

**CLC-7 rabbit pAb****Cat#: orb764868 (Manual)**

For research use only. Not intended for diagnostic use.

<b>Product Name</b>	CLC-7 rabbit pAb
<b>Host species</b>	Rabbit
<b>Applications</b>	WB;ELISA
<b>Species Cross-Reactivity</b>	Human;Mouse;Rat
<b>Recommended dilutions</b>	Western Blot: 1/500 - 1/2000. ELISA: 1/40000. Not yet tested in other applications.
<b>Immunogen</b>	The antiserum was produced against synthesized peptide derived from human CLCN7. AA range:10-59
<b>Specificity</b>	CLC-7 Polyclonal Antibody detects endogenous levels of CLC-7 protein.
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide..
<b>Storage</b>	Store at -20°C. Avoid repeated freeze-thaw cycles.
<b>Protein Name</b>	H(+)/Cl(-) exchange transporter 7
<b>Gene Name</b>	CLCN7
<b>Cellular localization</b>	Lysosome membrane ; Multi-pass membrane protein .
<b>Purification</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Clonality</b>	Polyclonal

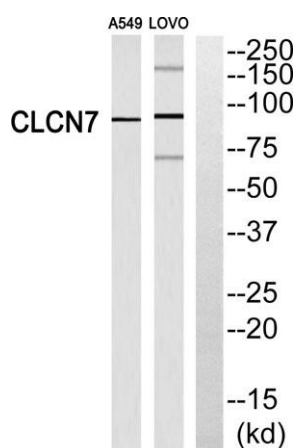
<b>Concentration</b>	1 mg/ml
<b>Observed band</b>	90kD
<b>Human Gene ID</b>	1186
<b>Human Swiss-Prot Number</b>	P51798
<b>Alternative Names</b>	CLCN7; H(+)/Cl(-) exchange transporter 7; Chloride channel 7 alpha subunit; Chloride channel protein 7; CLC-7

### Background

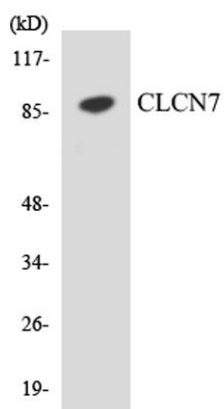
chloride voltage-gated channel 7 (CLCN7) Homo sapiens The product of this gene belongs to the CLC chloride channel family of proteins. Chloride channels play important roles in the plasma membrane and in intracellular organelles. This gene encodes chloride channel 7. Defects in this gene are the cause 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 autosomal 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. [provided by RefSeq, Jul 2008],



**Western Blot analysis of A549 cells using CLC-7 Polyclonal Antibody diluted at 1:500**



Western blot analysis of CLCN7 Antibody. The lane on the right is blocked with the CLCN7 peptide.



Western blot analysis of the lysates from COLO205 cells using CLCN7 antibody.