

Product datasheet for RC219081L4V

OriGene Technologies, Inc.

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CACNA2D1 (NM_000722) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: CACNA2D1 (NM_000722) Human Tagged ORF Clone Lentiviral Particle

Symbol: CACNA2D1

Synonyms: CACNA2; CACNL2A; CCHL2A; LINC01112; lncRNA-N3

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

ACCN: NM_000722 **ORF Size:** 3273 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC219081).

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 000722.2

RefSeq Size: 3822 bp
RefSeq ORF: 3276 bp

Locus ID: 781

 UniProt ID:
 P54289

 Cytogenetics:
 7q21.11

Domains: VWA, Cache

Protein Families: Druggable Genome, Ion Channels: Other





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

cardiomyopathy, Hypertrophic cardiomyopathy (HCM), MAPK signaling pathway

MW: 123 kDa

Gene Summary: The preproprotein encoded by this gene is cleaved into multiple chains that comprise the

alpha-2 and delta subunits of the voltage-dependent calcium channel complex. Calcium channels mediate the influx of calcium ions into the cell upon membrane polarization. Mutations in this gene can cause cardiac deficiencies, including Brugada syndrome and short QT syndrome. Alternate splicing results in multiple transcript variants, some of which may

lack the delta subunit portion. [provided by RefSeq, Nov 2014]