CLCN7 Rabbit Polyclonal Antibody

CAB12422

Product Information

Size:

20uL, 50uL, 100uL, 200uL

Observed MW:

Calculated MW:

86kDa/88kDa

Applications:

IF

Reactivity:

Human

Antibody Information

Recommended dilutions:
IF 1:50 - 1:100

Source:
Rabbit

Isotype:
IgG

Purification:
Affinity purification

Protein Background

The product of this gene belongs to the CLC chloride channel family of proteins. Chloride channels play important roles in the plasma membrane and in intracellular organelles. This gene encodes chloride channel 7. Defects in this gene are the cause of osteopetrosis autosomal recessive type 4 (OPTB4), also called infantile malignant osteopetrosis type 2 as well as the cause of autosomal dominant osteopetrosis type 2 (OPTA2), also called autosomal dominant Albers-Schonberg disease or marble disease autosoml dominant. Osteopetrosis is a rare genetic disease characterized by abnormally dense bone, due to defective resorption of immature bone. OPTA2 is the most common form of osteopetrosis, occurring in adolescence or adulthood.

Immunogen information

Gene ID:
1186

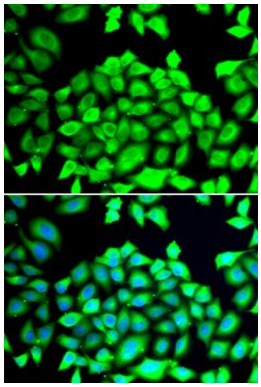
Uniprot
P51798

Synonyms:
CLCN7; CLC-7; CLC7; OPTA2; OPTB4; PPP1R63

Immunogen:
Recombinant fusion protein containing a sequence corresponding to amino acids 626-805 of human CLCN7 (NP_001278.1).

Storage:
Store at -20℃. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

Product Images



Immunofluorescence analysis of HeLa cells using CLCN7 antibody (CAB12422). Blue: DAPI for nuclear staining.

