

Product datasheet for **SC113923**

CLN6 (NM_017882) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	CLN6 (NM_017882) Human Untagged Clone
Tag:	Tag Free
Symbol:	CLN6
Synonyms:	CLN4A; HsT18960; nclf
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL5</u>
E. coli Selection:	Ampicillin (100 ug/mL)



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Fully Sequenced ORF:

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>OriGene sequence for NM_017882 edited
GAATTCGGCACGAGGGGGCAGCCGAGCTGAACCGTCTTCTCCTCGGAAAGGCAGGGCCG
AGGGGCTGCGGGGCAGCCATGGAGGCGACGCGGAGGGCGGCAGCACCTGGGAGCGACGGG
CGGCCCAGGCGCGCAGCTGGGCGCTCCTTCTGCAGGCCAGGCATGGCTCTGTGAGCGC
TGATGAGGCTGCCCGACGGCTCCCTTCCACCTCGACCTCTGGTTCTACTTCACACTGCA
GAACTGGGTTCTGGACTTTGGGCGTCCCATTGCCATGCTGGTATTCCTCTCGAGTGGTT
TCCACTCAACAAGCCAGTGTGGGGACTACTTCCACATGGCCTACAACGTCATCACGCC
CTTTCTTGTCAAGCTCATCGAGCGGTCCCCCGCACCTGCCACGCTCCATCACGTA
CGTGAGCATCATCATCTTTCATCATGGGTGCCAGCATCCACCTGGTGGGTGACTCTGTCAA
CCACCGCTGCTCTTCACTGGCTACCAGCACCACTGTCTGTCCGTGAGAACCCCATCAT
CAAGAATCTCAAGCCGGAGACGCTGATCGACTCCTTTGAGCTGCTCTACTATTATGATGA
GTACCTGGGTCACTGCATGTGGTACATCCCCTTCTTCTCATCCTCTTTCATGTACTTCAG
CGGCTGCTTACTGCCTTAAAGCTGAGAGCTTGATTCCAGGGCCTGCCCTGCTCCTGGT
GGCACCCAGTGGCTGTACTACTGGTACCTGGTACCGAGGGCCAGATCTTCATCCTCTT
CATCTTACCTTCTTCGCCATGTGGCCCTCGCTGCACCAGAAGCGCAAGCGCCTCTT
CCTGGACAGCAACGGCCTTTCCTTCTCCTCCTTCGCACTGACCCTCTTGCTGTGGC
GCTCTGGGTGCGCTGGCTGTGGAATGACCCTGTTCTCAGGAAGAAGTACCCGGGTGCAT
CTACGTCCCTGAGCCCTGGGCTTTTACACCCCTCACGTGAGCAGTGGCCTGAGTCCC
TGGCACCAAGGCTCTGGCCTCTGCTGGGTGGGAGGGTGGGCCATGGAGGGCATCTGAATA
CAGGAGTAGGGGGGGTGTGGGTGTGAACCAGAGACCGAGAGCATGAGTGGGGTGTGCCT
CGTGTGCGTGGATTGCTGTGTGTGTGTGTCTTGTATATGTGTGCGCAGAGTGCATCAT
TTTCAGACTCTACTATTTCCGTCAAGTTTCTGTTTGATTTGGATCATCTCAGGATCGGAT
TCTGTTTTAGAGTGTCTGCGCCAGGATCCGGGCCCTGCCCTCCTCTGCACCTGACCA
CACTCCCTACTCAGGGCTAGTCTGTTCTTCCCGGACATCTTCTGGTAGCGGTGCAGGAGA
GGGCTGGGTGGGCGAGAGGCCAGGAGGGGACCTGGTGTGTACCTGCCACCCACCTGGCT
CATCCCTCAGGCCACCCTGACCCTACATTACATAGGTTACGTGACGCTACTGTGGCTGT
TGAGCAAAGCATTTCTCCTTTCTGGCCTCATTGCACTAGATGGGCCTGTGGTCCAAAG
TAGGTGAGTAGGTTGGGTTGCTGACACCCCTGGGTGACGCTTTGGGACAGATGAGTGG
CTCTGCTCTGCACTGCCCTCTCCCTGCCTGGGGGCTATGTGCACTCCAGACCCCTGCC
AGGCTCAGGCCCATGAGGTATGGAGACACCCTGGCCCCAGGAGCTGGAGGCACCGCCCA
CTCCCTGGCATTCCAGCTTTCAGGTGACCCCTCTCTACCCAAAGCTCTGTCCCCTGC
TCCCACTCCAGAAGAAGTGCAGGACGCTTTCGGGCGAGCCTAGCCACAGGCTTTGAGCGC
CTGCATTCCTGGGGGCTGGAGGGTGGGGTGCCAAAGGCCCTGAGCAAAAGCCAGAGCTCC
TCTCATCAAAGCCTTTACAAGGTGCTGGGCCAGAGGCTTTGCCCTTGACAGAGTGGCCCA
GGGTTTCAAGGGAGGAGGAACCTCCCCCTACCTAGGACCTTCTGTGGGGGGTCTACAG
AGTCAGGGACAGAAGGGAAGGGACCCACAGGAAGTCACAGTGGTGGCCAGGGATGTGTCA
GCCCCAGCCACGGGGACGCGGATTCAAGAATGAAGTAAATACAGTCACAGCCCCAAAA
AAAAAAAAAAAAAACTCGAC
    
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5' Read Nucleotide Sequence:	<p>>OriGene 5' read for NM_017882 unedited</p> <pre>GGGTGTTTAGATTTGTATACGACTCACTATAGGCGGCCGGAATCGGCACGAGGGGGCAG CCGAGCTGAACCGGTCTCTTCTCGAAGGCAGGGCCGAGGGGCTGCGGGGCAGCCATG GAGGCGACGCGGAGGCGGCAGCACCTGGGAGCGACGGGCGGCCAGGCGCGCAGCTGGGC GCCTCCTTCTGCAGGCCAGGCATGGCTCTGTGAGCGCTGATGAGGCTGCCCCACGGCT CCCTTCCACCTCGACCTCTGGTTCTACTTCACACTGCAGAAGTGGGTTCTGGACTTTGGG CGTCCCATTGCCATGCTGGTATTCCCTCTCGAGTGGTTTTCCACTCAACAAGCCAGTGT GGGGACTACTTCCACATGGCCTACAACGTCATCACGCCCTTTCTTTGCTCAAGCTCATC GAGCGGTCCCCCGCACCTGCCACGCTCCATCACGTACGTGAGCATCATCATCTTCATC ATGGGTGCCAGCATCCACCTGGTGGTGACTCTGTCAACCACCGCTGCTTTCAGTGGC TACCAGCACCACTGTCTGTCCGTGAGAACCCCATCATCAAGAATCTCAAGCCGGAGACG CTGATCGACTCCTTTGAGCTGCTCTACTATTATGATGAGTACCTGGGTCAGTGCATGTGG TACATCCCCTTCTTCTCATCTTTCATGTACTTCAGCGGCTGCTTTACTGCCTCTAAA GCTGAGAGCTTGATTCCAGGGCCTGCCCTGCTCCTGGTGGCACCCAGTGGCCTGTACTAC TGGTACCTGGTCAACGAGGCCAGATCTTCATCCTTTCATCTTACCTTCTCGCCATG CTGGCCCTCGTCTGCACCAAAAGCGCAAGCGCCTCTTCTCTG</pre>
3' Read Nucleotide Sequence:	<p>>OriGene 3' read for NM_017882 unedited</p> <pre>GTACCAAAGCGGCNCTCNAGGTNCGAGAAAAAATCTTTTTTTTTTGGGGCTGTGACTGTT TTTACTTCATTCTTGAATCCCGCGTCCCGTGGCTGGGGGCTGACACATCCCTGGGCACC ACTGTGACTTCTGTGGGTCCCTCCCTTCTGTCCCTGACTCTGTAGACCCCCACAGGA AGGGTCTTAGGTAGGGGAGGGTCTTCTCCTCNTAAACCCTGGGCCACTCTGTCAAGGC AAAGCCTCTGGGCCAGCACCTTGTAAGGCTTTGATGAGAGGAGCTCTGGCTTTTGCTC AGGGCCTTTGGCACCCACCTCCAGCCCCAGGAATGCAGGCGCTCAAAGCCTGTGGCT AGGCTGCCGAAGCACGTGCCCGAGTTCTTCTGGAGTGGGAGCAGGGGGACAGAGCTTTG GGTAGAGGAGGGTACCTGCAAAGCTGGAATGCCAGGGGAGTGGGCGGTGCCTCCAGCTC CTGGGGGGCAGGGTGTCTCCATACCTCATGGGCCTGAGCTGGGCAGGGGTCTGGAGTGC ACATAGCCCCAGGCAGGGAGAGGGCAGTGACAGGACAGAGCCACTCATCTGTCCCAAAG CTGACCCAAAGGGGTGTGAGCAACCCCAACCTACTGACCTACTTTGGACCACAGGCCAT GTAGTGCAAATGAGGCCCAAGGAGAAATGCTCTGCTCAACAGCCACAGTAGGCTGAC GTAACCTATGTAATGTATGGTCAAGGTGGGCCCTGAAGGATGAGCCACGTGGTGGGCAGG TGACACACCAGGTCCCCCTCTGGGCCTTTGGCCACCCAGCCCTCTCCTGCACGTTAC CANAAGATGTCCCGGAAGACAGACTACCCTGAGTTAGGAGTGTGGCCAAAGTGACAA GAGGGCCAGGGCCCCGATCCTGCCACAACACTCTTAAAN</pre>
Restriction Sites:	NotI-NotI
ACCN:	NM_017882
Insert Size:	2350 bp
OTI Disclaimer:	Our molecular clone sequence data has been matched to the reference identifier above as a point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative RNA splicing form or single nucleotide polymorphism (SNP).
Components:	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).

Reconstitution Method:	<ol style="list-style-type: none">1. Centrifuge at 5,000xg for 5min.2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.3. Close the tube and incubate for 10 minutes at room temperature.4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.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.
RefSeq:	NM_017882.1 , NP_060352.1
RefSeq Size:	2227 bp
RefSeq ORF:	936 bp
Locus ID:	54982
UniProt ID:	Q9NWW5
Cytogenetics:	15q23
Protein Families:	Transmembrane
Gene Summary:	<p>This gene is one of eight which have been associated with neuronal ceroid lipofuscinoses (NCL). Also referred to as Batten disease, NCL comprises a class of autosomal recessive, neurodegenerative disorders affecting children. The genes responsible likely encode proteins involved in the degradation of post-translationally modified proteins in lysosomes. The primary defect in NCL disorders is thought to be associated with lysosomal storage function. [provided by RefSeq, Oct 2008]</p>