

## KCNJ11 抗原（重组蛋白）

中文名称： KCNJ11 抗原（重组蛋白）

英文名称： KCNJ11 Antigen (Recombinant Protein)

别名： BIR; HHF2; PHHI; IKATP; TNDM3; KIR6.2

储存： 冷冻（-20℃）

相关类别： 抗原

概述：

Fusion protein corresponding to a region derived from 167-390 amino acids of human KCNJ11

技术规格：

<b>Full name:</b>	potassium inwardly-rectifying channel, subfamily J, member 11
<b>Synonyms:</b>	BIR; HHF2; PHHI; IKATP; TNDM3; KIR6.2
<b>Swissprot:</b>	Q14654
<b>Gene Accession:</b>	BC112358
<b>Purity:</b>	>85%, as determined by Coomassie blue stained SDS-PAGE
<b>Expression system:</b>	Escherichia coli
<b>Tags:</b>	His tag C-Terminus, GST tag N-Terminus
<b>Background:</b>	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 unregula

ted insulin secretion. Defects in this gene may also contribute to a utosomal dominant non-insulin-dependent diabetes mellitus type II (NIDDM), transient neonatal diabetes mellitus type 3 (TNDM3), and permanent neonatal diabetes mellitus (PNDM). Multiple alternatively spliced transcript variants that encode different protein isoforms have been described for this gene.