

## Product datasheet for RC221533L4V

## OriGene Technologies, Inc.

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## OTOL1 (NM\_001080440) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** OTOL1 (NM\_001080440) Human Tagged ORF Clone Lentiviral Particle

Symbol: OTOL1

Synonyms: C1QTNF15; C1QTNF16

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM 001080440

ORF Size: 1431 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC221533).

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:** NM 001080440.1, NP 001073909.1

 RefSeq Size:
 1434 bp

 RefSeq ORF:
 1434 bp

 Locus ID:
 131149

 UniProt ID:
 A6NHN0

 Cytogenetics:
 3q26.1

**Protein Families:** Transmembrane

MW: 49.2 kDa







## **Gene Summary:**

This gene encodes a secreted glycoprotein with a C-terminal complement Cq1-like globular domain that belongs to the C1q/tumor necrosis factor-related protein (CTRP) family. The encoded protein is expressed in the inner ear and forms a multimeric complex called the otoconia, together with cerebellin-1 and otoconin-90, as part of the otoconial membrane. It contains extensive posttranslational modifications including hydroxylated prolines and glycosylated lysines. Naturally occurring mutations in this gene are associated with abnormal otoconia formation and balance deficits resulting from vestibular dysfunction. [provided by RefSeq, Jul 2017]