



# Mouse anti-Human KCNQ1 monoclonal antibody, clone T48B-20 (CABT-B10507)

This product is for research use only and is not intended for diagnostic use.

## PRODUCT INFORMATION

<b>Specificity</b>	Detects ~75KDa.
<b>Immunogen</b>	Recombinant fusion protein corresponding to amino acids 2-101 of human KCNQ1.
<b>Isotype</b>	IgG1
<b>Source/Host</b>	Mouse
<b>Species Reactivity</b>	Human
<b>Clone</b>	T48B-20
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	WB, IHC, IF, IP
<b>Format</b>	Liquid
<b>Buffer</b>	In PBS, pH 7.4 (50% glycerol, 0.09% sodium azide)
<b>Storage</b>	Store at -20°C. Aliquot to avoid repeated freezing and thawing.

## BACKGROUND

<b>Introduction</b>	This gene encodes a voltage-gated potassium channel required for repolarization phase of the cardiac action potential. This protein can form heteromultimers with two other potassium channel proteins, KCNE1 and KCNE3. Mutations in this gene are associated with hereditary long QT syndrome 1 (also known as Romano-Ward syndrome), Jervell and Lange-Nielsen syndrome, and familial atrial fibrillation. This gene exhibits tissue-specific imprinting, with
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preferential expression from the maternal allele in some tissues, and biallelic expression in others. This gene is located in a region of chromosome 11 amongst other imprinted genes that are associated with Beckwith-Wiedemann syndrome (BWS), and itself has been shown to be disrupted by chromosomal rearrangements in patients with BWS. Alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Aug 2011]

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<b>Keywords</b>	KCNQ1; potassium channel, voltage gated KQT-like subfamily Q, member 1; LQT; RWS; WRS; LQT1; SQT2; ATFB1; ATFB3; JLNS1; KCNA8; KCNA9; Kv1.9; Kv7.1; KVLQT1; potassium voltage-gated channel subfamily KQT member 1; slow delayed rectifier channel subunit; voltage-gated potassium channel subunit Kv7.1; kidney and cardiac voltage dependend K+ channel; potassium voltage-gated channel, KQT-like subfamily, member 1; IKs producing slow voltage-gated potassium channel subunit alpha KvLQT1;
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## GENE INFORMATION

<b>Entrez Gene ID</b>	<a href="#">3784</a>
<b>UniProt ID</b>	<a href="#">Q96AI9</a>
<b>Pathway</b>	Cholinergic synapse;Gastric acid secretion;Neuronal System;Pancreatic secretion;Potassium Channels;
<b>Function</b>	calmodulin binding;contributes_to delayed rectifier potassium channel activity;outward rectifier potassium channel activity;voltage-gated potassium channel activity;

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