

## Product datasheet for RC207778L3V

## 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

## SCN2B (NM\_004588) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** SCN2B (NM\_004588) Human Tagged ORF Clone Lentiviral Particle

Symbol: SCN2B
Synonyms: ATFB14

Mammalian Cell

Puromycin

Selection:

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

Tag:Myc-DDKACCN:NM\_004588

ORF Size: 645 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC207778).

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 004588.3

 RefSeq Size:
 4922 bp

 RefSeq ORF:
 648 bp

 Locus ID:
 6327

 UniProt ID:
 060939

 Cytogenetics:
 11q23.3

**Protein Families:** Druggable Genome, Ion Channels: Sodium, Transmembrane

MW: 24.3 kDa







**Gene Summary:** 

The protein encoded by this gene is the beta 2 subunit of the type II voltage-gated sodium channel. The encoded protein is involved in cell-cell adhesion and cell migration. Defects in this gene can be a cause of Brugada Syndrome, atrial fibrillation, or sudden infant death syndrome. [provided by RefSeq, Jul 2015]