

Product datasheet for RC204131L3V

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

SLC33A1 (NM_004733) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: SLC33A1 (NM_004733) Human Tagged ORF Clone Lentiviral Particle

Symbol: SLC33A1

Synonyms: ACATN; AT-1; AT1; CCHLND; SPG42

Mammalian Cell

Selection:

Puromycin

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

Tag: Myc-DDK
ACCN: NM 004733

ORF Size: 1647 bp

ORF Nucleotide

The ODE

Sequence:

The ORF insert of this clone is exactly the same as(RC204131).

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 accurring variations (e.g. polymorphisms), each with its own valid existence. This

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 004733.2

 RefSeq Size:
 3811 bp

 RefSeq ORF:
 1650 bp

 Locus ID:
 9197

 UniProt ID:
 000400

Cytogenetics: 3q25.31

Protein Families: Transmembrane

Protein Pathways: Glycosphingolipid biosynthesis - ganglio series, Metabolic pathways





SLC33A1 (NM_004733) Human Tagged ORF Clone Lentiviral Particle - RC204131L3V

MW: 60.9 kDa

Gene Summary: The protein encoded by this gene is required for the formation of O-acetylated (Ac)

gangliosides. The encoded protein is predicted to contain 6 to 10 transmembrane domains, and a leucine zipper motif in transmembrane domain III. Defects in this gene have been reported to cause spastic paraplegia autosomal dominant type 42 (SPG42) in one Chinese family, but not in similar patients of European descent. Two transcript variants encoding the

same protein have been found for this gene. [provided by RefSeq, Jul 2010]