

Anti-KCNJ12 Antibody



Description This gene encodes an inwardly rectifying K+ channel which may be

blocked by divalent cations. This protein is thought to be one of multiple inwardly rectifying channels which contribute to the cardiac inward rectifier current (IK1). The gene is located within the Smith-Magenis

syndrome region on chromosome 17.

Model STJ116587

Host Rabbit

Reactivity Mouse, Rat

Applications WB

Immunogen Recombinant fusion protein containing a sequence corresponding to amino

acids 354-433 of human KCNJ12 (NP_066292.2).

Gene ID 3768

Gene Symbol KCNJ12

Dilution range WB 1:500 - 1:2000

Purification Affinity purification

Note For Research Use Only (RUO).

Protein Name ATP-sensitive inward rectifier potassium channel 12 Inward rectifier K(+

channel Kir2.2 IRK-2 Inward rectifier K(+ channel Kir2.2v Potassium channel

inwardly rectifying subfamily J member 12

Molecular Weight 49.001 kDa

Clonality Polyclonal

Conjugation Unconjugated

Isotype IgG

Formulation PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

Storage Instruction Store at -20C. Avoid freeze / thaw cycles.

Database Links HGNC:62580MIM:602323Reactome:R-HSA-1296041

Alternative Names ATP-sensitive inward rectifier potassium channel 12 Inward rectifier K(+

channel Kir2.2 IRK-2 Inward rectifier K(+ channel Kir2.2v Potassium channel

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Function Inward rectifying potassium channel that is activated by phosphatidylinositol

4,5-bisphosphate and that probably participates in controlling the resting membrane potential in electrically excitable cells, Probably participates in establishing action potential waveform and excitability of neuronal and muscle tissues, Inward rectifier potassium channels are characterized by a greater tendency to allow potassium to flow into the cell rather than out of it, Their voltage dependence is regulated by the concentration of extracellular

potassium

Cellular Localization Membrane

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