abcam

Product datasheet

Recombinant Human Tropomyosin 2 protein ab103503

1 Image

Description

Product name Recombinant Human Tropomyosin 2 protein

Purity > 90 % SDS-PAGE.

ab103503 was purified by using anion-exchange chromatography (DEAE sepharose resin) and

gel-filtration chromatography (Sephacryl S-200) with 20mM Tris pH 7.5, 2mM EDTA.

Expression system Escherichia coli

Accession P07951-2

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence MGSSHHHHHHSSGLVPRGSHMDAIKKKMQMLKLDKEN

AIDRAEQAEADKK

QAEDRCKQLEEEQQALQKKLKGTEDEVEKYSESVKEAQ

EKLEQAEKKATD

AEADVASLNRRIQLVEEELDRAQERLATALQKLEEAEKAA

DESERGMKVI

ENRAMKDEEKMELQEMQLKEAKHIAEDSDRKYEEVARKL

VILEGELERSE

ERAEVAESRARQLEEELRTMDQALKSLMASEEEYSTKED

KYEEEIKLLEE

KLKEAETRAEFAERSVAKLEKTIDDLEETLASAKEENVEIH

QTLDQTLLE LNNL

Predicted molecular weight 35 kDa including tags

Amino acids 1 to 284

Tags His tag N-Terminus

Specifications

Our <u>Abpromise guarantee</u> covers the use of ab103503 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

Mass Spectrometry

1

Mass spectrometry

MALDI-TOF

Form

Liquid

Preparation and Storage

Stability and Storage

Shipped at $4\,^\circ\text{C}$. Upon delivery aliquot and store at -20 $^\circ\text{C}$ or -80 $^\circ\text{C}$. Avoid repeated freeze / thaw

cycles.

pH: 8.00

Constituents: 0.0154% DTT, 0.316% Tris HCl, 30% Glycerol (glycerin, glycerine), 0.58% Sodium

chloride

General Info

Function

Binds to actin filaments in muscle and non-muscle cells. Plays a central role, in association with the troponin complex, in the calcium dependent regulation of vertebrate striated muscle contraction. Smooth muscle contraction is regulated by interaction with caldesmon. In non-muscle cells is implicated in stabilizing cytoskeleton actin filaments. The non-muscle isoform may have a role in agonist-mediated receptor internalization.

Tissue specificity

Present in primary breast cancer tissue, absent from normal breast tissue.

Involvement in disease

Nemaline myopathy 4 (NEM4) [MIM:609285]: A form of nemaline myopathy. Nemaline myopathies are muscular disorders characterized by muscle weakness of varying severity and onset, and abnormal thread-or rod-like structures in muscle fibers on histologic examination. Nemaline myopathy type 4 presents from infancy to childhood with hypotonia and moderate-to-severe proximal weakness with minimal or no progression. Major motor milestones are delayed but independent ambulation is usually achieved, although a wheelchair may be needed in later life. Note=The disease is caused by mutations affecting the gene represented in this entry. Arthrogryposis, distal, 1A (DA1A) [MIM:108120]: A form of distal arthrogryposis, a disease characterized by congenital joint contractures that mainly involve two or more distal parts of the

limbs, in the absence of a primary neurological or muscle disease. Distal arthrogryposis type 1 is characterized largely by camptodactyly and clubfoot. Hypoplasia and/or absence of some interphalangeal creases is common. The shoulders and hips are less frequently affected. Note=The disease is caused by mutations affecting the gene represented in this entry.

Sequence similarities

Belongs to the tropomyosin family.

Domain

The molecule is in a coiled coil structure that is formed by 2 polypeptide chains. The sequence

exhibits a prominent seven-residues periodicity.

Post-translational modifications

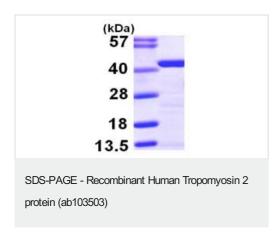
Phosphorylated on Ser-61 by PIK3CG. Phosphorylation on Ser-61 is required for ADRB2

internalization.

Cellular localization

Cytoplasm > cytoskeleton.

Images



15% SDS-PAGE analysis of 3µg ab103503.

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