

## Product datasheet for **SC322900**

### CLDN19 (NM\_001123395) Human Untagged Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	CLDN19 (NM_001123395) Human Untagged Clone
Tag:	Tag Free
Symbol:	CLDN19
Synonyms:	HOMG5
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
Fully Sequenced ORF:	>SC322900 representing NM_001123395. Blue=Insert sequence Red=Cloning site Green=Tag(s)

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GCTCGTTTAGTGAACCGTCAGAATTTTGTAAACGACTCACTATAGGGCGCCGGGAATTCGTCGACTG
GATCCGGTACCGAGGAGATCTGCCGCCCGGATCGCC
ATGGCCAACTCAGGCCTCCAGCTCCTGGGCTACTTCTTGGCCCTGGGTGGCTGGGTGGGCATCATTGCT
AGCACAGCCCTGCCACAGTGAAGCAGTCTTCTACGCAGGCGACGCCATCATCACTGCCGTGGGCCTC
TATGAAGGGCTCTGGATGTCTGCGCCTCCAGAGCACTGGGCAAGTGCAGTGAAGCTCTACGACTCG
CTGCTCGCCCTGGACGGTCACATCCAATCAGCGCGGGCCCTGATGGTGGTGGCCGTGCTCCTGGGCTTC
GTGGCCATGGTCTCAGCGTAGTTGGCATGAAGTGTACGCGGGTGGGAGACAGCAACCCATTGCCAAG
GGCCGTGTTGCCATCGCCGGGGAGCCCTTTCATCCTGGCAGGCCTCTGCACTTTGACTGCTGTCTCG
TGGTATGCCACCCCTGGTACCCAGGAGTTCTTCAACCAAGCACACCTGTCAATGCCAGGTATGAATTT
GGCCAGCCCTGTTCTGGTGGGCTGGGCTCAGCTGGCCTGGCCGTGCTGGGCGGCTCCTTCTCTGCTGC
ACATGCCCGGAGCCAGAGAGACCAACAGCAGCCACAGCCCTATCGGCCTGGACCTCTGCTGCTGCC
CGAGAGTACGCTGA
ACGCGTACGCGGCCGCTCGAGCAGAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGAT
TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC
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Restriction Sites:	Sgfl-MluI
Plasmid Map:	<input type="checkbox"/>
ACCN:	NM_001123395
Insert Size:	636 bp



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<b>OTI Disclaimer:</b>	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).
<b>OTI Annotation:</b>	This TrueClone is provided through our Custom Cloning Process that includes sub-cloning into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.
<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><a href="#">NM_001123395.1</a></u>
<b>RefSeq Size:</b>	3602 bp
<b>RefSeq ORF:</b>	636 bp
<b>Locus ID:</b>	149461
<b>UniProt ID:</b>	<u><a href="#">Q8N6F1</a></u>
<b>Cytogenetics:</b>	1p34.2
<b>Protein Families:</b>	Transmembrane
<b>Protein Pathways:</b>	Cell adhesion molecules (CAMs), Leukocyte transendothelial migration, Tight junction
<b>MW:</b>	22.1 kDa
<b>Gene Summary:</b>	<p>The product of this gene belongs to the claudin family. It plays a major role in tight junction-specific obliteration of the intercellular space, through calcium-independent cell-adhesion activity. Defects in this gene are the cause of hypomagnesemia renal with ocular involvement (HOMGO). HOMGO is a progressive renal disease characterized by primary renal magnesium wasting with hypomagnesemia, hypercalciuria and nephrocalcinosis associated with severe ocular abnormalities such as bilateral chorioretinal scars, macular colobomata, significant myopia and nystagmus. Alternatively spliced transcript variants encoding distinct isoforms have been identified for this gene. [provided by RefSeq, Jun 2010]</p> <p>Transcript Variant: This variant (2) contains an additional segment in the coding region compared to variant 1. The resulting isoform (b) contains a shorter and distinct C-terminus compared to isoform a.</p>