

Product datasheet for **SC326387**

PCDH15 (NM_001142770) Human Untagged Clone

Product data:

Product Type: Expression Plasmids
Product Name: PCDH15 (NM_001142770) Human Untagged Clone
Tag: Tag Free
Symbol: PCDH15
Synonyms: CDHR15; DFNB23; USH1F
Vector: pCMV6 series
Fully Sequenced ORF: >NCBI ORF sequence for NM_001142770, the custom clone sequence may differ by one or more nucleotides

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ATGTTTCGACAGTTTTATCTCTGGACATGTTTAGCTTCAGGGATCATCCTGGGCTCTCTC
TTTGAAATCTGCTTGGGCCAGTATGATGATGATTGCAAAGTAGCTAGGGGAGGACCACCA
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AATGACACCTTTGAAATCCCCTAATGTTGACTGGAAATATAGTGTAAAGGAAGAGGCTC
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CTGAATGAGAGGGCAACCACCACCACCTCTCACAGTGGATGTTCTGGATGGAGATGAC
TTGGGTCCAATGTTTCTCCTTGTGTCTTGTGCCAAACACTCGTGATTGCCGTCCACTC
ACTTATCAAGCTGCCATACCTGAGTTGAGAACTCCGGAAGAAGTGAACCCATTATTGTT
ACGCCACCAATCCAAGCATTGATCAGGACCGGAATATTCAACCGCCATCAGATAGGCCA
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CATCCTAGGACAGCAGAAGTCTCCTGGAGCCAGTAAACAGAGACTTTCACCAGAAA
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CAAGGCTATATCCTGGAATCTGCCCCAGTGGGAGCAACCATTTCCGACAGTCTCAATTTG
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CGTACCTCACCTTACTTCAACAGTGGACAGGGAAGAAGCAAACTTACACCTTTTCCG
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GACATGAGACCTGGGACAGTGTACATACAGTCACTGCAGTCGACGCAGACGAAGGGTCA
AATGGGGAGATCACATATGAAATCCTTGTGGGGCTCAGGGAGACTTCATCATCAATAAA
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CTCACGGTCCAAGCAGCGGATAATGCTCCTCCTGCAGAGCGAAGGAACTCCATCTGCACT
 GTGTATATTGAAGTGCTCCACCAAATAATCAAAGCCCTCCTCGCTTCCCACAGCTGATG
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 GCCTTCATCATATCCTCTGCTGCATTCTGCCATCTTGGTGGTTTTGGTCAGCTACAGA
 CAGTTTAAAGTACGTCAAGCTGAGTGTACAAAGACTGCACGAATTCAGGCCGCTTACCC
 GCGGCTAAACCAGCAGTCCCGCTCCTGCACCAGTGGCAGCGCCCCGCCGCCGCCGCG
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 ATGGGTATGAAATGCCTCAATATGGGAGTCCCGTCGATTGTTACCACCAGCTGGACAGG
 AGGAATATGGTGAAGTGGTGGTGAAGCTGAGGAAGAATATGAGGAGGAAGAGGAAGAGC
 CAAAGAAAATTAACAAAGGTTGAAATTAGAGAGCCTAGTGAGGAGGAAGAAG

Restriction Sites:

Please inquire

ACCN:

NM_001142770

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).
OTI Annotation:	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.
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:	<u>NM_001142770.1</u> , <u>NP_001136242.1</u>
RefSeq Size:	5803 bp
RefSeq ORF:	4620 bp
Locus ID:	65217
Cytogenetics:	10q21.1
Protein Families:	Druggable Genome, Transmembrane
Gene Summary:	<p>This gene is a member of the cadherin superfamily. Family members encode integral membrane proteins that mediate calcium-dependent cell-cell adhesion. It plays an essential role in maintenance of normal retinal and cochlear function. Mutations in this gene result in hearing loss and Usher Syndrome Type 1F (USH1F). Extensive alternative splicing resulting in multiple isoforms has been observed in the mouse ortholog. Similar alternatively spliced transcripts are inferred to occur in human, and additional variants are likely to occur. [provided by RefSeq, Dec 2008]</p> <p>Transcript Variant: This variant (J) lacks an alternate in-frame exon in the 5' coding region and has a distinct 3' splice pattern, compared to variant A. The resulting isoform (CD2-2) lacks a 5-aa segment in the 5' coding region and has a distinct and longer C-terminus, compared to isoform CD1-1. Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments.</p>