abcam

Product datasheet

Anti-CYP7B1 antibody [EPR8395] ab138497

重组 RabMAb

★★★★★ 1 Abreviews 10 References 3 图像

概述

产品名称 Anti-CYP7B1抗体[EPR8395]

描述 兔单克隆抗体[EPR8395] to CYP7B1

宿主 Rabbit

经测试应用 适用于: WB

不适用于: Flow Cyt,ICC/IF,IHC-P or IP

种属反应性 与反应: Human

免疫原 Synthetic peptide within Human CYP7B1 aa 400-500. The exact sequence is proprietary.

阳性对照 PC3, Human fetal liver, and Human fetal kidney lysates

常规说明 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**.

Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with

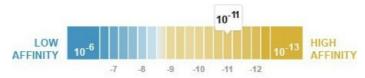
these species. Please contact us for more information.

性能

形式 Liquid

存放说明 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

解离常数(Kn) $K_D = 4.10 \times 10^{-11} M$



Learn more about K_D

存储溶液 pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 9% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA, 50% Tissue culture

supernatant

纯**度** Tissue culture supernatant

克隆 单克隆

克隆编号 EPR8395

同种型 IgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab138497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	****(1)	1/1000 - 1/10000. Detects a band of approximately 55 kDa (predicted molecular weight: 58 kDa).

应用说明 Is unsuitable for Flow Cyt,ICC/IF,IHC-P or IP.

靶标

组织特异性 Brain, testis, ovary, prostate, liver, colon, kidney, and small intestine.

通路 Lipid metabolism; bile acid biosynthesis.

疾病相关 Defects in CYP7B1 are the cause of spastic paraplegia autosomal recessive type 5A (SPG5A)

[MIM:270800]. Spastic paraplegia is a neurodegenerative disorder characterized by a slow, gradual, progressive weakness and spasticity of the lower limbs. Rate of progression and the severity of symptoms are quite variable. Initial symptoms may include difficulty with balance, weakness and stiffness in the legs, muscle spasms, and dragging the toes when walking. In some forms of the disorder, bladder symptoms (such as incontinence) may appear, or the weakness

and stiffness may spread to other parts of the body.

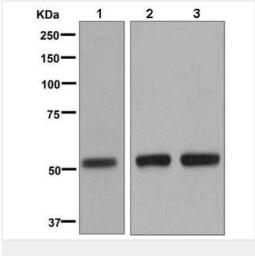
Defects in CYP7B1 are the cause of congenital bile acid synthesis defect type 3 (CBAS3) [MIM:613812]. Clinical features include severe cholestasis, cirrhosis and liver synthetic failure.

Hepatic microsomal oxysterol 7-alpha-hydroxylase activity is undetectable.

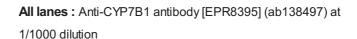
序列相似性 Belongs to the cytochrome P450 family.

细**胞定位** Endoplasmic reticulum membrane. Microsome membrane.

图片



Western blot - Anti-CYP7B1 antibody [EPR8395] (ab138497)



Lane 1: PC3 lysate

Lane 2 : Human fetal liver lysate

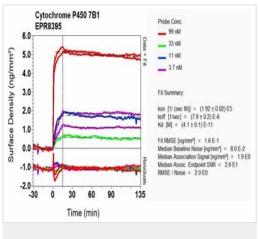
Lane 3 : Human fetal kidney lysate

Lysates/proteins at 30 µg per lane.

Secondary

All lanes: HRP labelled goat anti-rabbit at 1/2000 dilution

Predicted band size: 58 kDa **Observed band size:** 55 kDa



Ol-RD Scanning - Anti-CYP7B1 antibody [EPR8395] (ab138497)

Equilibrium disassociation constant (K_D)

Learn more about K_D

Click here to learn more about K_D



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