

## Product datasheet for RC230017L4V

#### OriGene Technologies, Inc.

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### Cadherin like 23 (CDH23) (NM 001171932) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

Product Name: Cadherin like 23 (CDH23) (NM 001171932) Human Tagged ORF Clone Lentiviral Particle

Symbol: Cadherin like 23

Synonyms: CDHR23; PITA5; USH1D

Mammalian Cell

Selection:

Puromycin

**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

**ACCN:** NM\_001171932

ORF Size: 1218 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC230017).

Sequence:
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.

**RefSeq:** <u>NM 001171932.1</u>, <u>NP 001165403.1</u>

 RefSeq ORF:
 1221 bp

 Locus ID:
 64072

 UniProt ID:
 Q9H251

 Cytogenetics:
 10q22.1

**Protein Families:** Transmembrane

MW: 45 kDa





# Cadherin like 23 (CDH23) (NM\_001171932) Human Tagged ORF Clone Lentiviral Particle – RC230017L4V

#### **Gene Summary:**

This gene is a member of the cadherin superfamily, whose genes encode calcium dependent cell-cell adhesion glycoproteins. The encoded protein is thought to be involved in stereocilia organization and hair bundle formation. The gene is located in a region containing the human deafness loci DFNB12 and USH1D. Usher syndrome 1D and nonsyndromic autosomal recessive deafness DFNB12 are caused by allelic mutations of this cadherin-like gene. Upregulation of this gene may also be associated with breast cancer. Alternative splice variants encoding different isoforms have been described. [provided by RefSeq, May 2013]