

## Product datasheet for RC201969L2V

## 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

## PLOD1 (NM\_000302) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: PLOD1 (NM 000302) Human Tagged ORF Clone Lentiviral Particle

Symbol: PLOD1

**Synonyms:** EDS6; EDSKCL1; LH; LH1; LLH; PLOD

Mammalian Cell

Selection:

None

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

Tag: mGFP

ACCN: NM\_000302

ORF Size: 2181 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC201969).

OTI Disclaimer:

Sequence:

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 000302.2

 RefSeq Size:
 3047 bp

 RefSeq ORF:
 2184 bp

 Locus ID:
 5351

 UniProt ID:
 Q02809

 Cytogenetics:
 1p36.22

Domains: 20G-Fell\_Oxy, P4Hc
Protein Families: Druggable Genome





## PLOD1 (NM\_000302) Human Tagged ORF Clone Lentiviral Particle - RC201969L2V

**Protein Pathways:** Lysine degradation

MW: 83.6 kDa

**Gene Summary:** Lysyl hydroxylase is a membrane-bound homodimeric protein localized to the cisternae of

the endoplasmic reticulum. The enzyme (cofactors iron and ascorbate) catalyzes the hydroxylation of lysyl residues in collagen-like peptides. The resultant hydroxylysyl groups are attachment sites for carbohydrates in collagen and thus are critical for the stability of intermolecular crosslinks. Some patients with Ehlers-Danlos syndrome type VI have

deficiencies in lysyl hydroxylase activity. Two transcript variants encoding different isoforms

have been found for this gene. [provided by RefSeq, Oct 2015]