

Rabbit polyclonal antibody to human KCNN4 (11-28): Affinity purified

Catalogue No.: R-1068-100

Description: THIS PRODUCT IS TEMPORARILY OUT OF STOCK. PLEASE REFER TO THE "REPLACED

BY" FIELD BELOW TO LOCATE THE CURRENT BIOSENSIS PRODUCT TO MEET YOUR RESEARCH NEEDS. KCNN4 is a multi-pass membrane protein and belongs to the potassium channel KCNN family. KCNN4 forms a voltage-independent potassium channel that is activated

by intracellular calcium. The channel is blocked by clotrimazole and charybdotoxin.

Batch No.: See product label

Unit size: 100 ug

Antigen: A synthetic peptide corresponding to a region (11-28) from human KCNN4. To enhance the

immunological response, this peptide was coupled to carrier protein BSA.

Other Names: SK4; KCa4; IKCa1; IK1; Putative Gardos channel; KCNN4; IK1; IKCA1; KCA4; SK4; KCA3.1;

Intermediate conductance calcium-activated potassium channel protein 4;

Accession: O15554 KCNN4_HUMAN;

Produced in: Rabbit

Purity: Affinity purified on antigen column

Applications: Western Blotting (WB). A concentration of 0.1-0.5 ug/mL is recommended for WB. Human

KCNN4 has a predicted length of 427 residues and MW of 48 kDa. Biosensis recommends

optimal dilutions/concentrations should be determined by the end user.

Specificity: The specificity of this antibody has been confirmed by WB against the antigen.

Cross-reactivity: Human; rat; predicted to react with mouse due to sequence homology;

Form: Lyophilised with 5mg BSA, 0.9mg NaCl, 0.2mg Na2HPO4, 0.05mg Thimerosal, 0.05mg NaN3

Reconstitute in 100 uL of sterile distilled water to achieve an antibody concentration of 1

mg/mL. Centrifuge to remove any insoluble material.

Storage: At least 12 months after purchase at 2-8C (lyophilized formulations). After reconstitution,

aliquot and store at -20C for a higher stability. Avoid freeze-thaw cycles

Expiry Date: 12 months after purchase

Specific References: 1. Ruggieri P et al (2012) The inhibition of KCa3.1 channels activity reduces cell motility in

glioblastoma derived cancer stem cells.

PLoS One. 2012;7(10):e47825.