

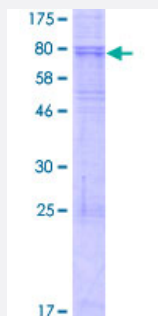
Full-Length

KCNJ1 (Human) Recombinant Protein (P01)

Catalog # H00003758-P01

Size 25 ug, 10 ug

Applications



Specification

Product Description

Human KCNJ1 full-length ORF (NP_722449.1, 1 a.a. - 389 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence

MPTVYLCSEQIRVLTESMFKHLRKWVVTRFFGHSRQRARLVSKDGRCNIEFGNVEAQSRFFVVDI
WTTVLDLKWRYKMTIFITAFLLGFWFFFGLLWYAVAYHKDLPEFHPSANHTPCVENINGLTSAFLFS
LETQVTIGYGFRVTEQCATAIFLLIFQSILGVIINSFMCGAILAKISRPKKRAKTITFSKNAVISKRGK
LCLLIRVANLRKSLIGSHYGLKLLKTTVTPEGETILDQININFVVDAGNENLFFISPLTYHVIDHNSPFF
HMAAETLLQQDFELVVFLDGTVESTSATCQVRTSYVPEEVLWGYRFAPVSKTKEGKYRVDHFHNF
SKTVEVETPHCAMCLYNEKDVRARMKRGYDNPFISEVNETDDTKM

Host

Wheat Germ (in vitro)

Theoretical MW (kDa)

71

Interspecies Antigen Sequence

Mouse (91)

Preparation Method

[in vitro wheat germ expression system](#)

Purification

Glutathione Sepharose 4 Fast Flow

Quality Control Testing

12.5% SDS-PAGE Stained with Coomassie Blue.

Storage Buffer

50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.

Storage Instruction

Store at -80°C. Aliquot to avoid repeated freezing and thawing.

Note

Best use within three months from the date of receipt of this protein.

Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

Gene Info — KCNJ1

Entrez GeneID[3758](#)**GeneBank Accession#**[NM_153765.1](#)**Protein Accession#**[NP_722449.1](#)**Gene Name**

KCNJ1

Gene Alias

KIR1.1, ROMK, ROMK1

Gene Description

potassium inwardly-rectifying channel, subfamily J, member 1

Omim ID[241200 600359](#)**Gene Ontology**[Hyperlink](#)**Gene Summary**

Potassium channels are present in most mammalian cells, where they participate in a wide range of physiologic responses. The protein encoded by this gene is an integral membrane protein and inward-rectifier type potassium channel. It is activated by internal ATP and probably plays an important role in potassium homeostasis. The encoded protein has a greater tendency to allow potassium to flow into a cell rather than out of a cell. Mutations in this gene have been associated with antenatal Bartter syndrome, which is characterized by salt wasting, hypokalemic alkalosis, hypercalciuria, and low blood pressure. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq]

Other Designations

ATP-regulated potassium channel ROM-K|ATP-sensitive inward rectifier potassium channel 1|OTTHUMP00000045938|inwardly rectifying K+ channel|potassium inwardly-rectifying channel J1

Disease

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