

OriGene Technologies, Inc.

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Product datasheet for PH310938

GIRK2 (KCNJ6) (NM_002240) Human Mass Spec Standard

Product data:

Product Type:	Mass Spec Standards
Description:	KCNJ6 MS Standard C13 and N15-labeled recombinant protein (NP_002231)
Species:	Human
Expression Host:	HEK293
Expression cDNA Clone or AA Sequence:	RC210938
Predicted MW:	48.5 kDa
Protein Sequence:	>RC210938 protein sequence <mark>Red=</mark> Cloning site Green=Tags(s)
	MAKLTESMTNVLEGDSMDQDVESPVAIHQPKLPKQARDDLPRHISRDRTKRKIQRYVRKDGKCNVHHGNV RETYRYLTDIFTTLVDLKWRFNLLIFVMVYTVTWLFFGMIWWLIAYIRGDMDHIEDPSWTPCVTNLNGFV SAFLFSIETETTIGYGYRVITDKCPEGIILLLIQSVLGSIVNAFMVGCMFVKISQPKKRAETLVFSTHAV ISMRDGKLCLMFRVGDLRNSHIVEASIRAKLIKSKQTSEGEFIPLNQTDINVGYYTGDDRLFLVSPLIIS HEINQQSPFWEISKAQLPKEELEIVVILEGMVEATGMTCQARSSYITSEILWGYRFTPVLTLEDGFYEVD YNSFHETYETSTPSLSAKELAELASRAELPLSWSVSSKLNQHAELETEEEEKNLEEQTERNGDVANLENE SKV
	TRTRPLEQKLISEEDLAANDILDYKDDDDKV
Tag:	C-Myc/DDK
Purity:	> 80% as determined by SDS-PAGE and Coomassie blue staining
Concentration:	>0.05 µg/µL as determined by microplate BCA method
Labeling Method:	Labeled with [U- 13C6, 15N4]-L-Arginine and [U- 13C6, 15N2]-L-Lysine
Buffer:	25 mM Tris-HCl, 100 mM glycine, pH 7.3
Storage:	Store at -80°C. Avoid repeated freeze-thaw cycles.
Stability:	Stable for 3 months from receipt of products under proper storage and handling conditions.
RefSeq:	<u>NP 002231</u>
RefSeq Size:	2537
RefSeq ORF:	1269
Synonyms:	BIR1; GIRK-2; GIRK2; hiGIRK2; KATP-2; KATP2; KCNJ7; KIR3.2; KPLBS



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	GIRK2 (KCNJ6) (NM_002240) Human Mass Spec Standard – PH310938		
Locus ID:	3763		
UniProt ID:	<u>P48051</u>		
Cytogenetics:	21q22.13		
Summary:	This gene encodes a member of the G protein-coupled inwardly-rectifying potassium channel family of inward rectifier potassium channels. This type of potassium channel allows a greater flow of potassium into the cell than out of it. These proteins modulate many physiological processes, including heart rate in cardiac cells and circuit activity in neuronal cells, through G- protein coupled receptor stimulation. Mutations in this gene are associated with Keppen- Lubinsky Syndrome, a rare condition characterized by severe developmental delay, facial dysmorphism, and intellectual disability. [provided by RefSeq, Apr 2015]		
Protein Families	: Druggable Genome, Ion Channels: Potassium, Transmembrane		

Product images:

116	_	-	
66	_	-	
45	_	-	
35	_	-	
25	_	-	
18	_	-	
14	-	-	

Coomassie blue staining of purified KCNJ6 protein (Cat# [TP310938]). The protein was produced from HEK293T cells transfected with KCNJ6 cDNA clone (Cat# [RC210938]) using MegaTran 2.0 (Cat# [TT210002]).

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