abcam

Product datasheet

Anti-GCDH/GCD antibody [EPR14864(B)] ab182152

Recombinant RabMAb

4 Images

Overview

Product name Anti-GCDH/GCD antibody [EPR14864(B)]

Description Rabbit monoclonal [EPR14864(B)] to GCDH/GCD

Host species Rabbit

Tested applications Suitable for: WB, Flow Cyt (Intra)

Reacts with: Human Species reactivity

Predicted to work with: Mouse, Rat

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control Human fetal liver and HepG2 cell lysates; PC3 cells.

General notes This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply - Animal-free production

For more information see here.

Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb patents**.

Properties

Form Liquid

Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long Storage instructions

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol (glycerin, glycerine), 59% PBS, 0.05% BSA

Purity Tissue culture supernatant

Clonality Monoclonal EPR14864(B) Clone number

Isotype lgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab182152 in the following tested applications.

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

Application	Abreviews	Notes
WB		1/1000 - 1/10000. Detects a band of approximately 48 kDa (predicted molecular weight: 48 kDa).
Flow Cyt (Intra)		1/80. ab172730 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.

Target

Function 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.

Tissue specificity Isoform 1 and isoform 2 are expressed in fibroblasts and liver.

Pathway Amino-acid metabolism; lysine degradation.

Amino-acid metabolism; tryptophan metabolism.

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

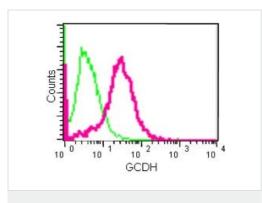
autosomal recessive metabolic disorder characterized by progressive dystonia and athetosis due

to gliosis and neuronal loss in the basal ganglia.

Sequence similarities Belongs to the acyl-CoA dehydrogenase family.

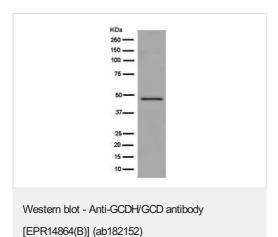
Cellular localization Mitochondrion matrix.

Images



Flow Cytometry (Intracellular) - Anti-GCDH/GCD antibody [EPR14864(B)] (ab182152)

Intracellular flow cytometric analysis of PC3 cells (paraformaldehyde-fixed, 2%) labeling GCDH/GCDwith ab182152 at 1/80 dilution (red) or a rabbit lgG (negative) (green), followed by Goat anti rabbit lgG (FITC) secondary at 1/150 dilution.

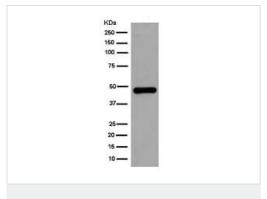


Anti-GCDH/GCD antibody [EPR14864(B)] (ab182152) at 1/5000 dilution + HepG2 cell lysate at 20 µg

Secondary

Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugate at 1/1000 dilution

Predicted band size: 48 kDa



Anti-GCDH/GCD antibody [EPR14864(B)] (ab182152) at 1/20000 dilution + Human fetal liver lysate at 20 μg

Secondary

Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugate at 1/1000 dilution

Predicted band size: 48 kDa

Western blot - Anti-GCDH/GCD antibody [EPR14864(B)] (ab182152)



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

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