abcam

Product datasheet

Recombinant Human Cardiac Troponin T protein ab86685

2 Images

Description

Product name Recombinant Human Cardiac Troponin T protein

Purity > 90 % SDS-PAGE.

ab86685 is purified using conventional chromatography techniques.

Expression system Escherichia coli

Accession P45379-11

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence MGSSHHHHHH SSGLVPRGSH MSDIEEVVEE

YEEEEQEEAA VEEQEEAAEE DAEAEAETEE TRAEEDEEEE EAKEAEDGPM EESKPKPRSF MPNLVPPKIP DGERVDFDDI HRKRMEKDLN

ELQALIEAHF ENRKKEEEEL VSLKDRIERR RAERAEQQRI

RNEREKERQN RLAEERARRE EEENRRKAED EARKKKALSN MMHFGGYIQK TERKSGKRQT

EREKKKKILA ERRKVLAIDH LNEDQLREKA KELWQSIYNL

EAEKFDLQEK FKQQKYEINV LRNRINDNQK

VSKTRGKAKV TGRWK

Predicted molecular weight 36 kDa

Amino acids 1 to 285

Tags His tag N-Terminus

Additional sequence information | Isoform 11 of Cardiac Troponin T.

Specifications

Our Abpromise guarantee covers the use of ab86685 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

Sandwich ELISA

1

Form Liquid

Preparation and Storage

Stability and Storage

Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw

cycles.

pH: 8.00

Preservative: 0.0006% Imidazole

Constituents: 0.24% Tris HCI, 50% Glycerol, 1.16% Sodium chloride, 0.0017% PMSF, 0.02%

DTT

General Info

Function

Troponin T is the tropomyosin-binding subunit of troponin, the thin filament regulatory complex which confers calcium-sensitivity to striated muscle actomyosin ATPase activity.

Tissue specificity

Heart. The fetal heart shows a greater expression in the atrium than in the ventricle, while the adult heart shows a greater expression in the ventricle than in the atrium. Isoform 6 predominates in normal adult heart. Isoforms 1, 7 and 8 are expressed in fetal heart. Isoform 7 is also expressed in failing adult heart.

Involvement in disease

Defects in TNNT2 are the cause of cardiomyopathy familial hypertrophic type 2 (CMH2) [MIM:115195]. Familial hypertrophic cardiomyopathy is a hereditary heart disorder characterized by ventricular hypertrophy, which is usually asymmetric and often involves the interventricular septum. The symptoms include dyspnea, syncope, collapse, palpitations, and chest pain. They can be readily provoked by exercise. The disorder has inter- and intrafamilial variability ranging from benign to malignant forms with high risk of cardiac failure and sudden cardiac death.

Defects in TNNT2 are the cause of cardiomyopathy dilated type 1D (CMD1D) [MIM:601494].

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.

Defects in TNNT2 are the cause of cardiomyopathy familial restrictive type 3 (RCM3)

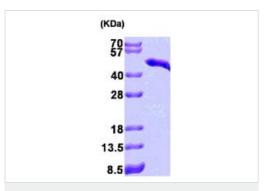
[MIM:612422]. Restrictive cardiomyopathy is a heart disorder characterized by impaired filling of the ventricles with reduced diastolic volume, in the presence of normal or near normal wall thickness and systolic function.

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Sequence similarities

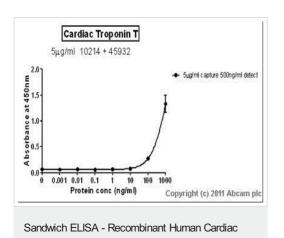
Belongs to the troponin T family.

Images



SDS-PAGE - Recombinant Human Cardiac Troponin T protein (ab86685)





Troponin T protein (ab86685)

Standard Curve for Cardiac Troponin T (Analyte: <u>Cardiac</u>

<u>Troponin T protein (His tag) (ab86685)</u>); dilution range 1pg/ml to

1ug/ml using Capture Antibody <u>Mouse monoclonal [1F11] to</u>

<u>Cardiac Troponin T (ab10214)</u> at 5ug/ml and Detector Antibody

<u>Rabbit polyclonal to cardiac Troponin T (ab45932)</u> at 0.5ug/ml.

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

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