

## Product datasheet for RC214259L4V

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## Gliomedin (GLDN) (NM\_181789) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: Gliomedin (GLDN) (NM 181789) Human Tagged ORF Clone Lentiviral Particle

Symbol: Gliomedir

Synonyms: CLOM; COLM; CRG-L2; CRGL2; LCCS11; UNC-112; UNC-122

**Mammalian Cell** 

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_181789 **ORF Size:** 1653 bp

**ORF Nucleotide** 

1000 00

Sequence:

The ORF insert of this clone is exactly the same as(RC214259).

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.

**RefSeg:** NM 181789.2

 RefSeq Size:
 5084 bp

 RefSeq ORF:
 1656 bp

 Locus ID:
 342035

 UniProt ID:
 Q6ZMI3

 Cytogenetics:
 15q21.2

**Protein Families:** Transmembrane

MW: 58.8 kDa







## **Gene Summary:**

This gene encodes a protein that contains olfactomedin-like and collagen-like domains. The encoded protein, which exists in both transmembrane and secreted forms, promotes formation of the nodes of Ranvier in the peripheral nervous system. Mutations in this gene cause a form of lethal congenital contracture syndrome in human patients. Autoantibodies to the encoded protein have been identified in sera form patients with multifocal motor neuropathy. [provided by RefSeq, May 2017]