

## **Product datasheet for SC202925**

## 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

## alpha Sarcoglycan (SGCA) (NM\_000023) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

**Product Name:** alpha Sarcoglycan (SGCA) (NM\_000023) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: SGCA

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

**ACCN:** NM\_000023

**Insert Size:** 259 bp

Insert Sequence: >SC202925 3'UTR clone of NM\_000023

The sequence shown below is from the reference sequence of NM\_000023. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

CAGGTGCCCCTCATTCTGGACCAGCACTGACAGCCTAGCCAGTGGTTCCAGGTCCAGCCCTGACTTCAT CCTCCCTTCTCTGTCCACACCACGAGTGGCACATCCCACCTGCTGATTCCAGCTCCTGGCCCTCCTGGAACCCAGGCTCTAAACAAGCAGGGAGAGGGGGTGGGGTGGGGTGAGAGTGTGTGGAGTAAGGACATTCAG

AATAAATATCTGCTGCTCGCTCACCAATTGCTGCTGGCAGCCTCTCCCGTC

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul

**OTI Disclaimer:** Our molecular clone sequence data has been matched to the sequence identifier above as a

point of reference. Note that the complete sequence of this clone is largely the same as the

reference sequence but may contain minor differences, e.g., single nucleotide

polymorphisms (SNPs).

**Components:** The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

**RefSeq:** <u>NM 000023.4</u>





## alpha Sarcoglycan (SGCA) (NM\_000023) Human 3' UTR Clone - SC202925

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]

**Locus ID:** 6442 **MW:** 9.3