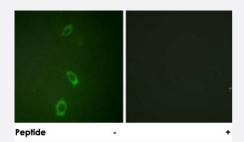


KCNJ1 polyclonal antibody

Catalog # PAB18418 Size 100 ug

Applications



Immunofluorescence

Immunofluorescence analysis of A-549 cells, using KCNJ1 polyclonal antibody (Cat # PAB18418).

Peptide "+" means "peptide blocking".

Specification	
Product Description	Rabbit polyclonal antibody raised against synthetic peptide of KCNJ1.
Immunogen	A synthetic peptide corresponding to human KCNJ1.
Host	Rabbit
Reactivity	Human, Mouse, Rat
Specificity	This antibody is specific to KCNJ1.
Form	Liquid
Purification	Affinity purification
Concentration	1 mg/mL
Recommend Usage	Immunofluorescence (1:500-1:1000) ELISA (1:10000) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS, 150mM NaCl, pH 7.4 (50% glycerol, 0.02% sodium azide)



Product Information

Storage Instruction	Store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

Applications

Immunofluorescence

Immunofluorescence analysis of A-549 cells, using KCNJ1 polyclonal antibody (Cat # PAB18418). Peptide "+" means "peptide blocking".

• Enzyme-linked Immunoabsorbent Assay

Gene Info — KCNJ1	
Entrez GenelD	3758
Protein Accession#	P48048
Gene Name	KCNJ1
Gene Alias	KIR1.1, ROMK, ROMK1
Gene Description	potassium inwardly-rectifying channel, subfamily J, member 1
Omim ID	<u>241200</u> <u>600359</u>
Gene Ontology	<u>Hyperlink</u>
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 potassi um to flow into a cell rather than out of a cell. Mutations in this gene have been associated with ant enatal Bartter syndrome, which is characterized by salt wasting, hypokalemic alkalosis, hypercalci uria, and low blood pressure. Multiple transcript variants encoding different isoforms have been fo und for this gene. [provided by RefSeq

Disease



- Cardiovascular Diseases
- Diabetes Mellitus
- Edema
- Genetic Predisposition to Disease
- Hyperparathyroidism
- Hypertension
- Hypotension
- Tobacco Use Disorder