

Product Data Sheet

Anti-Kir6.2 Antibody

Catalog # Source Reactivity Applications

CPA4045 Rabbit H, M, R, B, P, Rb WB, IH, IF/IC

Description Rabbit polyclonal antibody to Kir6.2

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

region of human Kir6.2. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of Kir6.2 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), IF/IC (1/100 - 1/500)

Gene Symbol KCNJ11

Alternative Names ATP-sensitive inward rectifier potassium channel 11; IKATP; Inward rectifier K(+)

channel Kir6.2; Potassium channel, inwardly rectifying subfamily J member 11

Entrez Gene 3767 (Human); 16514 (Mouse); 83535 (Rat)

SwissProt Q14654 (Human); Q61743 (Mouse); P70673 (Rat)

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

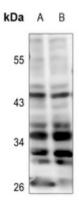
freeze/thaw cycles.

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

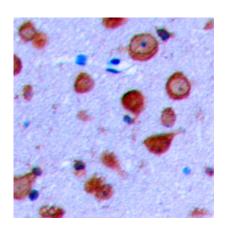
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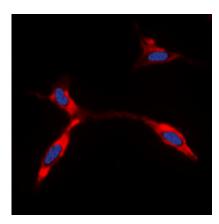
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Western blot analysis of Kir6.2 expression in HEK293T (A), K562 (B) whole cell lysates. (Predicted band size: 43 kD; Observed band size: 44; 31 kD)



Immunohistochemical analysis of Kir6.2 staining in human brain cortex 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 Kir6.2 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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