

Product datasheet for **SC323009**

Dystrobrevin alpha (DTNA) (NM_001128175) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	Dystrobrevin alpha (DTNA) (NM_001128175) Human Untagged Clone
Tag:	Tag Free
Symbol:	DTNA
Synonyms:	D18S892E; DRP3; DTN; DTN-A; LVNC1
Vector:	<u>pCMV6 series</u>
Fully Sequenced ORF:	>NCBI ORF sequence for NM_001128175, the custom clone sequence may differ by one or more nucleotides ATGATTGAAGATAGTGGGAAAAGAGGAAATACCATGGCAGAAAGAAGACAGCTGTTTGCA GAGATGAGGGCTCAAGATCTGGATCGCATCCGACTCTCCACCTACAGAACAGCATGCAAG CTTAGGTTTGTTCAGAAGAAATGCAATTTGCACCTGGTGGACATATGGAATGTCATAGAA GCATTGCGGGAAAATGCTCTGAACAACCTGGACCCAAACACTGAACTCAACGTGTCCCGC TTAGAGGCTGTGCTCTCCACTATTTTTACCAGCTCAACAAACGGATGCCAACCACTCAC CAAATCCATGTGGAGCAGTCCATCAGCCTCCTCCTTAACCTCCTGCTTGCAGCGTTTGAT CCGGAAGGCCATGGTAAATTTAGTATTTGCTGTCAAAATGGCTTTAGCCACATTGTGT GGAGGGAAGATCATGGACAAATTAAGATATATTTTCTCAATGATTTCTGACTCCAGTGGG GTGATGGTTTATGGACGATATGACCAATTCCTTCGGGAAGTTCTCAAACCTACCCACGGCA GTTTTTGAAGGTCCTTCATTTGGTTACACAGAACAGTCAGCCAGATCCTGTTTCTCCAA CAGAAAAAGTCACGTTAAATGGTTTCTTGACACGCTTATGTCAGATCCTCCCCCGCAG TGTCTGGTCTGGTTGCCTCTTCTGCATCGACTAGCAAATGTGGAAAAATGCTTCCATCCG GTTGAGTGTTCTACTGCCACAGTGAGAGTATGATGGGATTTTCGCTACCGATGCCAACAG TGTCACAATTACCAGCTCTGTCAGGACTGCTTCTGGAGGGGACATGCCGGTGGTTTCTCAT AGCAACCAGCACCAATGAAAGAGTACACGTCATGGAAATCACCTGCTAAGAAGCTGACT AATGCATTAAGCAAGTCCCTGAGCTGTGCTTCCAGCCGTGAACCTTTGCACCCCATGTTT CCAGATCAGCCTGAGAAGCCACTCAACTTGGCTCACATCGTGCCTCCAGACCTGTAACC AGCATGAACGACACCCTGTTCTCCACTCTGTTCCCTCCTCAGGAAGTCTTTTATTACC AGGAGCTCGGACGGTGTCTTTGGTGGATGCGTC
Restriction Sites:	Please inquire
ACCN:	NM_001128175
OTI Disclaimer:	Our molecular clone sequence data has been matched to the reference identifier above as a point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative RNA splicing form or single nucleotide polymorphism (SNP).



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OTI Annotation:	This TrueClone is provided through our Custom Cloning Process that includes sub-cloning into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.
Components:	The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).
Reconstitution Method:	<ol style="list-style-type: none">1. Centrifuge at 5,000xg for 5min.2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.3. Close the tube and incubate for 10 minutes at room temperature.4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.
RefSeq:	<u>NM_001128175.1, NP_001121647.1</u>
RefSeq Size:	1728 bp
RefSeq ORF:	1116 bp
Locus ID:	1837
UniProt ID:	<u>Q9Y4J8</u>
Cytogenetics:	18q12.1
Protein Families:	Druggable Genome
Gene Summary:	<p>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]</p> <p>Transcript Variant: This variant (9) lacks an internal coding exon and multiple exons from the 3' end, and contains an alternate 3' exon, compared to variant 1. The resulting isoform (9) lacks an internal segment and has a much shorter and distinct C-terminus, as compared to isoform 1.</p>