

## Product datasheet for RC205277L3V

## OriGene Technologies, Inc.

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## POMT1 (NM\_001077365) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** POMT1 (NM\_001077365) Human Tagged ORF Clone Lentiviral Particle

Symbol: POMT

Synonyms: LGMD2K; LGMDR11; MDDGA1; MDDGB1; MDDGC1; RT

**Mammalian Cell** 

Selection:

Puromycin

**Vector:** pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK

**ACCN:** NM\_001077365

ORF Size: 2175 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC205277).

Sequence:

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.

**RefSeq:** <u>NM 001077365.1</u>, <u>NP 001070833.1</u>

 RefSeq Size:
 3080 bp

 RefSeq ORF:
 2178 bp

 Locus ID:
 10585

 UniProt ID:
 Q9Y6A1

 Cytogenetics:
 9q34.13

**Protein Families:** Transmembrane

**Protein Pathways:** O-Mannosyl glycan biosynthesis





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**MW:** 82.4 kDa

**Gene Summary:** The protein encoded by this gene is an O-mannosyltransferase that requires interaction with

the product of the POMT2 gene for enzymatic function. The encoded protein is found in the membrane of the endoplasmic reticulum. Defects in this gene are a cause of Walker-Warburg syndrome (WWS) and limb-girdle muscular dystrophy type 2K (LGMD2K). Several transcript variants encoding different isoforms have been found for this gene.[provided by RefSeq, Oct

2008]