abcam

Product datasheet

Recombinant Human ACADS/SCAD protein ab95383

1 Image

Description

Product name Recombinant Human ACADS/SCAD protein

Purity > 95 % SDS-PAGE.

ab95383 is purified using conventional chromatography techniques.

Expression system Escherichia coli

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence MGSSHHHHHH SSGLVPRGSH MLHTIYQSVE

LPETHQMLLQ TCRDFAEKEL FPIAAQVDKE HLFPAAQVKK MGGLGLLAMD VPEELGGAGL

DYLAYAIAME EISRGCASTG VIMSVNNSLY LGPILKFGSK

EQKQAWVTPF TSGDKIGCFA LSEPGNGSDA GAASTTARAE GDSWVLNGTK AWITNAWEAS AAVVFASTDR ALQNKSISAF LVPMPTPGLT

LGKKEDKLGI RGSSTANLIF EDCRIPKDSI LGEPGMGEKI

AMQTLDMGRI GIASQALGIA QTALDCAVNY AENRMAFGAP LTKLQVIQFK LADMALALES ARLLTWRAAM LKDNKKPFIK EAAMAKLAAS

EAATAISHQA IQILGGMGYV TEMPAERHYR DARITEIYEG

TSEIQRLVIA GHLLRSYRS

Specifications

Our Abpromise guarantee covers the use of ab95383 in the following tested applications.

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

Applications Mass Spectrometry

SDS-PAGE

Form Liquid

Additional notes Previously labelled as ACADS.

Preparation and Storage

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

cycles.

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Constituents: 0.0154% DTT, 0.316% Tris HCl, 20% Glycerol (glycerin, glycerine), 0.58% Sodium

chloride

General Info

Pathway Lipid metabolism; mitochondrial fatty acid beta-oxidation.

Involvement in disease Defects in ACADS are the cause of acyl-CoA dehydrogenase short-chain deficiency (ACADSD)

[MIM:201470]. It is an autosomal recessive disorder resulting in acute acidosis and muscle

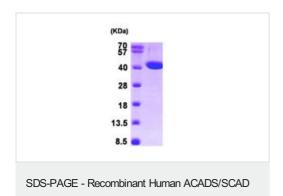
weakness in infants, and a form of lipid-storage myopathy in adults.

Sequence similaritiesBelongs to the acyl-CoA dehydrogenase family.

Cellular localization Mitochondrion matrix.

Images

protein (ab95383)



15% SDS-PAGE analysis of 3µg ab95383.

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

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