

Product datasheet for RC213549L1V

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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

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

NM 032980

Mammalian Cell

Selection:

ACCN:

None

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

Myc-DDK Tag:

ORF Size: 1173 bp

ORF Nucleotide

Sequence:

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

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: 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





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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]