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Diagnostik & molekulare Diagnostik



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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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KCNE1 (Human) Matched Antibody Pair

Catalog # : H00003753-AP11

規格 : [1 Set]

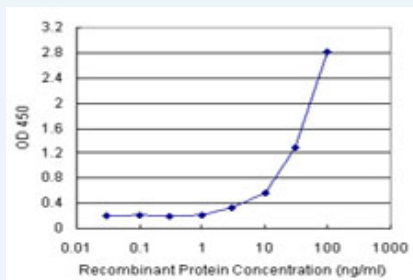
[List All](#)

Specification

Product Description: This antibody pair set comes with matched antibody pair to detect and quantify protein level of human KCNE1.

Reactivity: Human

Quality Control Testing: Standard curve using recombinant protein (H00003753-P01) as an analyte.



Sandwich ELISA detection sensitivity ranging from 3 ng/ml to 100 ng/ml.

Supplied Product: Antibody pair set content:
 1. Capture antibody: rabbit MaxPab® affinity purified polyclonal anti-KCNE1 (100 ug)
 2. Detection antibody: mouse monoclonal anti-KCNE1, IgG1 Kappa (20 ug)
 *Reagents are sufficient for at least 1-2 x 96 well plates using recommended protocols.

Storage Instruction: Store reagents of the antibody pair set at -20°C or lower. Please aliquot to avoid repeated freeze thaw cycle. Reagents should be returned to -20°C storage immediately after use.

MSDS:  [Download](#)

Applications

ELISA Pair (Recombinant protein)

 [Protocol Download](#)

Gene Information

Entrez GeneID: [3753](#)

Gene Name: KCNE1

Gene Alias: FLJ18426, FLJ38123, FLJ94103, ISK, JLNS, JLNS2, LQT2/5, LQT5, MGC33114, MinK

Gene Description: potassium voltage-gated channel, Isk-related family, member 1

Omim ID: [176261](#), [220400](#)

Gene Ontology: [Hyperlink](#)

Application Image

ELISA Pair (Recombinant protein)

Gene Summary: The product of this gene belongs to the potassium channel KCNE family. Potassium ion channels are essential to many cellular functions and show a high degree of diversity, varying in their electrophysiologic and pharmacologic properties. This gene encodes a transmembrane protein known to associate with the product of the KVLQT1 gene to form the delayed rectifier potassium channel. Mutation in this gene are associated with both Jervell and Lange-Nielsen and Romano-Ward forms of long-QT syndrome. Alternatively spliced transcript variants encoding the same protein have been identified. [provided by RefSeq]

Other Designations: lKs producing slow voltage-gated potassium channel subunit beta Mink,OTTHUMP00000108623,OTTHUMP00000108625,OTTHUMP00000108626,cardiac delayed rectifier potassium channel protein,delayed rectifier potassium channel subunit lK,minimal potassium channel,pot

Related Disease

[Arrhythmia](#) [Arrhythmias](#), [Cardiac Atrial Fibrillation](#) [Atrioventricular Block](#) [Brugada Syndrome](#) [Cardiomyopathies](#) [Cardiovascular Diseases](#) [Channelopathies](#) [Chromosome Deletion](#) [Cognition](#) [Death, Sudden](#) [Death, Sudden, Cardiac](#) [Diabetes Mellitus, Type 2](#) [Edema](#) [Gastroparesis](#) [Genetic Predisposition to Disease](#) [Hearing Loss](#) [Hearing Loss, Noise-Induced](#) [Heart Arrest](#)

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