

## Product datasheet for RC202452L2V

## OriGene Technologies, Inc.

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## **GM2A (NM\_000405) Human Tagged ORF Clone Lentiviral Particle**

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** GM2A (NM\_000405) Human Tagged ORF Clone Lentiviral Particle

Symbol: GM2A

**Synonyms:** GM2-AP; SAP-3

Mammalian Cell

Selection:

None

**Vector:** pLenti-C-mGFP (PS100071)

Tag: mGFP

**ACCN:** NM\_000405

ORF Size: 579 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC202452).

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 000405.3

 RefSeq Size:
 3690 bp

 RefSeq ORF:
 582 bp

 Locus ID:
 2760

 UniProt ID:
 P17900

 Cytogenetics:
 5q33.1

Domains: ML

**Protein Families:** Druggable Genome





## GM2A (NM\_000405) Human Tagged ORF Clone Lentiviral Particle - RC202452L2V

Protein Pathways: Lysosome

MW: 20.8 kDa

Gene Summary: This gene encodes a small glycolipid transport protein which acts as a substrate specific co-

factor for the lysosomal enzyme beta-hexosaminidase A. Beta-hexosaminidase A, together with GM2 ganglioside activator, catalyzes the degradation of the ganglioside GM2, and other molecules containing terminal N-acetyl hexosamines. Mutations in this gene result in GM2-gangliosidosis type AB or the AB variant of Tay-Sachs disease. Alternative splicing results in

multiple transcript variants. [provided by RefSeq, Nov 2009]