

Anti-KCNE1 (Potassium voltage-gated channel subfamily E member 1) Polyclonal Antibody

Cat. #: 60B882

Description:

KCNE1 (Potassium voltage-gated channel subfamily E member 1) is an ancillary protein that assembles as a beta subunit with a voltage-gated potassium channel complex of pore-forming alpha subunits. KCNE1 modulates the gating kinetics and enhances stability of the channel complex. The KCNE1 assembled with KCNQ1/KVLQT1 is proposed to form the slowly activating delayed rectifier cardiac potassium (IKs) channel. The outward current reaches its steady state only after 50 seconds. The KCNE1 assembled with KCNH2/HERG may modulate the rapidly activating component of the delayed rectifying potassium current in heart (IKr). KCNE1 is associated with KCNQ1/KVLQT1 and KCNH2/HERG. Defects in KCNE1 are a cause of the autosomal recessive Jervell and Lange-Nielsen syndrome (JLNS). JLNS comprises profound congenital sensorineural deafness associated with syncopal episodes. These are caused by ventricular tachyarrhythmia secondary to abnormal repolarization, manifested by a prolonged QT interval on the electrocardiogram.

Immunogen/Specificity:

Polyclonal antibody produced in rabbits immunizing with a synthetic peptide corresponding to N-terminal residues of human KCNE1 (Potassium voltage-gated channel subfamily E member 1)

References

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- Tyson, J., et al, *Hum. Mol. Genet.* 6 (12), 2179-2185 (1997)
- Schulze-Bahr, E., et al, *Nat. Genet.* 17 (3), 267-268 (1997)
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Clone Number:

Isotype:

Species: Human

Storage and Stability: at -20°C

Storage buffer:

This antibody is stored in PBS, 0.01% sodium azide and 50% glycerol.

Preparation:

Purified by antigen-specific affinity chromatography.

Applications :

ELISA

Western Blotting (1 µg/ml for 2hrs)