

Product datasheet for RC220291L3V

OriGene Technologies, Inc.

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ST3GAL3 (NM_174967) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: ST3GAL3 (NM_174967) Human Tagged ORF Clone Lentiviral Particle

Symbol: ST3GAL3

Synonyms: DEE15; EIEE15; MRT12; SIAT6; ST3GALII; ST3Gal III; ST3GalIII; ST3N

Mammalian Cell

Selection:

Puromycin

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

Tag: Myc-DDK

ACCN: NM_174967

ORF Size: 465 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC220291).

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 174967.2

 RefSeq Size:
 1653 bp

 RefSeq ORF:
 468 bp

 Locus ID:
 6487

 UniProt ID:
 Q11203

Cytogenetics: 1p34.1

Protein Families: Secreted Protein, Transmembrane





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Protein Pathways: Glycosphingolipid biosynthesis - lacto and neolacto series, Keratan sulfate biosynthesis,

Metabolic pathways

MW: 17.4 kDa

Gene Summary: The protein encoded by this gene is a type II membrane protein that catalyzes the transfer of

sialic acid from CMP-sialic acid to galactose-containing substrates. The encoded protein is normally found in the Golgi apparatus but can be proteolytically processed to a soluble form. This protein is a member of glycosyltransferase family 29. Mutations in this gene have been associated with a form of autosomal recessive nonsymdromic cognitive disability as well as infantile epileptic encephalopathy. Multiple transcript variants encoding several different

isoforms have been found for this gene. [provided by RefSeq, Jul 2017]