

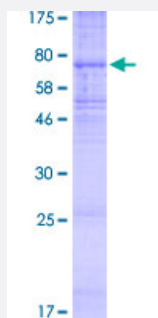
Full-Length

KCNJ11 (Human) Recombinant Protein (P01)

Catalog # H00003767-P01

Size 25 ug, 10 ug

Applications



Specification

Product Description

Human KCNJ11 full-length ORF (NP_000516.3, 1 a.a. - 390 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence

MLSRKGIIPEEYVLTRLAEDPAKPRYRARQRRARFVSKKGNCNVAHKNIREQGRFLQDVFTTLVDL
KWPHTLLIFTMSFLCSWLLFAMAWWLIFAHGDLPSEGTAEPVTSIHSFSSAFLFSIEVQVTIGF
GGRMVTEECPLAILIVQNIVGLMINAIMLGCIFMKTAQAHRAETLIFSKHAVIALRHGRLCFMLRVG
DLRKSMIISATIHMQVVRKTTSPGEVVPLHQVDIPMENGVGGSIFLVAPLIYHVIDANSPLYDLAP
SDLHHHQDLEIVILEGVVETTGITQARTSYLADEILWGQRFVPMVAEEDGRYSVDYSKFGNTVKVP
TPLCTARQLDEDHSLLEALTLASARGPLRKRSVPMKAKPKFSISPDSLS

Host

Wheat Germ (in vitro)

Theoretical MW (kDa)

69.9

Interspecies Antigen Sequence

Mouse (96); Rat (96)

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 — KCNJ11

Entrez GeneID[3767](#)**GeneBank Accession#**[NM_000525.3](#)**Protein Accession#**[NP_000516.3](#)**Gene Name**

KCNJ11

Gene Alias

BIR, HHF2, IKATP, KIR6.2, MGC133230, PHHI, TNDM3

Gene Description

potassium inwardly-rectifying channel, subfamily J, member 11

Omim ID[125853](#) [600937](#) [601820](#) [606176](#) [610582](#)**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. The encoded protein, which has a greater tendency to allow potassium to flow into a cell rather than out of a cell, is controlled by G-proteins and is found associated with the sulfonylurea receptor SUR. Mutations in this gene are a cause of familial persistent hyperinsulinemic hypoglycemia of infancy (PHHI), an autosomal recessive disorder characterized by unregulated insulin secretion. Defects in this gene may also contribute to autosomal dominant non-insulin-dependent diabetes mellitus type II (NIDDM), transient neonatal diabetes mellitus type 3 (TNDM3), and permanent neonatal diabetes mellitus (PNDM). [provided by RefSeq]

Other Designations

ATP-sensitive inward rectifier potassium channel 11|beta-cell inward rectifier subunit|inwardly rectifying potassium channel KIR6.2|potassium inwardly-rectifying channel J11

Pathway

- [Type II diabetes mellitus](#)

Disease

- [Alzheimer disease](#)
- [Atherosclerosis](#)
- [Birth Weight](#)
- [Calcinosis](#)
- [Carcinoma](#)
- [Cardiovascular Diseases](#)
- [Coronary Artery Disease](#)
- [Coronary Disease](#)
- [Death](#)
- [Diabetes](#)
- [Diabetes Complications](#)
- [Diabetes Mellitus](#)
- [Diabetic Angiopathies](#)
- [Diabetic Nephropathies](#)
- [Disease Progression](#)
- [Edema](#)
- [Epilepsy](#)
- [Fetal Diseases](#)
- [Genetic Predisposition to Disease](#)
- [Glucose Intolerance](#)
- [Glucose Metabolism Disorders](#)

- [HIV Infections](#)
- [Hyperglycemia](#)
- [Hyperinsulinism](#)
- [Hypertension](#)
- [Hypoglycemia](#)
- [Insulin Resistance](#)
- [Kidney Failure](#)
- [Malnutrition](#)
- [Metabolic Syndrome X](#)
- [Myocardial Infarction](#)
- [Neoplasms](#)
- [Obesity](#)
- [Osteoporosis](#)
- [Persistent Hyperinsulinemia Hypoglycemia of Infancy](#)
- [Polycystic Ovary Syndrome](#)
- [Prostatic Neoplasms](#)
- [Starvation](#)
- [Weight Gain](#)