antibodies -online.com





anti-CLCNKB antibody (AA 51-150) (AbBy Fluor® 350)



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|--------|-----|------|-----|
| | N/P | r\/I | i⊢₩ |

| Quantity: | 100 μL |
|----------------------|---|
| Target: | CLCNKB |
| Binding Specificity: | AA 51-150 |
| Reactivity: | Human |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This CLCNKB antibody is conjugated to AbBy Fluor® 350 |
| Application: | Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |

Product Details

| Immunogen: | KLH conjugated synthetic peptide derived from human CLCNKB |
|-----------------------|--|
| Isotype: | IgG |
| Predicted Reactivity: | Human,Mouse,Rat,Dog,Cow,Pig,Rabbit |
| Purification: | Purified by Protein A. |

Target Details

| Target: | CLCNKB |
|-------------------|---|
| Alternative Name: | CLCNKB (CLCNKB Products) |
| Background: | Synonyms: Bartter syndrome type 3, Chloride channel Kb, Chloride channel kidney B, Chloride |

channel protein CIC-Kb, Chloride channel voltage sensitive Kb, CIC K2, CIC-K2, CICK2, CLCKB, CLCKB_HUMAN, CLCNKB, hCIC Kb, hClCKb, MGC24087, OTTHUMP00000011120, OTTHUMP00000011121, RP11 5P18.8.

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. 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. Mutations in this gene cause type III Barter?s syndrome which is characterized by renal salt-wasting and low blood pressure.

Application Details

| Application Notes: | IF(IHC-P) 1:50-200 |
|--------------------|-----------------------|
| | IF(IHC-F) 1:50-200 |
| | IF(ICC) 1:50-200 |
| Restrictions: | For Research Use only |

Handling

| Format: | Liquid |
|--------------------|--|
| Concentration: | 1 μg/μL |
| Buffer: | Aqueous buffered solution containing 0.01M TBS (pH 7.4) with 1 % BSA, 0.03 % Proclin300 and 50 % Glycerol. |
| Preservative: | ProClin |
| Precaution of Use: | This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be handled by trained staff only. |
| Storage: | -20 °C |
| Storage Comment: | Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles. |

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Expiry Date:

12 months