abcam

Product datasheet

Anti-Abhd5/CGI-58 antibody ab111984

1 References 3 图像

概述

产**品名称** Anti-Abhd5/CGI-58抗体

描述 山羊多克隆抗体to Abhd5/CGI-58

宿主 Goat

适用于: WB, IHC-P

种属反应性 与反应: Mouse, Human

预测可用于: Rat, Dog, Pig 🔷

免疫原 Synthetic peptide corresponding to Human Abhd5/CGI-58 aa 150-250 (internal sequence).

Database link: **Q8WTS1**

Run BLAST with
Run BLAST with

常规说明

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

性能

形式 Liquid

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

term.

存储溶液 pH: 7.30

Preservative: 0.02% Sodium azide

Constituents: 0.5% BSA, 99% Tris buffered saline

纯**度** Immunogen affinity purified

克隆 多克隆

同种型 IgG

应用

1

The Abpromise guarantee

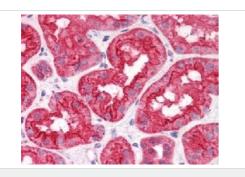
Abpromise™承诺保证使用ab111984于以下的经测试应用

"应用说明"部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB		Use a concentration of 0.2 - 0.6 µg/ml. Predicted molecular weight: 39 kDa.
IHC-P		Use a concentration of 3.75 μg/ml.

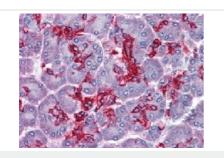
靶标	
功能	Lysophosphatidic acid acyltransferase which functions in phosphatidic acid biosynthesis. May regulate the cellular storage of triacylglycerol through activation of the phospholipase PNPLA2. Involved in keratinocyte differentiation.
组织 特异性	Widely expressed in various tissues, including lymphocytes, liver, skeletal muscle and brain. Expressed by upper epidermal layers and dermal fibroblasts in skin, hepatocytes and neurons (at protein level).
疾病相关	Defects in ABHD5 are the cause of Chanarin-Dorfman syndrome (CDS) [MIM:275630]; also called triglyceride storage disease with impaired long-chain fatty acid oxidation or neutral lipid storage disease with ichthyosis. CDS is an autosomal recessive inborn error of lipid metabolism with multisystemic accumulation of triglycerides although plasma concentrations are normal. Clinical characteristics are congenital generalized ichthyosis, vacuolated leukocytes, hepatomegaly, myopathy, cataracts, neurosensory hearing loss and developmental delay. The disorder presents at birth with generalized, fine, white scaling of the skin and a variable degree of erythema resembling non-bullous congenital ichthyosiform erythroderma.
序列相似性	Belongs to the peptidase S33 family. ABHD4/ABHD5 subfamily.
发 展 阶段	Detected in fetal epidermis from 49 to 135 days estimated gestational age (at protein level).
结 构域	The HXXXXD motif is essential for acyltransferase activity and may constitute the binding site for the phosphate moiety of the glycerol-3-phosphate.
细胞定位	Cytoplasm. Lipid droplet. Colocalized with PLIN and ADRP on the surface of lipid droplets. The localization is dependent upon the metabolic status of the adipocytes and the activity of PKA.

图片



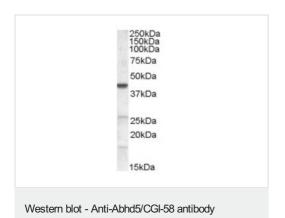
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Abhd5/CGI-58 antibody (ab111984)

ab111984, at $3.75 \mu g/ml$, staining Abhd5/CGI-58 in formalin-fixed, paraffin-embedded Human kidney tissue by Immunohistochemistry.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Abhd5/CGI-58 antibody (ab111984)

ab111984, at 3.75 μ g/ml, staining Abhd5/CGI-58 in formalin-fixed, paraffin-embedded Human pancreas tissue by Immunohistochemistry.



(ab111984)

Anti-Abhd5/CGI-58 antibody (ab111984) at 0.2 μ g/ml + NIH3T3 lysate (in RIPA buffer) at 35 μ g

Developed using the ECL technique.

Predicted band size: 39 kDa

Primary incubation was 1 hour.

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

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