

## Product datasheet for RC222985L1V

### 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

# SPTLC1 (NM\_006415) Human Tagged ORF Clone Lentiviral Particle

#### **Product data:**

**Product Type:** Lentiviral Particles

**Product Name:** SPTLC1 (NM\_006415) Human Tagged ORF Clone Lentiviral Particle

Symbol: SPTLC1

Synonyms: HSAN1; HSN1; LBC1; LCB1; SPT1; SPTI

Mammalian Cell

Selection:

None

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

Tag: Myc-DDK
ACCN: NM 006415

ORF Size: 1419 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC222985).

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.

**RefSeg:** NM 006415.2

 RefSeq Size:
 2780 bp

 RefSeq ORF:
 1422 bp

 Locus ID:
 10558

 UniProt ID:
 015269

 Cytogenetics:
 9q22.31

**Domains:** aminotran\_1\_2

**Protein Families:** Druggable Genome, Transmembrane





#### SPTLC1 (NM\_006415) Human Tagged ORF Clone Lentiviral Particle - RC222985L1V

**Protein Pathways:** Metabolic pathways, Sphingolipid metabolism

**MW:** 52.6 kDa

**Gene Summary:** This gene encodes a member of the class-II pyridoxal-phosphate-dependent

aminotransferase family. The encoded protein is the long chain base subunit 1 of serine palmitoyltransferase. Serine palmitoyltransferase converts L-serine and palmitoyl-CoA to 3-oxosphinganine with pyridoxal 5'-phosphate and is the key enzyme in sphingolipid biosynthesis. Mutations in this gene were identified in patients with hereditary sensory

neuropathy type 1. Alternatively spliced variants encoding different isoforms have been identified. Pseudogenes of this gene have been defined on chromosomes 1, 6, 10, and 13.

[provided by RefSeq, Jul 2013]