

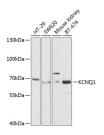
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# KCNQ1 Polyclonal Antibody

Catalog No. E-AB-60659 Reactivity H,M,R Store at -20°C. Avoid freeze / thaw cycles. **Storage** Host Rabbit **Applications Isotype IgG** 

**Note:** Centrifuge before opening to ensure complete recovery of vial contents.

# **Images**



Western blot analysis of extracts of various cell lines using KCNQ1 Polyclonal Antibody at dilution of 1:1000.

### **Immunogen Information**

**Immunogen** Recombinant fusion protein of human KCNQ1

(NP\_861463.1).

GeneID 3784 **Swissprot** P51787

**Synonyms** KCNQ1,ATFB1,ATFB3,JLNS1,KCNA8,KCNA9,KV

LQT1,Kv1.9,Kv7.1,LQT,LQT1,RWS,SQT2,WRS

#### **Product Information**

Calculated MW 61kDa/74kDa

**Observed MW** 70kDa

**Buffer** PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

**Purify** Affinity purification Dilution WB 1:500-1:2000

# **Background**

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 tissuespecific imprinting, with 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.