

OriGene Technologies, Inc.

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Product datasheet for RC202452L1V

GM2A (NM_000405) Human Tagged ORF Clone Lentiviral Particle

Product data:

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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-Myc-DDK (PS100064)
Tag:	Myc-DDK
ACCN:	NM_000405
ORF Size:	579 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC202452).
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. <u>More info</u>
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 000405.3</u>
RefSeq Size:	3690 bp
RefSeq ORF:	582 bp
Locus ID:	2760
UniProt ID:	<u>P17900</u>
Cytogenetics:	5q33.1
Domains:	ML
Protein Families:	Druggable Genome



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	GM2A (NM_000405) Human Tagged ORF Clone Lentiviral Particle – RC202452L1V
Protein Pathway	s: 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]

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