

KCNQ5 Antibody (C-term)

Affinity Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP16776b

Specification

KCNQ5 Antibody (C-term) - Product Information

Application WB,E
Primary Accession O9NR82

Other Accession NP 001153602.1, NP 001153604.1

Reactivity Human, Mouse

Host Rabbit
Clonality Polyclonal
Isotype Rabbit IgG
Calculated MW 102179
Antigen Region 781-809

KCNQ5 Antibody (C-term) - Additional Information

Gene ID 56479

Other Names

Potassium voltage-gated channel subfamily KQT member 5, KQT-like 5, Potassium channel subunit alpha KvLQT5, Voltage-gated potassium channel subunit Kv75, KCNQ5

Target/Specificity

This KCNQ5 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 781-809 amino acids from the C-terminal region of human KCNQ5.

Dilution

WB~~1:1000

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

KCNQ5 Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

KCNQ5 Antibody (C-term) - Protein Information

Name KCNQ5

Function Associates with KCNQ3 to form a potassium channel which contributes to M-type



current, a slowly activating and deactivating potassium conductance which plays a critical role in determining the subthreshold electrical excitability of neurons. Therefore, it is important in the regulation of neuronal excitability. May contribute, with other potassium channels, to the molecular diversity of a heterogeneous population of M-channels, varying in kinetic and pharmacological properties, which underlie this physiologically important current. Insensitive to tetraethylammonium, but inhibited by barium, linopirdine and XE991. Activated by niflumic acid and the anticonvulsant retigabine. As the native M-channel, the potassium channel composed of KCNQ3 and KCNQ5 is also suppressed by activation of the muscarinic acetylcholine receptor CHRM1.

Cellular Location

Cell membrane; Multi-pass membrane protein

Tissue Location

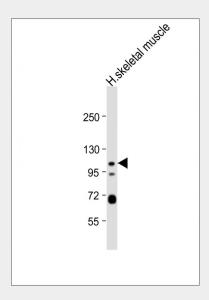
Strongly expressed in brain and skeletal muscle. In brain, expressed in cerebral cortex, occipital pole, frontal lobe and temporal lobe. Lower levels in hippocampus and putamen. Low to undetectable levels in medulla, cerebellum and thalamus

KCNQ5 Antibody (C-term) - Protocols

Provided below are standard protocols that you may find useful for product applications.

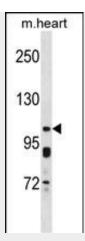
- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- <u>Immunofluorescence</u>
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

KCNQ5 Antibody (C-term) - Images



Anti-KCNQ5 Antibody (C-term) at 1:1000 dilution + human skeletal muscle lysate Lysates/proteins at $20~\mu g$ per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size : 102~kDa Blocking/Dilution buffer: 5% NFDM/TBST.





KCNQ5 Antibody (C-term) (Cat. #AP16776b) western blot analysis in mouse heart tissue lysates (35ug/lane). This demonstrates the KCNQ5 antibody detected the KCNQ5 protein (arrow).

KCNQ5 Antibody (C-term) - Background

This gene is a member of the KCNQ potassium channel gene family that is differentially expressed in subregions of the brain and in skeletal muscle. The protein encoded by this gene yields currents that activate slowly with depolarization and can form heteromeric channels with the protein encoded by the KCNQ3 gene. Currents expressed from this protein have voltage dependences and inhibitor sensitivities in common with M-currents. They are also inhibited by M1 muscarinic receptor activation. Multiple transcript variants encoding different isoforms have been found for this gene.

KCNQ5 Antibody (C-term) - References

Bailey, S.D., et al. Diabetes Care (2010) In press:
Rose, J.E., et al. Mol. Med. 16 (7-8), 247-253 (2010):
Talmud, P.J., et al. Am. J. Hum. Genet. 85(5):628-642(2009)
Roura-Ferrer, M., et al. Cell. Physiol. Biochem. 24 (5-6), 325-334 (2009):
Bal, M., et al. J. Biol. Chem. 283(45):30668-30676(2008)