

Product datasheet for RC211663L4V

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

ST3GAL5 (NM_001042437) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: ST3GAL5 (NM_001042437) Human Tagged ORF Clone Lentiviral Particle

Symbol: ST3GAL5

Synonyms: SATI; SIAT9; SIATGM3S; SPDRS; ST3Gal V; ST3GalV

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

ACCN: NM_001042437

ORF Size: 1185 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC211663).

OTI Disclaimer:

Cytogenetics:

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 001042437.1

 RefSeq Size:
 2262 bp

 RefSeq ORF:
 1188 bp

 Locus ID:
 8869

 UniProt ID:
 Q9UNP4

Protein Families: Transmembrane

2p11.2

Protein Pathways: Glycosphingolipid biosynthesis - ganglio series, Metabolic pathways





ORIGENE

MW: 45.4 kDa

Gene Summary:

Ganglioside GM3 is known to participate in the induction of cell differentiation, modulation of cell proliferation, maintenance of fibroblast morphology, signal transduction, and integrinmediated cell adhesion. The protein encoded by this gene is a type II membrane protein which catalyzes the formation of GM3 using lactosylceramide as the substrate. The encoded protein is a member of glycosyltransferase family 29 and may be localized to the Golgi apparatus. Mutation in this gene has been associated with Amish infantile epilepsy syndrome. Transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul 2008]