CLC-K (V-15): sc-21295



The Power to Question

BACKGROUND

The family of voltage-dependent chloride channels (CLCs) regulate cellular trafficking of chloride ions, a critical component of all living cells. CLCs regulate excitability in muscle and nerve cells, aid in organic solute transport, and maintain cellular volume. CLC-KA is a kidney-specific chloride channel that mediates transepithelial chloride transport in the thin ascending limb of the Henle loop in the inner medulla. CLC-KA plays a crucial role in urine concentration. The gene encoding human CLC-KA maps to chromosome 1p36.13. Mutations in this gene may be associated with nephrogenic diabetes insipidus in those cases where mutations in the vasopressin V2 receptor and the AQP2 water channel are lacking. CLC-KB mediates basolateral chloride ion efflux in the thick ascending limb and in more distal nephron segments. The gene encoding human CLC-KB maps to chromosome 1p36.13. Mutations in this gene cause type III Barter's syndrome which is characterized by renal saltwasting and low blood pressure.

REFERENCES

- Saito-Ohara, F., Uchida, S., Takeuchi, Y., Sasaki, S., Hayashi, A., Marumo, F. and Ikeuchi, T. 1996. Assignment of the genes encoding the human chloride channels, CLCNKA and CLCNKB, to 1p36 and of CLCN3 to 4q32-q33 by in situ hybridization. Genomics 36: 372-374.
- Simon, D.B., Bindra, R.S., Mansfield, T.A., Nelson-Williams, C., Mendonca, E., Stone, R., Schurman, S., Nayir, A., Alpay, H., Bakkaloglu, A., Rodriguez-Soriano, J., Morales, J.M., Sanjad, S.A., Taylor, C.M., Pilz, D., Brem, A., et al. 1997. Mutations in the chloride channel gene, CLCNKB, cause Bartter's syndrome type III. Nat. Genet. 17: 171-178.
- 3. Matsumura, Y., Uchida, S., Kondo, Y., Miyazaki, H., Ko, S.B., Hayama, A., Morimoto, T., Liu, W., Arisawa, M., Sasaki, S. and Marumo, F. 1999. Overt nephrogenic diabetes insipidus in mice lacking the CLC-K1 chloride channel. Nat. Genet. 21: 95-98.
- Gyömörey, K., Yeger, H., Ackerley, C., Garami, E. and Bear, C.E. 2000. Expression of the chloride channel CLC-2 in the murine small intestine epithelium. Am. J. Physiol. Cell Physiol. 279: C1787-C1794.
- Estevez, R., Boettger, T., Stein, V., Birkenhager, R., Otto, E., Hildebrandt, F. and Jentsch, T.J. 2001. Barttin is a CL⁻ channel β-subunit crucial for renal CL⁻ reabsorption and inner ear K⁺ secretion. Nature 414: 558-561.

CHROMOSOMAL LOCATION

Genetic locus: CLCNKB (human) mapping to 1p36.13; Clcnkb (mouse) mapping to 4 E1.

SOURCE

CLC-K (V-15) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of CLC-KB of rat origin.

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with <0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-21295 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

CLC-K (V-15) is recommended for detection of CLC-KA and CLC-KB of mouse, rat and, to a lesser extent, human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

CLC-K (V-15) is also recommended for detection of CLC-KA and CLC-KB in additional species, including canine and bovine.

Molecular Weight of CLC-K: 63 kDa.

Positive Controls: KNRK whole cell lysate: sc-2214 or NRK whole cell lysate: sc-364197.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat lgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat lgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use donkey anti-goat lgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat lgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

SELECT PRODUCT CITATIONS

 Capasso, G., Rizzo, M., Garavaglia, M.L., Trepiccione, F., Zacchia, M., Mugione, A., Ferrari, P., Paulmichl, M., Lang, F., Loffing, J., Carrel, M., Damiano, S., Wagner, C.A., Bianchi, G. and Meyer, G. 2008. Upregulation of apical sodium-chloride cotransporter and basolateral chloride channels is responsible for the maintenance of salt-sensitive hypertension. Am. J. Physiol. Renal Physiol. 295: F556-F567.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.

Santa Cruz Biotechnology, Inc. 1.800.457.3801 831.457.3801 **Europe** +00800 4573 8000 49 6221 4503 0 **www.scbt.com**