

Recombinant Human ACADM/MCAD protein ab99329

1 Image

Description

| | |
|----------------------------|--|
| Product name | Recombinant Human ACADM/MCAD protein |
| Purity | > 90 % SDS-PAGE. ab99329 is purified using conventional chromatography techniques. |
| Expression system | Escherichia coli |
| Accession | P11310 |
| Protein length | Full length protein |
| Animal free | No |
| Nature | Recombinant |
| Species | Human |
| Sequence | MGSSHHHHHHSSGLVPRGSHM KANRQREPGLGFSFEF TEQQKEFQATARK FAREEIIPVAAEYDKTGEYPVPLIRRAWELGLMNTHIPENC GGLGLGTFD ACLISEELAYGCTGVQTAIEGNSLGQMPIIIAGNDQQKKKYL GRMTEEPL MCAYCVTEPGAGSDVAGIKTKAEKKGDEYIINGQKMWITN GGKANWYFLL ARSDPDPKAPANKAFTGFIVEADTPGIQIGRKELNMGQRC SDTRGVFED VKVPKENVLIGDGAGFKVAMGAFDKTRPVVAAGAVGLAQ RALDEATKYAL ERKTFGKLLVEHQAI SFMLAEMAMKVELARMSYQRAAWE VDSGRRNTYYA SIAKAFAGDIANQLATDAVQILGGNGFNTEYPVEKLMRDAK YQIYEGTS QIQLIVAREHIDKYKN |
| Predicted molecular weight | 46 kDa including tags |
| Amino acids | 26 to 421 |
| Tags | His tag N-Terminus |

Specifications

Our **Abpromise guarantee** covers the use of **ab99329** in the following tested applications.

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

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|--------------------------|-------------------------------|
| Applications | SDS-PAGE |
| | Mass Spectrometry |
| Mass spectrometry | MALDI-TOF |
| Form | Liquid |
| Additional notes | Previously labelled as ACADM. |

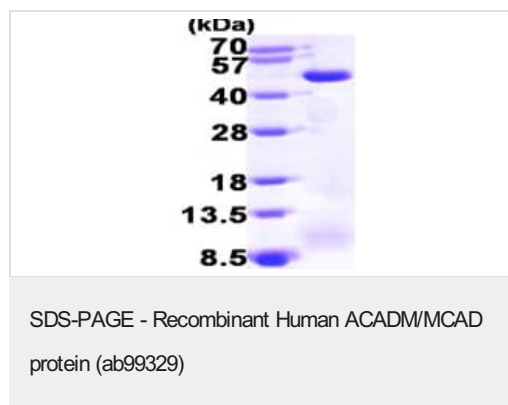
Preparation and Storage

| | |
|------------------------------|---|
| 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 |
| | Constituents: 0.316% Tris HCl, 20% Glycerol (glycerin, glycerine), 0.58% Sodium chloride |

General Info

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|-------------------------------|--|
| Function | This enzyme is specific for acyl chain lengths of 4 to 16. |
| Pathway | Lipid metabolism; mitochondrial fatty acid beta-oxidation. |
| Involvement in disease | Defects in ACADM are the cause of acyl-CoA dehydrogenase medium-chain deficiency (ACADM) [MIM:201450]. It is an autosomal recessive disease which causes fasting hypoglycemia, hepatic dysfunction, and encephalopathy, often resulting in death in infancy. |
| Sequence similarities | Belongs to the acyl-CoA dehydrogenase family. |
| Cellular localization | Mitochondrion matrix. |

Images



15% SDS-PAGE analysis of 3µg ab99329.

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

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