclase

Entrez Gene 111, 112 (Human); 224129, 11512 (Mouse); 64532 (Rat)

SwissProt 095622, O43306 (Human); P84309, Q01341 (Mouse); Q04400, Q03343 (Rat)

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

freeze/thaw cycles.

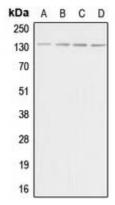
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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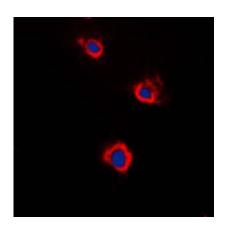
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Western blot analysis of Adenylate Cyclase 5/6 expression in HeLa (A), Raw264.7 (B), PC12 (C), rat brain (D) whole cell lysates.



Immunofluorescent analysis of Adenylate Cyclase 5/6 staining in Raw264.7 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 humidified 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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