

## **Product datasheet for SC209613**

## PCDH15 (NM 001142766) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

Product Name: PCDH15 (NM\_001142766) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: PCDH15

Synonyms: CDHR15; DFNB23; USH1F

**ACCN:** NM\_001142766

**Insert Size:** 789 bp

Insert Sequence: >SC209613 3'UTR clone of NM\_001142766

The sequence shown below is from the reference sequence of NM\_001142766. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

TTATAATGAAAATAAATCTTGAGTCGTTGA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

**Restriction Sites:** Sgfl-Mlul

**OTI Disclaimer:** Our molecular clone sequence data has been matched to the sequence identifier above as a

point of reference. Note that the complete sequence of this clone is largely the same as the

reference sequence but may contain minor differences, e.g., single nucleotide

polymorphisms (SNPs).



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## PCDH15 (NM\_001142766) Human 3' UTR Clone - SC209613

**Components:** The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

RefSeq: <u>NM 001142766.2</u>

**Summary:** 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 IF (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]

**Locus ID:** 65217 **MW:** 30.7