abcam

Product datasheet

Recombinant Human delta Sarcoglycan protein (denatured) ab140566

1 Image

Description

Product name Recombinant Human delta Sarcoglycan protein (denatured)

Purity > 85 % SDS-PAGE.

Expression system Escherichia coli

Accession Q92629

Protein length Protein fragment

Animal free No

Nature Recombinant

Species Human

Sequence MGSSHHHHHH SSGLVPRGSH RSHMKVMNFT

IDGMGNLRIT EKGLKLEGDS EFLQPLYAKE IQSRPGNALY FKSARNVTVN ILNDQTKVLT QLITGPKAVE AYGKKFEVKT

VSGKLLFSAD NNEVVVGAER LRVLGAEGTV FPKSIETPNV RADPFKELRL ESPTRSLVME

APKGVEINAE AGNMEATCRT ELRLESKDGE IKLDAAKIRL

PRLPHGSYTP TGTRQKVFEI CVCANGRLFL

SQAGAGSTCQ INTSVCL

Predicted molecular weight 28 kDa including tags

Amino acids 57 to 289

Tags His tag N-Terminus

Description Recombinant Human delta Sarcoglycan protein

Specifications

Our **Abpromise guarantee** covers the use of **ab140566** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications SDS-PAGE

Form Liquid

Preparation and Storage

1

Stability and Storage

Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.

pH: 8.00

Constituents: 2.4% Urea, 0.32% Tris HCI, 10% Glycerol (glycerin, glycerine)

General Info

Function Component of the sarcoglycan complex, a subcomplex of the dystrophin-glycoprotein complex

which forms a link between the F-actin cytoskeleton and the extracellular matrix.

Tissue specificity Most strongly expressed in skeletal and cardiac muscle. Also detected in smooth muscle. Weak

expression in brain and lung.

Involvement in diseaseDefects in SGCD are the cause of limb-girdle muscular dystrophy type 2F (LGMD2F)

[MIM:601287]. LGMD2F is an autosomal recessive disorder.

Defects in SGCD are the cause of cardiomyopathy dilated type 1L (CMD1L) [MIM:606685]. Dilated cardiomyopathy is a disorder characterized by ventricular dilation and impaired systolic function, resulting in congestive heart failure and arrhythmia. Patients are at risk of premature

death.

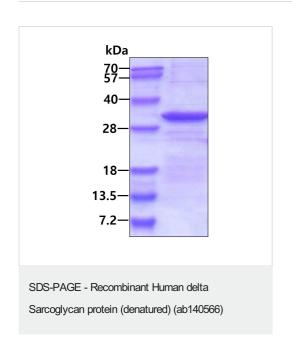
Sequence similaritiesBelongs to the sarcoglycan beta/delta/gamma/zeta family.

Post-translational Glycosylated.

modifications Disulfide bonds are present.

Cellular localization Cell membrane > sarcolemma. Cytoplasm > cytoskeleton.

Images



SDS-PAGE of ab140566 (3 μ g) under reducing condition and visualized by coomassie blue stain.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

Our Abpromise to you: Quality guaranteed and expert technical support

- · Replacement or refund for products not performing as stated on the datasheet
- Valid for 12 months from date of delivery
- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- · We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit https://www.abcam.com/abpromise or contact our technical team.

Terms and conditions

• Guarantee only valid for products bought direct from Abcam or one of our authorized distributors