

## Product datasheet for **RG226163**

### **CLC7 (CLCN7) (NM\_001114331) Human Tagged ORF Clone**

#### **Product data:**

Product Type:	Expression Plasmids
Product Name:	CLC7 (CLCN7) (NM_001114331) Human Tagged ORF Clone
Tag:	TurboGFP
Symbol:	CLC7
Synonyms:	CLC-7; CLC7; HOD; OPTA2; OPTB4; PPP1R63
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-AC-GFP (PS100010)
E. coli Selection:	Ampicillin (100 ug/mL)



[View online »](#)

**ORF Nucleotide Sequence:**

>RG226163 representing NM\_001114331  
 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC  
 GCC**CGGATCGCC**

ATGGCCAACGTCTCTAAGAAGGTGTCCTGGTCCGGCCGGGACCGGACGACGAGGAGGCGGCCGCTGC  
 TCGGAGGACGGCGCGGCCCGGGGGACCGCGCTGCTGAACGGGGCTGGGCCTGGGGCTGCGCGCCA  
 GGATATGGACCTCCACATCCCTTCCCCAAGGAGATCCCACACAACGAGAAGCTCCTGTCCCTCAAGTAT  
 GAGAGCTTGGACTATGACAACAGTGAGAACCAGCTGTTCTGGAGGAGGAGCGGCGGATCAATCACACGG  
 CCTTCCGACGGTGGAGATCAAGCGCTGGGTCACTGCGCCCTCATTGGGATCCTCACGGCCTCGTGCC  
 CTGCTTCAATGACATCGTGGTGGAAAACCTGGCTGGCCTCAAGTACAGGGTCAACAGGGCAATATCGAC  
 AAGTTCACAGAGAAGGGCGGACTGTCCTTCTCCCTGTTGCTGTGGCCACGCTGAACGCCGCTTCGTGC  
 TCGTGGGCTCTGTGATTGTGGCTTTCATAGAGCCGGTGGCTGCTGGCAGCGGAATCCCCAGATCAAGT  
 CTTCTCAACGGGGTGAAGATCCCCACGTGGTGGGCTCAAGACGTTGGTGATCAAAGTGTCCGGTGTG  
 ATCCTGTCCGTGGTGGGGGCTGGCCGTGGGAAAGGAAGGGCCGATGATCCACTCAGGTTCAAGTATTG  
 CCGCCGGGATCTCTCAGGGAAGGTCAACGTCACTGAAACGAGATTTCAAGATCTTCGAGTACTTCGCGAG  
 AGACACAGAGAAGCGGGACTTTCGTCTCCGACGGGGCTGCGGCCGAGTGTGAGCGGCTTTGGAGCCCC  
 GTGGGTGGGGCTCTGTTAGCTTGGAGGAGGTGCGTCTTCTGGAACGAGTTCCTGACCTGGAGGATCT  
 TCTTTGCTCCATGATCTCCACGTTACCCCTGAATTTGTTCTGAGCATTACCACGGGAACATGTGGGA  
 CCTGTCCAGCCCAGGCCATCAACTTCGGAAGGTTTACTCGGAGAAAATGGCCTACACGATCCACGAG  
 ATCCCGGCTTTCATCGCCATGGCGTGGTGGGCGGTGTGCTTGGAGCAGTGTCAATGCCTTGAAGTACT  
 GGCTGACCATGTTTGAATCAGTACATCCACCGCCCTGCCTGCAGGTGATTGAGGCCGCTGTGGTGGC  
 CGCGTACCGCCACAGTTGCCTTCGTGCTGATCTACTCGTCGCGGATTGCCAGCCCTGCAGGGGGC  
 TCCATGTCTACCCGCTGCAGCTCTTTGTGCAGATGGCGAGTAACTCCATGGCTGCGGCCTTCTTCA  
 ACACCCGGAGAAGAGCGTGGTGGAGCTTCCACGACCCGCCAGGCTCTACAACCCCTGACCCTCGG  
 CCTGTTACGCTGGTCTACTTCTTCTGGCCTGCTGGACCTACGGGCTCACGGTGTCTGCCGGGGTCTTC  
 ATCCCGTCCCTGCTCATCGGGGCTGCTGGGGCCGGCTTTGGGATCTCCCTGTCTACCTCACGGGG  
 CGGCGATCTGGGCGACCCCGCAAATACGCCCTGATGGGAGTGTGCCAGCTGGGCGGATTGTGCG  
 GATGACACTGAGCCTGACCGTCATCATGATGGAGGCCACCAGCAACGTGACCTACGGCTTCCCCATCATG  
 CTGGTGTGATGACCGCAAGATCGTGGGCGACGCTTTCATTGAGGGCCTGTACGACATGCACATTCAGC  
 TGCAGAGTGTGCCCTTCCCTGCACTGGGAGGCCCGGTACCTCACACTCACTCACTGCCAGGAGGTGAT  
 GAGCACACAGTACCTGCCTGAGGCGGCGTGAGAAGGTGCGGCTCATTGTGGACGTGTGAGCGACAG  
 GCGTCCAATCACAAACGGCTTCCCGTGGTGGAGCATGCCGATGACACCCAGCCTGCCCGGCTCCAGGGCC  
 TGATCCTGCGCTCCCAGCTCATCGTTCTCCTAAAGCACAAAGTGTGTTGGAGCGGTCCAACCTGGGCT  
 GGTACAGCGGCGCTGAGGCTGAAGGACTTCCGAGACGCTACCCGCGCTTCCACCCATCCAGTCCATC  
 CACGTGTCCAGGACGAGCGGGAGTGCACCATGGACCTCTCCGAGTTCATGAACCCCTCCCCTACACGG  
 TGCCCCAGGAGGCGTCTGCTCCACGGGTGTTCAAGCTGTTCCGGGCCCTGGGCTGCGGCACCTGGTGGT  
 GGTGGACAACCGCAATCAGGTTGTGGGTTGGTGACCAGGAAGGACCTCGCCAGGTACCGCTGGGAAAG  
 AGAGGCTTGGAGGAGCTCTGCTGGCCAGACG

**ACGCGT**ACGCGGCCGCTCGAG – GFP Tag – GTTTAA

**Protein Sequence:** >RG226163 representing NM\_001114331  
 Red=Cloning site Green=Tags(s)

```

MANVSKKVSWSGRDRDDEEAAPLLRRTARPGGGTPLLNGAGPGAARQMDPPHPFKEIPHNEKLLSLKY
ESLDYDENSENQLFEEERRINHTAFRTVEIKRWVICALIGILTGLVACFIDIVVENLAGLYRVIKGNID
KFTEKGLSFLSLLLWATLNAFVLVGSVIVAFIEPVAAGSGIPQIKCFLNGVKIPHVRLKTLVIKVS
VILSVVGGGLAVGKEGPMIHSVIAAGISQGRSTSLKRFKIFEFYFRDTEKRFVSAAGAAAGVSAAGF
VGGVLFSLLEEGASFVNQFLTWRIFASFMISTFTLNFVLSIYHGMMWDLSSPGLINFRDSEKMA
YTIHEIPVFIAMGVVGGVVGAVFNALNYWLTMFRIRYIHRPCLQVIEAVLVAAVTATVAFVLIYSS
RDCQPLQGGSMSYPLQLFCADGEYNSMAAAFFNTPEKSVVSLFHDPGSGYNPLTLGLFTLVYFFL
ACWYGLTVSAGVFIPLSLLIGAAWGRFLGISLSYLTGAAIWADPGKYALMGAAQLGGIVRMTLSL
TVIMMEATSNVTYGFPIMLVLMTAKIVGDVFI EGLYDMHIQLQSVPFHWEAPVTSLSLTAREVMST
PVTCLRRREKVGIVDVLSDTASNHNHGFVVEHADDTQPARLQGLILRSQLIVLLKHKVFVERSNL
GLVQRRRLKDFRDYAPRFPPIQSIHVSQDERECTMDLSEFMNPSPYTPQEASLPRVFKLFRALGLR
HLVVVDNRNQVGLVTRKDLARYRLGKRGLEELSLAQT
  
```

TRTRPLE - GFP Tag - V

**Restriction Sites:** SgfI-MluI

**Cloning Scheme:**



**ACCN:** NM\_001114331

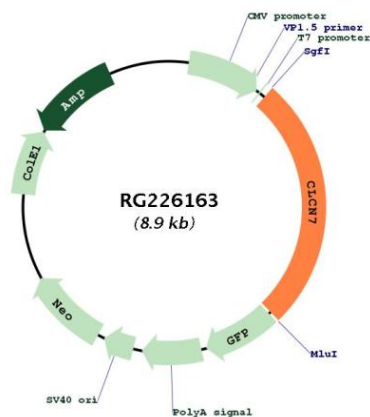
**ORF Size:** 2343 bp

**OTI Disclaimer:** The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. [More info](#)

**OTI Annotation:** This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.

<b>Components:</b>	The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).
<b>Reconstitution Method:</b>	<ol style="list-style-type: none"> <li>1. Centrifuge at 5,000xg for 5min.</li> <li>2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.</li> <li>3. Close the tube and incubate for 10 minutes at room temperature.</li> <li>4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.</li> <li>5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.</li> </ol>
<b>RefSeq:</b>	<u>NM_001114331.2, NP_001107803.1</u>
<b>RefSeq Size:</b>	4164 bp
<b>RefSeq ORF:</b>	2346 bp
<b>Locus ID:</b>	1186
<b>UniProt ID:</b>	<u>P51798</u>
<b>Cytogenetics:</b>	16p13.3
<b>Protein Families:</b>	Druggable Genome, Ion Channels: Other, Transmembrane
<b>Gene Summary:</b>	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]

**Product images:**



Circular map for RG226163

