

Product datasheet for RC204670L2V

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

SLC35A1 (NM_006416) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: SLC35A1 (NM 006416) Human Tagged ORF Clone Lentiviral Particle

Symbol: SLC35A1

Synonyms: CDG2F; CMPST; CST; hCST

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_006416 **ORF Size:** 1011 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC204670).

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 006416.2</u>

 RefSeq Size:
 1918 bp

 RefSeq ORF:
 1014 bp

 Locus ID:
 10559

 UniProt ID:
 P78382

 Cytogenetics:
 6q15

Domains: Nuc_sug_transp

Protein Families: Transmembrane





ORÏGENE

MW: 36.8 kDa

Gene Summary: The protein encoded by this gene is found in the membrane of the Golgi apparatus, where it

transports nucleotide sugars into the Golgi. One such nucleotide sugar is CMP-sialic acid, which is imported into the Golgi by the encoded protein and subsequently glycosylated. Defects in this gene are a cause of congenital disorder of glycosylation type 2F (CDG2F). Two transcript variants encoding different isoforms have been found for this gene.[provided by

RefSeq, Dec 2009]