abcam

Product datasheet

Anti-CYP7B1 antibody [EPR8395] ab138497



★★★★★ 1 Abreviews 10 References 3 Images

Overview

Product name Anti-CYP7B1 antibody [EPR8395]

Description Rabbit monoclonal [EPR8395] to CYP7B1

Host species Rabbit

Tested applications Suitable for: WB

Unsuitable for: Flow Cyt,ICC/IF,IHC-P or IP

Species reactivity Reacts with: Human

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

Positive control PC3, Human fetal liver, and Human fetal kidney lysates

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**.

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

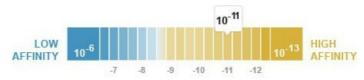
these species. Please contact us for more information.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

 $K_D = 4.10 \times 10^{-11} M$ Dissociation constant (K_D)



Learn more about K_D

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

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

supernatant

Purity Tissue culture supernatant

ClonalityMonoclonalClone numberEPR8395

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab138497 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)	1/1000 - 1/10000. Detects a band of approximately 55 kDa (predicted molecular weight: 58 kDa).

Application notes Is unsuitable for Flow Cyt,ICC/IF,IHC-P or IP.

Target

Tissue specificity Brain, testis, ovary, prostate, liver, colon, kidney, and small intestine.

Pathway Lipid metabolism; bile acid biosynthesis.

Involvement in disease 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.

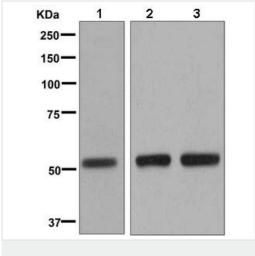
Defects in CYP7B1 are the cause of congenital bile acid synthesis defect type 3 (CBAS3) [MIM:613812]. Clinical features include severe cholestasis, cirrhosis and liver synthetic failure.

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

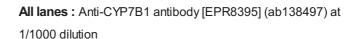
Sequence similaritiesBelongs to the cytochrome P450 family.

Cellular localization Endoplasmic reticulum membrane. Microsome membrane.

Images



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



Lane 1: PC3 lysate

Lane 2 : Human fetal liver lysate

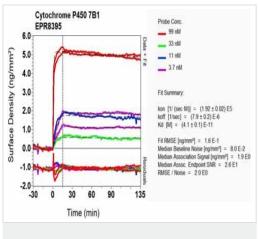
Lane 3 : Human fetal kidney lysate

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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