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Product Information

Anti-Potassium Channel $K_{Ca}1.1$ (BK Ca) (1184-1200)

(Large-conductance Ca^{2+} -activated Potassium Channel); MAXI K^+ ; KCNMA1

Developed in Rabbit, Affinity Isolated Antibody

Product Number **P 4872**

Product Description

Anti-Potassium Channel $K_{Ca}1.1$ (BK Ca) (1184-1200) was developed in rabbit using a synthetic peptide (C)STANRPNRPKSRESRDK corresponding to amino acid residues 1184-1200 of mouse $K_{Ca}1.1$ as the immunogen. This sequence has 100% homology with rat, while human, bovine, chicken and dog have 16/17 residues identical. The antibody was affinity isolated on immobilized immunogen.

Anti-Potassium Channel $K_{Ca}1.1$ (BK Ca) recognizes $K_{Ca}1.1$ from human and rat samples. It has been successfully used in Western blot with rat brain membranes, and in immunohistochemistry with rat brain sections.

The action of potassium (K^+) channels is regulated by voltage, calcium and a variety of neurotransmitters. Each subfamily generally consists of a primary pore forming α subunit that is associated with several regulatory subunits.¹ To date, some 70 different genes that encode the α subunits of K^+ channels have been identified. Recently, the crystal structure of the K^+ channels has been identified.²

The vast family of K^+ channels has been subdivided into the three main subfamilies: the 2 TM, 4 TM and 6 TM K^+ channels.³ The 6 TM family of K^+ channels includes the voltage-gated K^+ (K_V) channels, the KCNQ, the EAG, the calcium-activated K^+ (Slo) subfamily (which is actually a 7 TM not a 6 TM type of channel) and the calcium-activated SK subfamily.

Structurally, the calcium-activated K^+ channels can be divided into two groups: the small or intermediate conductance potassium channels (SK/IK), and the high conductance potassium channels (BK). BK channels are expressed in virtually all cell types where they cause hyperpolarization and help to connect between intracellular Ca^{2+} signaling pathways and membrane

excitability. $K_{Ca}1.1$ channels have a crucial role in smooth muscle contractility, neuronal spike shaping and neurotransmitter release.^{4,5}

Small or intermediate conductance calcium-activated K^+ channels (SK/IK) are responsible for the slow after-hyperpolarization following an action potential. These channels are distinguished from BK channels by their high sensitivity to intracellular calcium, low conductivity and weak or negligible voltage-dependence.⁶ Recently, four SK channels (SK1-4) have been cloned.⁷⁻⁹ SK1 and SK2 channels have highest densities in subregions of the hippocampus and neocortex, while SK3 channels are distributed more diffusely in these brain regions and predominantly expressed in phylogenetically older brain regions.¹⁰ The SK3 channels appear to be implicated in the regulation of electrical excitability and neurotransmitter release of monoaminergic neurons and have been implicated in schizophrenia, ataxia and anorexia nervosa.¹¹ The SK4 (also called IK Ca 1) possess intermediate conductivity and are expressed in many tissue types including red blood cells and T lymphocytes.¹²

Reagent

The antibody is supplied as lyophilized powder from phosphate buffered saline containing 1% bovine serum albumin and 0.025% sodium azide as preservative.

Precautions and Disclaimer

Due to the sodium azide content, a material safety data sheet (MSDS) for this product has been sent to the attention of the safety officer of your institution. Consult the MSDS for information regarding hazards and safe handling.

Preparation Instructions

Reconstitute the lyophilized vial with either 0.05 ml or 0.2 ml deionized water, depending on the package size. Further dilutions should be made using a carrier protein such as BSA (1%).

Storage/Stability

Store at -20 °C. For extended storage, freeze in working aliquots. Avoid repeated freezing and thawing. Storage in "frost-free" freezers is not recommended. Centrifuge before use. Working dilution samples should be discarded if not used within 12 hours.

Product Profile

The recommended working dilution is 1:500 for immunoblotting.

Note: In order to obtain best results in different techniques and preparations we recommend determining optimal working concentration by titration test.

References

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