

## Product datasheet for RC203275L2V

## 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

## SAT1 (NM\_002970) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: SAT1 (NM 002970) Human Tagged ORF Clone Lentiviral Particle

Symbol: SAT1

Synonyms: DC21; KFSD; KFSDX; SAT; SSAT; SSAT-1

Mammalian Cell

Selection:

None

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

Tag: mGFP

**ACCN:** NM\_002970

ORF Size: 513 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC203275).

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 002970.1

 RefSeq Size:
 1085 bp

 RefSeq ORF:
 516 bp

 Locus ID:
 6303

 UniProt ID:
 P21673

 Cytogenetics:
 Xp22.11

**Domains:** Acetyltransf

**Protein Families:** Druggable Genome





## SAT1 (NM\_002970) Human Tagged ORF Clone Lentiviral Particle - RC203275L2V

**Protein Pathways:** Arginine and proline metabolism, Metabolic pathways

MW: 20 kDa

**Gene Summary:** The protein encoded by this gene belongs to the acetyltransferase family, and is a rate-

limiting enzyme in the catabolic pathway of polyamine metabolism. It catalyzes the

acetylation of spermidine and spermine, and is involved in the regulation of the intracellular

concentration of polyamines and their transport out of cells. Defects in this gene are associated with keratosis follicularis spinulosa decalvans (KFSD). Alternatively spliced

transcripts have been found for this gene.[provided by RefSeq, Sep 2009]