

## Product datasheet for RC207610L1V

## OriGene Technologies, Inc.

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## PLOD2 (NM\_182943) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** PLOD2 (NM\_182943) Human Tagged ORF Clone Lentiviral Particle

Symbol: PLOD2

Synonyms: BRKS2; LH2; TLH

Mammalian Cell

Selection:

None

**Vector:** pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK
ACCN: NM 182943

ORF Size: 2274 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC207610).

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 182943.2

 RefSeq Size:
 4072 bp

 RefSeq ORF:
 2277 bp

 Locus ID:
 5352

 UniProt ID:
 000469

 Cytogenetics:
 3q24

**Protein Pathways:** Lysine degradation

**MW:** 87.1 kDa







## **Gene Summary:**

The protein encoded by this gene is a membrane-bound homodimeric enzyme that is localized to the cisternae of the rough 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 VIB have deficiencies in lysyl hydroxylase activity. Mutations in the coding region of this gene are associated with Bruck syndrome. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Jul 2008]