产品名称: KCE1L Rabbit Polyclonal Antibody

产品货号: APRab12922



# 产品概述 (Summary)

产品名称 (Production Name) KCE1L Rabbit Polyclonal Antibody

描述 (Description) Rabbit polyclonal Antibody

宿主 (Host) Rabbit

应用 (Application) WB,ELISA

种属反应性 (Reactivity) Human,Mouse

## 产品性能 (Performance)

偶联物 (Conjugation)Unconjugated修饰 (Modification)Unmodified

同种型 (Isotype) IgG

克隆 (Clonality) Polyclonal 形式 (Form) Liquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid 存放说明 (Storage)

freeze/thaw cycles.

储存溶液 (Buffer) Liquid in PBS containing 50% glycerol, and 0.02% New type preservative N.

纯化方式 (Purification) Affinity purification

# 免疫原信息 (Immunogen)

基因名 (Gene Name) KCNE1L AMMECR2

别名 (Alternative Names)

基因 ID (Gene ID) 23630.0

Q9UJ90.Synthesized peptide derived from human protein . at AA range: 40-蛋白ID (SwissProt ID)

120

## 产品应用 (Application)

**稀释比 (Dilution Ratio)** WB 1:500-1:2000,ELISA 1:5000-1:20000

蛋白分子量 (Molecular Weight) 15kDa

## 研究背景 (Background)

potassium voltage-gated channel subfamily E regulatory subunit 5(KCNE5) Homo sapiens Voltage-gated potassium (Kv)

Web:https://www.enkilife.cn E-mail:order@enkilife.cn (销售) tech@enkilife.cn (技支持) Tel:027-87002838



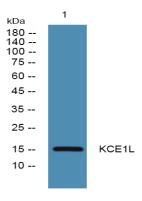
产品货号: APRab12922



channels represent the most complex class of voltage-gated ion channels from both functional and structural standpoints. 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.,

#### 研究领域 (Research Area)

## 图片 (Image Data)



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

#### 注意事项 (Note)

For research use only .

Web:https://www.enkilife.cn E-mail:order@enkilife.cn (销售) tech@enkilife.cn (技术支持) Tel:027-87002838