

### Product datasheet for RC213549L2V

#### 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

## Dystrobrevin alpha (DTNA) (NM 032980) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** Dystrobrevin alpha (DTNA) (NM\_032980) Human Tagged ORF Clone Lentiviral Particle

Symbol: Dystrobrevin alpha

Synonyms: D18S892E; DRP3; DTN; DTN-A; LVNC1

**Mammalian Cell** 

Selection:

None

**Vector:** pLenti-C-mGFP (PS100071)

Tag: mGFP

**ACCN:** NM\_032980 **ORF Size:** 1173 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC213549).

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 032980.2

 RefSeq Size:
 5221 bp

 RefSeq ORF:
 1176 bp

 Locus ID:
 1837

 UniProt ID:
 Q9Y4J8

 Cytogenetics:
 18q12.1

**Protein Families:** Druggable Genome

MW: 43.4 kDa





# Dystrobrevin alpha (DTNA) (NM\_032980) Human Tagged ORF Clone Lentiviral Particle – RC213549L2V

#### **Gene Summary:**

The protein encoded by this gene belongs to the dystrobrevin subfamily of the dystrophin family. This protein is a component of the dystrophin-associated protein complex (DPC), which consists of dystrophin and several integral and peripheral membrane proteins, including dystroglycans, sarcoglycans, syntrophins and alpha- and beta-dystrobrevin. The DPC localizes to the sarcolemma and its disruption is associated with various forms of muscular dystrophy. Mutations in this gene are associated with left ventricular noncompaction with congenital heart defects. Multiple alternatively spliced transcript variants encoding different isoforms have been identified for this gene. [provided by RefSeq, Jul 2008]