abcam

Product datasheet

Recombinant Human ACADVL/VLCAD protein ab98234

1图像

产 品名称 纯 度	重组人ACADVL/VLCAD蛋白		
纯 度	> 90 % SDS-PAGE		
	> 90 % SDS-PAGE.		
	ab98234 was purified using conventional chromatography techniques.		
表达系统	Escherichia coli		
Accession	<u>P49748</u>		
蛋白长度	Full length protein		
无 动 物成分	No		
性质	Recombinant		
种属	Human		
序列	MGSSHHHHHHSSGLVPRGSHMAGGAAQLALDKSDSLTRKKPAKAESKSFAVGMFKGQLTTDQVFPYPSVLNEEQTQFLKELVFFEEVNDPAKNDALEMVEETTWQGLKELGAFGLQVPSELGGVGLCRLVEIVGMHDLGVGTTLGAHQSIGFKGILLFGTKAQKEKYLPKLAAAFCLTEPSSGSDAASIRTSAVPSPCGKYYTLNGSKLWISNGGLAFAKTPVTDPATGAVKEKITAFVVERGFGGITHGPPEKKMGIKASNFDGVRVPSENVLGEVGSGFKVAMHILNNGRFGMAAALAGTMRGIIHATNRTQFGEKIHHFGLIQEKLARMVMLQYVTESMAYMVSANMDQQIEAAISKIFGSEAAWKVTDECIQIMGGMGFMKEPGVERVLRDLREGTNDILRLFVALQGCMDKGKELSGLGSALKNPFGNAGLLLGEAGRAGLGSGLSLSGLVHPELSRSGELAVRALEQFATVVEAKLIKHKKQFLLQRLADGAIDLYAMVVVLSRASRSLSEGHPTAQHEKMLCDTWARIREGMAALQSDPWQQELYRNFKSISKALVERGGVVTSNPLGF	EPVSR NTQYA SGETV DIFTV TAEVF AKAVD GATDF LFRIF KQLRR GIVNE	

预测分子量	69 kDa including tags
氨基酸	41 to 655
标签	His tag N-Terminus

技术指标

Our Abpromise guarantee covers the use of ab98234 in the following tested applications.

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

应 用	SDS-PAGE
形式	Liquid
补 充 说 明	Previously labelled as ACADVL.
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: 0.0154% DTT, 0.316% Tris HCI, 0.0292% EDTA, 10% Glycerol (glycerin, glycerine), 0.58% Sodium chloride
常规信息	
功能	Active toward esters of long-chain and very long chain fatty acids such as palmitoyl-CoA, mysritoyl-CoA and stearoyl-CoA. Can accomodate substrate acyl chain lengths as long as 24 carbons, but shows little activity for substrates of less than 12 carbons.
通路	Lipid metabolism; mitochondrial fatty acid beta-oxidation.
疾病相关	Defects in ACADVL are the cause of acyl-CoA dehydrogenase very long chain deficiency (ACADVLD) [MIM:201475]. ACADVLD is an autosomal recessive disease which leads to impaired long-chain fatty acid beta-oxidation. It is clinically heterogeneous, with three major
	phenotypes: a severe childhood form, with early onset, high mortality, and high incidence of cardiomyopathy; a milder childhood form, with later onset, usually with hypoketotic hypoglycemia as the main presenting feature, low mortality, and rare cardiomyopathy; and an adult form, with isolated skeletal muscle involvement, rhabdomyolysis, and myoglobinuria, usually triggered by exercise or fasting.
序列相似性	cardiomyopathy; a milder childhood form, with later onset, usually with hypoketotic hypoglycemia as the main presenting feature, low mortality, and rare cardiomyopathy; and an adult form, with isolated skeletal muscle involvement, rhabdomyolysis, and myoglobinuria, usually triggered by



15% SDS-PAGE analysis of 3µg ab98234

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