

## Product datasheet for RC214967L2V

## OriGene Technologies, Inc.

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## **ROR1 (NM\_005012) Human Tagged ORF Clone Lentiviral Particle**

## **Product data:**

**Product Type:** Lentiviral Particles

**Product Name:** ROR1 (NM\_005012) Human Tagged ORF Clone Lentiviral Particle

Symbol: ROR<sup>2</sup>

**Synonyms:** dJ537F10.1; NTRKR1

Mammalian Cell

Selection:

None

**Vector:** pLenti-C-mGFP (PS100071)

Tag: mGFP

**ACCN:** NM\_005012 **ORF Size:** 2811 bp

**ORF Nucleotide** 

2011 00

Sequence:
OTI Disclaimer:

The ORF insert of this clone is exactly the same as(RC214967).

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.

**RefSeg:** NM 005012.1

 RefSeq Size:
 3358 bp

 RefSeq ORF:
 2814 bp

 Locus ID:
 4919

 UniProt ID:
 Q01973

 Cytogenetics:
 1p31.3

**Domains:** KR, FRI, pkinase, TyrKc, S\_TKc, ig, IGc2, IG

**Protein Families:** Druggable Genome, Protein Kinase, Transmembrane





MW: 104.1 kDa

**Gene Summary:** 

This gene encodes a receptor tyrosine kinase-like orphan receptor that modulates neurite growth in the central nervous system. The encoded protein is a glycosylated type I membrane protein that belongs to the ROR subfamily of cell surface receptors. It is a pseudokinase that lacks catalytic activity and may interact with the non-canonical Wnt signalling pathway. This gene is highly expressed during early embryonic development but expressed at very low levels in adult tissues. Increased expression of this gene is associated with B-cell chronic lymphocytic leukaemia. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Jun 2012]