

GCDH Rabbit pAb

货号: **AYP13202**

产品信息

反应	Human,Mouse,Rat
宿主	Rabbit
克隆性	Polyclonal
预测反应	WB: Homo sapiens
应用	WB
推荐浓度	WB: 1:200 - 1:2000
理论分子量	47kDa/48kDa
实测分子量	48kDa
形式	Liquid
保存条件	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3.
偶联物	Unconjugated
阳性对照	HL-60,LO2,MCF7,BT-474,Mouse liver,Rat liver,Rat kidney
细胞定位	Mitochondrion matrix
纯化	Affinity purification

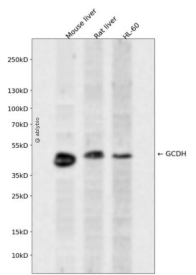
抗原信息

抗原信息	Recombinant fusion protein containing a sequence corresponding to amino acids 149-438 of human GCDH (NP_000150.1).
序列	MHPIYAYGSEEQRQKYLPLAKGELLGCFGLTEPNSGSDPSSMETRAHYNSSNKSYYTLNGTKTWITNSPMADLFVWWAR CEDGCIRGFLLLEKGMRGLSAPRIQGGKFSLRASATGMIIMDGVPEENVLPGASSLGGPFGCLNNARYGIAWGVLGASEF CLHTARQYALDRMQFGVPLARNQLIQKKLADMLTEITLGLHAQLGRLKDQDKAAPEMVSLKRNCGKALDIARQARD MLGGNGISDEYHVIRHAMNLEAVNTYEGTHDIHALILGRAITGIQAF TASK

靶点信息

研究背景	The protein encoded by this gene belongs to the acyl-CoA dehydrogenase family. It 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. The enzyme exists in the mitochondrial matrix as a homotetramer of 45-kD subunits. Mutations in this gene result in the metabolic disorder glutaric aciduria type 1, which is also known as glutaric acidemia type I. Alternative splicing of this gene results in multiple transcript variants. A related pseudogene has been identified on chromosome 12.
基因ID	2639
基因名	GCDH
Swiss	Q92947
别名	GCDH;ACAD5;GCD

产品验证



Western blot analysis of GCDH expressed in Mouse liver, Rat liver, HL-60 using GCDH Rabbit pAb at 1:1000. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) at 1:5000. Lysates/proteins: 30ug per lane. Blocking buffer: 5% non-fat dry milk in TBST. Detection: ECL Enhanced Kit. Exposure time: 120s.

实验步骤

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