# abcam

## Product datasheet

## Recombinant Human GCDH/GCD protein ab98118

## 1 图像

描述

产品名称 重组人GCDH/GCD蛋白

纯**度** > 90 % SDS-PAGE.

ab98118 is purified by using conventional chromatography techniques.

表达系统 Escherichia coli

Accession Q92947

**蛋白长度** Full length protein

无动物成分 No

性质 Recombinant

种属 Human

序列 MGSSHHHHHHSSGLVPRGSHMRPEFDWQDPLVLEEQLTTD

EILIRDTFRT

YCQERLMPRILLANRNEVFHREIISEMGELGVLGPTIKGYGC

AGVSSVAY

GLLARELERVDSGYRSAMSVQSSLVMHPIYAYGSEEQRQKYL

PQLAKGEL

LGCFGLTEPNSGSDPSSMETRAHYNSSNKSYTLNGTKTWITN

**SPMADLFV** 

VWARCEDGCIRGFLLEKGMRGLSAPRIQGKFSLRASATGMII

**MDGVEVPE** 

ENVLPGASSLGGPFGCLNNARYGIAWGVLGASEFCLHTARQY

ALDRMQFG

VPLARNQLIQKKLADMLTEITLGLHACLQLGRLKDQDKAAPE

**MVSLLKRN** 

NCGKALDIARQARDMLGGNGISDEYHVIRHAMNLEAVNTYEG

THDIHALI LGRAITGIQAFTASK

预**测分子量** 46 kDa including tags

**氨基酸** 45 to 438

标签 His tag N-Terminus

技术指标

Our Abpromise guarantee covers the use of ab98118 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 This product was previously labelled as GCDH

#### 制备和贮存

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

cycles.

pH: 8.00

Constituents: 0.077% DTT, 0.316% Tris HCI, 20% Glycerol (glycerin, glycerine), 1.16% Sodium

chloride

#### 常规信息

功能 Catalyzes the oxidative decarboxylation of glutaryl-CoA to crotonyl-CoA and CO(2) in the

degradative pathway of L-lysine, L-hydroxylysine, and L-tryptophan metabolism. It uses electron

transfer flavoprotein as its electron acceptor. Isoform Short is inactive.

组织特异性 Isoform 1 and isoform 2 are expressed in fibroblasts and liver.

通路 Amino-acid metabolism; lysine degradation.

Amino-acid metabolism; tryptophan metabolism.

疾病相关 Defects in GCDH are the cause of glutaric aciduria type 1 (GA1) [MIM:231670]. GA1 is an

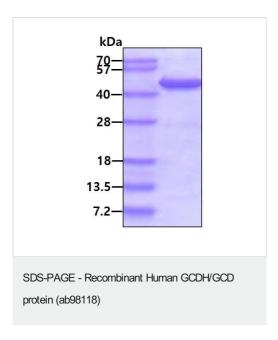
autosomal recessive metabolic disorder characterized by progressive dystonia and athetosis due

to gliosis and neuronal loss in the basal ganglia.

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

细**胞定位** Mitochondrion matrix.

## 图片



SDS-PAGE analysis of ab98118 (3µg) under reducing condition and visualized by coomassie blue stain.

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

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