

## Product datasheet for RC204729L3V

## OriGene Technologies, Inc.

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## delta Sarcoglycan (SGCD) (NM 000337) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** delta Sarcoglycan (SGCD) (NM\_000337) Human Tagged ORF Clone Lentiviral Particle

**Symbol:** delta Sarcoglycan

Synonyms: 35DAG; CMD1L; DAGD; LGMDR6; SG-delta; SGCDP; SGD

**Mammalian Cell** 

Selection:

ACCN:

Puromycin

**Vector:** pLenti-C-Myc-DDK-P2A-Puro (PS100092)

NM 000337

Tag: Myc-DDK

ORF Size: 870 bp

**ORF Nucleotide** 

Sequence:

The ORF insert of this clone is exactly the same as(RC204729).

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 000337.4

 RefSeq Size:
 9805 bp

 RefSeq ORF:
 873 bp

 Locus ID:
 6444

 UniProt ID:
 Q92629

**Cytogenetics:** 5q33.2-q33.3 **Domains:** sarcoglycan

**Protein Families:** Transmembrane





## delta Sarcoglycan (SGCD) (NM\_000337) Human Tagged ORF Clone Lentiviral Particle – RC204729L3V

Protein Pathways: Arrhythmogenic right ventricular cardiomyopathy (ARVC), Dilated cardiomyopathy,

Hypertrophic cardiomyopathy (HCM), Viral myocarditis

MW: 32.2 kDa

**Gene Summary:** The protein encoded by this gene is one of the four known components of the sarcoglycan

complex, which is a subcomplex of the dystrophin-glycoprotein complex (DGC). DGC forms a link between the F-actin cytoskeleton and the extracellular matrix. This protein is expressed most abundantly in skeletal and cardiac muscle. Mutations in this gene have been associated with autosomal recessive limb-girdle muscular dystrophy and dilated cardiomyopathy. Alternatively spliced transcript variants encoding distinct isoforms have been observed for

this gene. [provided by RefSeq, Jul 2008]