

## OriGene Technologies, Inc.

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## Product datasheet for RC201684L4V

## SACM1L (NM\_014016) Human Tagged ORF Clone Lentiviral Particle

## **Product data:**

Product Type:	Lentiviral Particles
Product Name:	SACM1L (NM_014016) Human Tagged ORF Clone Lentiviral Particle
Symbol:	SACM1L
Synonyms:	SAC1
Mammalian Cell Selection:	Puromycin
Vector:	pLenti-C-mGFP-P2A-Puro (PS100093)
Tag:	mGFP
ACCN:	NM_014016
ORF Size:	1761 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC201684).
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 014016.2</u>
RefSeq Size:	3550 bp
RefSeq ORF:	1764 bp
Locus ID:	22908
UniProt ID:	<u>Q9NTJ5</u>
Cytogenetics:	3p21.31
Domains:	Syja_N
Protein Families:	Druggable Genome, Transmembrane



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	SACM1L (NM_014016) Human Tagged ORF Clone Lentiviral Particle – RC201684L4V
MW:	66.8 kDa
Gene Summary:	This gene encodes an integral membrane protein, which is localized to the endoplasmic reticulum, and functions as a phosphoinositide phosphatase that hydrolyzes phosphatidylinositol 3-phosphate, phosphatidylinositol 4-phosphate, and phosphatidylinositol 3,5-bisphosphate. Deletion of this gene in mouse results in preimplantation lethality. Other studies suggest that this gene is also involved in the organization of golgi membranes and mitotic spindles. Alternatively spliced transcript variants have been found for this gene. A C-terminally extended isoform is also predicted to be produced by the use of an alternative in-frame, downstream translation termination codon via a stop codon readthrough mechanism.[provided by RefSeq, Dec 2017]

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