

### **Product Data Sheet**

# **Anti-Kir3.3 Antibody**

Catalog # Source Reactivity Applications

CPA1649 Rabbit H, M, R WB, IH

**Description** Rabbit polyclonal antibody to Kir3.3

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

region of human Kir3.3. The exact sequence is proprietary.

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

**Specificity** Recognizes endogenous levels of Kir3.3 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 KCNJ9

Alternative Names GIRK3; G protein-activated inward rectifier potassium channel 3; GIRK-3; Inward

rectifier K(+) channel Kir3.3; Potassium channel, inwardly rectifying subfamily J

member 9

Entrez Gene 3765 (Human); 16524 (Mouse); 116560 (Rat)

SwissProt Q92806 (Human); P48543 (Mouse); Q63511 (Rat)

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

freeze/thaw cycles.

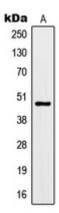
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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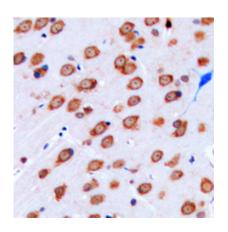
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Western blot analysis of Kir3.3 expression in rat brain (A) whole cell lysates. (Predicted band size: 44 kD; Observed band size: 48 kD)



Immunohistochemical analysis of Kir3.3 staining in human brain 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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