abcam

Product datasheet

Recombinant Human GCDH/GCD protein ab98118

画像数1

製品の詳細

製品名 Recombinant Human GCDH/GCD protein

精製度 > 90 % SDS-PAGE.

ab98118 is purified by using conventional chromatography techniques.

発現系 Escherichia coli

アクセッション番号 <u>Q92947</u>

タンパク質長 Full length protein

Animal free No

由来 Recombinant

生物種 Human

配列 MGSSHHHHHHSSGLVPRGSHMRPEFDWQDPLVLEEQLTTD

EILIRDTFRT

YCQERLMPRILLANRNEVFHREIISEMGELGVLGPTIKGYGC

AGVSSVAY

GLLARELERVDSGYRSAMSVQSSLVMHPIYAYGSEEQRQKYL

PQLAKGEL

 ${\tt LGCFGLTEPNSGSDPSSMETRAHYNSSNKSYTLNGTKTWITN}$

SPMADLFV

VWARCEDGCIRGFLLEKGMRGLSAPRIQGKFSLRASATGMII

MDGVEVPE

ENVLPGASSLGGPFGCLNNARYGIAWGVLGASEFCLHTARQY

ALDRMQFG

VPLARNQLIQKKLADMLTEITLGLHACLQLGRLKDQDKAAPE

MVSLLKRN

NCGKALDIARQARDMLGGNGISDEYHVIRHAMNLEAVNTYEG

THDIHALI LGRAITGIQAFTASK

予測される分子量 46 kDa including tags

領域 45 to 438

サブ His tag N-Terminus

特性

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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アプリケーション SDS-PAGE

Mass Spectrometry

質量分析 MALDI-TOF

製品の状態 Liquid

備考 This product was previously labelled as GCDH

前処理および保存

保存方法および安定性 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

関連情報

機能 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.

組織特異性 Isoform 1 and isoform 2 are expressed in fibroblasts and liver.

אליכיל Amino-acid metabolism; lysine degradation.

Amino-acid metabolism; tryptophan metabolism.

関連疾患 Defects in GCDH are the cause of glutaric aciduria type 1 (GA1) [MIM:231670]. GA1 is an

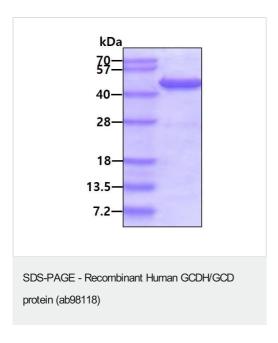
autosomal recessive metabolic disorder characterized by progressive dystonia and athetosis due

to gliosis and neuronal loss in the basal ganglia.

配列類似性 Belongs to the acyl-CoA dehydrogenase family.

細胞内局在 Mitochondrion matrix.

画像



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

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