



This product is for research use only and is not intended for diagnostic use.

## **PRODUCT INFORMATION**

Antigen Description	The CLCN family of voltage-dependent chloride channel genes comprises nine members (CLCN1-7, Ka and Kb) which demonstrate quite diverse functional characteristics while sharing significant sequence homology. The protein encoded by this gene regulates the electric excitability of the skeletal muscle membrane. Mutations in this gene cause two forms of inherited human muscle disorders: recessive generalized myotonia congenita (Becker) and dominant myotonia (Thomsen).
Immunogen	A synthetic peptide of human CLCN1 is used for rabbit immunization.
Isotype	IgG
Source/Host	Rabbit
Species Reactivity	Human
Purification	Protein A
Conjugate	Unconjugated
Applications	Western Blot (Transfected lysate); ELISA
Buffer	In 1x PBS, pH 7.4
Preservative	None
Storage	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

## **GENE INFORMATION**

Gene Name	CLCN1 chloride channel, voltage-sensitive 1 [ Homo sapiens ]
Official Symbol	CLCN1
Synonyms	CLCN1; chloride channel, voltage-sensitive 1; chloride channel 1, skeletal muscle; chloride channel protein 1; CIC 1; CLC1; Thomsen disease; autosomal dominant; clC-1; chloride channel protein, skeletal muscle; MGC138361; MGC142055;
Entrez Gene ID	1180
Protein Refseq	<u>NP_000074</u>
UniProt ID	<u>P35523</u>
Chromosome Location	7q12
Function	chloride channel activity; voltage-gated chloride channel activity; voltage-gated ion channel activity;