

Product Name: KCE1L Rabbit Polyclonal Antibody
Catalog #: APRab12922



Summary

| | |
|------------------------|----------------------------------|
| Production Name | KCE1L Rabbit Polyclonal Antibody |
| Description | Rabbit Polyclonal Antibody |
| Host | Rabbit |
| Application | WB |
| Reactivity | Human,Mouse |

Performance

| | |
|---------------------|--|
| Conjugation | Unconjugated |
| Modification | Unmodified |
| Isotype | IgG |
| Clonality | Polyclonal |
| Form | Liquid |
| Storage | Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles. |
| Buffer | Liquid in PBS containing 50% glycerol, and 0.02% New type preservative N. |
| Purification | Affinity purification |

Immunogen

| | |
|--------------------------|---|
| Gene Name | KCNE1L AMMECR2 |
| Alternative Names | |
| Gene ID | 23630.0 |
| SwissProt ID | Q9UJ90.Synthesized peptide derived from human protein . at AA range: 40-120 |

Application

| | |
|-------------------------|----------------------------------|
| Dilution Ratio | WB 1:500-2000 ELISA 1:5000-20000 |
| Molecular Weight | 15kD |

Background

potassium voltage-gated channel subfamily E regulatory subunit 5(KCNE5) Homo sapiens Voltage-gated potassium (Kv) channels represent the most complex class of voltage-gated ion channels from both functional and structural standpoints.

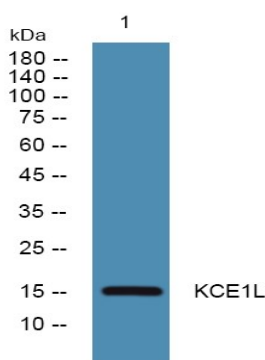
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Their diverse functions include regulating neurotransmitter release, heart rate, insulin secretion, neuronal excitability, epithelial electrolyte transport, smooth muscle contraction, and cell volume. This gene encodes a membrane protein which has sequence similarity to the KCNE1 gene product, a member of the potassium channel, voltage-gated, isk-related subfamily. This intronless gene is deleted in AMME contiguous gene syndrome and may be involved in the cardiac and neurologic abnormalities found in the AMME contiguous gene syndrome. [provided by RefSeq, Jul 2008],disease:Defects in KCNE1L may be a cause of AMME complex [MIM:300194]; also known as Alport syndrome, mental retardation, midface hypoplasia and elliptocytosis, and of additional mild abnormalities of the heart. The AMME complex is a contiguous gene deletion syndrome.,similarity:Belongs to the potassium channel KCNE family.,tissue specificity:Highly expressed in heart, skeletal muscle, brain, spinal cord and placenta.,

Research Area

Image Data



Western blot analysis of lysates from K562 cells, primary antibody was diluted at 1:1000, 4° over night

Note

For research use only.