

Datasheet for ABIN1395631  
**anti-KCNJ11 antibody (pThr224) (Biotin)**



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## Overview

Quantity:	100 µL
Target:	KCNJ11
Binding Specificity:	pThr224
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This KCNJ11 antibody is conjugated to Biotin
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (Paraffin-embedded Sections) (IHC (p)), Immunohistochemistry (Frozen Sections) (IHC (fro))

## Product Details

Immunogen:	KLH conjugated synthetic phosphopeptide derived from human Kir6.2 around the phosphorylation site of Thr224
Isotype:	IgG
Predicted Reactivity:	Human, Mouse, Rat, Dog, Cow, Sheep, Pig, Rabbit
Purification:	Purified by Protein A.

## Target Details

Target:	KCNJ11
Alternative Name:	Kir6.2 ( <a href="#">KCNJ11 Products</a> )

## Target Details

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**Background:** Synonyms: p-Kir6.2 phospho T224, ATP sensitive inward rectifier potassium channel 11, Beta cell inward rectifier subunit, mBIR, BIR, HHF 2, HHF2, IKATP, Inward rectifier K<sup>+</sup> channel Kir6.2, Inwardly rectifying potassium channel KIR6.2, IRK 11, IRK11, KCNJ11, Kir 6.2, Kir6.2, MGC133230, PPHI, Potassium channel, inwardly rectifying subfamily J member 11, Potassium inwardly rectifying channel J11, TNDM 3, TNDM3.

Background: 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). Multiple alternatively spliced transcript variants that encode different protein isoforms have been described for this gene. [provided by RefSeq]

**Pathways:** [Negative Regulation of Hormone Secretion](#)

## Application Details

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**Application Notes:** WB 1:300-5000  
IHC-P 1:200-400  
IHC-F 1:100-500

**Restrictions:** For Research Use only

## Handling

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**Format:** Liquid

**Concentration:** 1 µg/µL

**Buffer:** Aqueous buffered solution containing 0.01M TBS (pH 7.4) with 1 % BSA, 0.03 % Proclin300 and 50 % Glycerol.

**Preservative:** ProClin

**Precaution of Use:** This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be handled by trained staff only.

## Handling

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Storage: -20 °C

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Storage Comment: Store at -20°C for 12 months.

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Expiry Date: 12 months