

## Product datasheet for RC211493L1V

## 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

## **CACNB2 (NM 201596) Human Tagged ORF Clone Lentiviral Particle**

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** CACNB2 (NM\_201596) Human Tagged ORF Clone Lentiviral Particle

Symbol:

CAB2; CACNLB2; CAVB2; MYSB Synonyms:

**Mammalian Cell** 

Selection:

None

Vector: pLenti-C-Myc-DDK (PS100064)

Myc-DDK Tag: NM 201596 ACCN: **ORF Size:** 1980 bp

**ORF Nucleotide** 

OTI Disclaimer:

Sequence:

The ORF insert of this clone is exactly the same as(RC211493).

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.

RefSeq: NM 201596.1

RefSeq Size: 3445 bp RefSeq ORF: 1983 bp

Locus ID: 783

**UniProt ID:** Q08289

Cytogenetics: 10p12.33-p12.31

**Protein Families:** Druggable Genome, Ion Channels: Other





## CACNB2 (NM\_201596) Human Tagged ORF Clone Lentiviral Particle - RC211493L1V

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

cardiomyopathy, Hypertrophic cardiomyopathy (HCM), MAPK signaling pathway

**MW:** 73.4 kDa

**Gene Summary:** This gene encodes a subunit of a voltage-dependent calcium channel protein that is a

member of the voltage-gated calcium channel superfamily. The gene product was originally identified as an antigen target in Lambert-Eaton myasthenic syndrome, an autoimmune disorder. Mutations in this gene are associated with Brugada syndrome. Alternatively spliced variants encoding different isoforms have been described. [provided by RefSeq, Feb 2013]