# abcam

### Product datasheet

## Recombinant Human ACADL/LCAD protein ab113579

#### 1 图像

描述

产品名称 重组人ACADL/LCAD蛋白

纯**度** > 85 % SDS-PAGE.

> 85 % by SDS - PAGE. ab113579 is purified using conventional chromatography techniques.

表达系统 Escherichia coli

Accession P28330

**蛋白长度** Full length protein

无动物成分 No

性质 Recombinant

种属 Human

序列 MGSSHHHHHHSSGLVPRGSHMGGEERLETPSAKKLTDIGIRR

IFSPEHDI

FRKSVRKFFQEEVIPHHSEWEKAGEVSREVWEKAGKQGLLGV

NIAEHLGG

IGGDLYSAAIVWEEQAYSNCSGPGFSIHSGIVMSYITNHGSE

**EQIKHFIP** 

 ${\tt QMTAGKCIGAIAMTEPGAGSDLQGIKTNAKKDGSDWILNGSK}$ 

VFISNGSL

SDVVIVVAVTNHEAPSPAHGISLFLVENGMKGFIKGRKLHKM

GLKAQDTA

ELFFEDIRLPASALLGEENKGFYYIMKELPQERLLIADVAIS

ASEFMFEE

TRNYVKQRKAFGKTVAHLQTVQHKLAELKTHICVTRAFVDNC

LQLHEAKR

LDSATACMAKYWASELQNSVAYDCVQLHGGWGYMWEYPIAKA

YVDARVQP IYGGTNEIMKELIAREIVFDK

预**测分子量** 47 kDa including tags

**氨基酸** 31 to 430

标签 His tag N-Terminus

技术指标

Our Abpromise quarantee covers the use of ab113579 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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应用 SDS-PAGE

Mass Spectrometry

质**谱法** MALDI-TOF

形式 Liquid

**补充说明** Previously labelled as ACADL.

#### 制备和贮存

稳定性和存储 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: 0.02% DTT, 0.32% Tris HCI, 10% Glycerol (glycerin, glycerine), 0.88% Sodium

chloride

#### 常规信息

通路 Lipid metabolism; mitochondrial fatty acid beta-oxidation.

疾病相关 Defects in ACADL are a cause of acyl-CoA dehydrogenase very long-chain deficiency

(ACADVLD) [MIM:201475]. An inborn error of mitochondrial fatty acid beta-oxidation which leads to impaired long-chain fatty acid beta-oxidation. It is clinically heterogeneous, with three major phenotypes: a severe childhood form characterized by early onset, high mortality and high incidence of cardiomyopathy; a milder childhood form with later onset, characterized by hypoketotic hypoglycemia, low mortality and rare cardiomyopathy; an adult form, with isolated skeletal muscle involvement, rhabdomyolysis and myoglobinuria, usually triggered by exercise or

fasting.

**序列相似性** Belongs to the acyl-CoA dehydrogenase family.

细胞定位 Mitochondrion matrix.

#### 图片



3ug by SDS-PAGE under reducing conditions and visualized by coomassie blue stain.

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