

Product datasheet for RC209565L3V

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

SCN1B (NM_001037) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: SCN1B (NM_001037) Human Tagged ORF Clone Lentiviral Particle

Symbol: SCN1B

Synonyms: ATFB13; BRGDA5; DEE52; EIEE52; GEFSP1

Mammalian Cell

Selection:

Puromycin

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

Tag: Myc-DDK
ACCN: NM 001037

ORF Size: 654 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC209565).

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.

RefSeq: <u>NM 001037.3</u>

 RefSeq Size:
 1521 bp

 RefSeq ORF:
 657 bp

 Locus ID:
 6324

 UniProt ID:
 Q07699

 Cytogenetics:
 19q13.11

Domains: ig

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





ORIGENE

MW: 24.5 kDa

Gene Summary:

Voltage-gated sodium channels are heteromeric proteins that function in the generation and propagation of action potentials in muscle and neuronal cells. They are composed of one alpha and two beta subunits, where the alpha subunit provides channel activity and the beta-1 subunit modulates the kinetics of channel inactivation. This gene encodes a sodium channel beta-1 subunit. Mutations in this gene result in generalized epilepsy with febrile seizures plus, Brugada syndrome 5, and defects in cardiac conduction. Multiple transcript variants encoding different isoforms have been found for this gene.[provided by RefSeq, Oct 2009]