

OriGene Technologies, Inc.

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Product datasheet for RC219081L3V

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-Myc-DDK-P2A-Puro (PS100092) |
| Tag: | Myc-DDK |
| ACCN: | NM_000722 |
| ORF Size: | 3273 bp |
| ORF Nucleotide Sequence: | The ORF insert of this clone is exactly the same as(RC219081). |
| 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 000722.2</u> |
| RefSeq Size: | 3822 bp |
| RefSeq ORF: | 3276 bp |
| Locus ID: | 781 |
| UniProt ID: | <u>P54289</u> |
| Cytogenetics: | 7q21.11 |
| Domains: | VWA, Cache |
| Protein Families: | Druggable Genome, Ion Channels: Other |



This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2022 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US 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]

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