

Product datasheet for SC325682

OriGene Technologies, Inc.

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alpha Sarcoglycan (SGCA) (NM_001135697) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: alpha Sarcoglycan (SGCA) (NM_001135697) Human Untagged Clone

Tag: Tag Free Symbol: SGCA

Synonyms: 50DAG; adhalin; ADL; DAG2; DMDA2; LGMD2D; LGMDR3; SCARMD1

Vector: pCMV6 series

Fully Sequenced ORF: >NCBI ORF sequence for NM_001135697, the custom clone sequence may differ by one or

more nucleotides

GACCAGCAC

Restriction Sites: Please inquire **ACCN:** NM 001135697

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).

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:

- 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: NM 001135697.1, NP 001129169.1

 RefSeq Size:
 1069 bp

 RefSeq ORF:
 792 bp

 Locus ID:
 6442

 UniProt ID:
 Q16586

 Cytogenetics:
 17q21.33

Protein Families: Druggable Genome, Transmembrane

Protein Pathways: Arrhythmogenic right ventricular cardiomyopathy (ARVC), Dilated cardiomyopathy,

Hypertrophic cardiomyopathy (HCM), Viral myocarditis

Gene Summary: This gene encodes a component of the dystrophin-glycoprotein complex (DGC), which is

critical to the stability of muscle fiber membranes and to the linking of the actin cytoskeleton to the extracellular matrix. Its expression is thought to be restricted to striated muscle.

Mutations in this gene result in type 2D autosomal recessive limb-girdle muscular dystrophy. Multiple transcript variants encoding different isoforms have been found for this gene.

[provided by RefSeq, Oct 2008]

Transcript Variant: This variant (2) lacks two alternate in-frame exons in the central coding region, compared to variant 1. The resulting isoform (2) lacks a segment of the sarcoglycan

domain including the transmembrane domain, compared to isoform 1.