

## Product datasheet for **RG240261**

### OTOG (NM\_001292063) Human Tagged ORF Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	OTOG (NM_001292063) Human Tagged ORF Clone
Tag:	TurboGFP
Symbol:	OTOG
Synonyms:	DFNB18B; MLEMP; OTGN
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-AC-GFP (PS100010)
E. coli Selection:	Ampicillin (100 ug/mL)
ORF Nucleotide Sequence:	>RG240261 representing NM_001292063. Blue=ORF Red=Cloning site Green=Tag(s)

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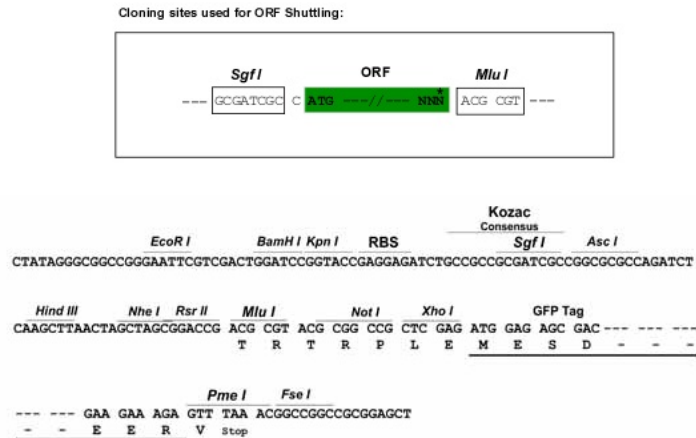
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**Protein Sequence:**

>Peptide sequence encoded by RG240261  
 Blue=ORF Red=Cloning site Green=Tag(s)

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Restriction Sites: Sgfl-Mlul

**Cloning Scheme:**


**ACCN:** NM\_001292063

**ORF Size:** 8739 bp

**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.

**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).

**RefSeq:** [NM\\_001292063.2](#)

**RefSeq Size:** 8742 bp

**RefSeq ORF:** 8742 bp

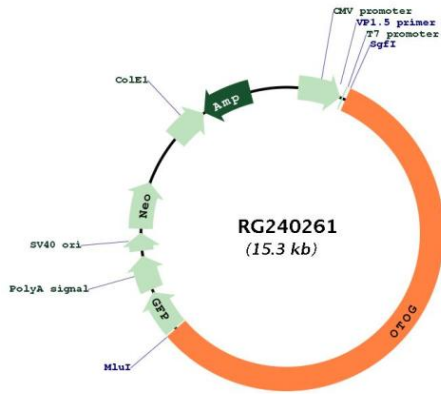
**Locus ID:** 340990

**Cytogenetics:** 11p15.1

**MW:** 313.8 kDa

**Gene Summary:** The protein encoded by this gene is a component of the acellular membranes of the inner ear. Disruption of the orthologous mouse gene shows that it plays a role in auditory and vestibular functions. It is involved in fibrillar network organization, the anchoring of otoconial membranes and cupulae to the neuroepithelia, and likely in sound stimulation resistance. Mutations in this gene cause autosomal recessive nonsyndromic deafness, type 18B. Alternative splicing of this gene results in multiple transcript variants. [provided by RefSeq, May 2014]

Product images:



Circular map for RG240261