abcam

Product datasheet

Anti-CPT1A antibody [8F6AE9] ab128568



★★★★★ 5 Abreviews 148 References 7 图像

概述

产品名称 Anti-CPT1A抗体[8F6AE9]

小鼠单克隆抗体[8F6AE9] to CPT1A

宿主 Mouse

经测试应用 适用于: Flow Cyt, IHC-P, WB, ICC/IF, In-Cell ELISA

种属反应性 与反应: Mouse, Rat, Human

免疫原 Recombinant fragment, corresponding to amino acids 489-773 of Human CPT1A.

阳性对照 Partial Human Recombinant CPT1A protein; HepG2, H9C2, H4IIE Whole Cell Lysates; Rat heart

mitochondrial, Human liver, Rat liver and Mouse liver homogenates; H9C2 cells; HeLa cells IHC-

P: human normal kidney FFPE tissue sections

常规说明 This monoclonal antibody to CPT1A has been knockout validated in Western blot. The expected

band for CPT1A was observed in wild type cells and the band was not seen in CPT1A knockout

cells.

This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or

conjugation for your experiments, please contact orders@abcam.com.

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. Do Not Freeze.

存储溶液 pH: 7.4

Preservative: 0.02% Sodium azide Constituent: HEPES buffered saline

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纯**度** Ammonium Sulphate Precipitation

纯**化**说明 Purity is near homogeneity as judged by SDS-PAGE. The antibody was produced in vitro using

hybridomas grown in serum-free medium, and then concentrated by ammonium sulfate

precipitation.

 克隆
 单克隆

 克隆编号
 8F6AE9

 同种型
 IgG2b

 轻链类型
 kappa

应用

The Abpromise guarantee Abpromise™承诺保证使用ab128568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
Flow Cyt		Use at an assay dependent concentration.
IHC-P	★★★★ ☆ (1)	Use a concentration of 0.5 μ g/ml. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
WB	★★★★★ (3)	Use a concentration of 1 µg/ml. Predicted molecular weight: 88 kDa.
ICC/IF	*** <u>*</u>	Use a concentration of 0.5 - 5 µg/ml.
In-Cell ELISA		Use a concentration of 0.1 - 1 µg/ml.

靶标

组织特异性 Strong expression in kidney and heart, and lower in liver and skeletal muscle.

通路 Lipid metabolism; fatty acid beta-oxidation.

疾病相关 Defects in CPT1A are the cause of carnitine palmitoy/transferase 1A deficiency (CPT1AD)

[MIM:255120]; also known as CPT-I deficiency or CPT1A deficiency. CPT1AD is a rare

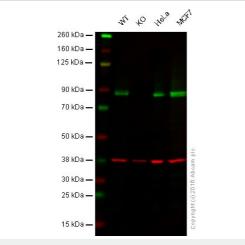
autosomal recessive metabolic disorder of long-chain fatty acid oxidation characterized by severe episodes of hypoketotic hypoglycemia usually occurring after fasting or illness. Onset is in infancy

or early childhood.

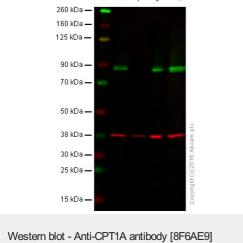
序列相似性 Belongs to the carnitine/choline acetyltransferase family.

细**胞定位** Mitochondrion outer membrane.

图片



(ab128568)



Immunocytochemistry/ Immunofluorescence - Anti-CPT1A antibody [8F6AE9] (ab128568)

Lane 1: Wild-type HAP1 cell lysate (20 µg)

Lane 2: CPT1A knockout HAP1 cell lysate (20 µg)

Lane 3: HeLa cell lysate (20 µg)

Lane 4: MCF-7 cell lysate (20 µg)

Lanes 1 - 4: Merged signal (red and green). Green - ab128568 observed at 88 kDa. Red - loading control, ab181602, observed at 37 kDa.

ab128568 specifically detected the expected band for CPT1A in wild-type HAP1 cells. No band was observed in CPT1A knockout cells. Wild-type and CPT1A knockout samples were subjected to SDS-PAGE. ab128568 and ab181602 (loading control to GAPDH) were diluted at 1 µg/mL and 1/10000 respectively and incubated overnight at 4°C. Blots were developed with Goat anti-Mouse IgG H&L (IRDye[®] 800CW) preadsorbed (<u>ab216772</u>) and Goat Anti-Rabbit lgG H&L (IRDye® 680RD) preadsorbed (ab216777) secondary antibodies at 1/10000 dilution for 1 hour at room temperature before imaging.

Immunofluorescent staining of CPT1A in H9C2 cells (rat) using ab128568.

Fixation: 4% paraformaldehyde PBS fixed for 20 minutes

Permebilization: 0.1% Triton X-100 PBS for 30 minutes at room

temperature while rocking

Blocking: 2x Sigma Block 0.1% Triton X-100 H2O for 2 hours at

room temperature while rocking

Primary antibodies: Anti-CPT1A antibody (ab128568) 0.5 ug/mL 1x Sigma Block with 0.1% Triton X-100 incubated overnight at 4 °C.

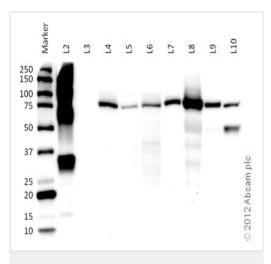
Washing: 3x 1% NGS 10 minutes/wash.

Secondary antibodies: Alexa 488 GAM 1:1000 diluted in 1% NGS with 0.1% Triton X-100 PBS incubated for 2 hours at room temperature while rocking.

Washing: 3x 1% NGS 10 minutes/wash.

DAPI: 20 ng/mL in 1% NGS, 0.1% Triton X-100 PBS.

Washing: 1x 1% NGS 10 minutes/wash.



Western blot - Anti-CPT1A antibody [8F6AE9] (ab128568)

All lanes: Anti-CPT1A antibody [8F6AE9] (ab128568) at 1 μg/ml

Lane 1: Marker

Lane 2: Partial Human Recombinant CPT1A protein (ab128569)

at 0.1 µg

Lane 3: Full Length Human Recombinant OTC protein at 0.1 µg

Lane 4: HepG2 (Human hepatocellular carcinoma cell line) Whole

Cell Lysate at 20 µg

Lane 5: H9C2 (Rat cardiomyoblast cell line) Whole Cell Lysate at

20 µց

Lane 6: H4IIE (Rat hepatoma cell line) Whole Cell Lysate at 20 µg

Lane 7: RHM (Rat heart mitochondrial homogenate) at 20 µg

Lane 8: HLH (Human liver homogenate) at 20 µg

Lane 9: RLH (Rat liver homogenate) at 20 µg

Lane 10: MLH (Mouse liver homogenate) at 20 µg

Secondary

All lanes: Goat polyclonal to Mouse IgG - H&L - Pre-Adsorbed

(HRP) at 1/10000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 88 kDa Observed band size: 88 kDa

Exposure time: 1 minute

Predicted partial recombinant protein band size: 32 kDa

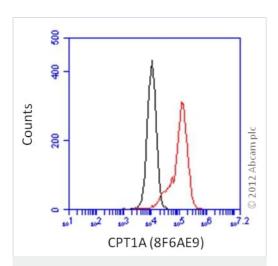
Observed band size: 32 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CPT1A antibody [8F6AE9] (ab128568)

IHC image of CPT1A staining in human normal kidney formalin fixed paraffin embedded tissue section, performed on a Leica BondTM system using the standard protocol F. The section was pretreated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab128568, 0.5µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.



Flow Cytometry - Anti-CPT1A antibody [8F6AE9] (ab128568)

Overlay histogram showing HeLa cells stained with ab128568 (red line) or no primary antibody (black line). The cells were fixed with 4% paraformaldehyde (15 min) and then permeabilized with 0.1% Triton X-100 in PBS, 3% BSA for 10 min. The cells were then incubated in 3% BSA in PBS for 10 minutes to block non-specific protein-protein interactions followed by the antibody (ab128568, 1µg/mL) for 60 min at 22°C. The secondary antibody used was DyLight® 488 goat anti-mouse lgG (H+L) (ab96871) at 1/1000 dilution for 30 min.



Western blot - Anti-CPT1A antibody [8F6AE9] (ab128568)

All lanes: Anti-CPT1A antibody [8F6AE9] (ab128568) at 1 µg/ml

Lane 1: Marker

Lane 2 : HHH (Human heart homogenate) RIPA Extract

Immunoprecipitated with no primary antibody

Lane 3 : HHH (Human heart homogenate) RIPA Extract Immunoprecipitated with Anti-CPT1A antibody 8F6AE9 (ab128568)

Lane 4: HHH (Human heart homogenate) RIPA Extract 20 µg

Secondary

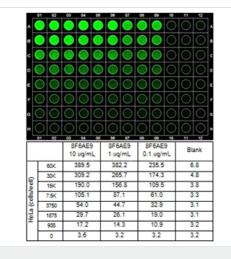
All lanes : Goat polyclonal to Mouse $\lg G - H\&L - Pre-Adsorbed$ (HRP) at 1:10000

Developed using the ECL technique.

Performed under non-reducing conditions.

Predicted band size: 88 kDa

Exposure time: 1 minute



In-Cell ELISA - Anti-CPT1A antibody [8F6AE9] (ab128568)

In-Cell ELISA for Anti-CPT1A antibody (ab128568) stained HeLa cells (human)

Seeding: HeLa cells seeded in a 1:2 dilution series starting at 60,000 cells/well across Row A, 30,000 cells/well across Row B, etc. Row H contains no cells.

Table: Example In-Cell ELISA Average Data from shown plate.

Fixation: 4% paraformaldehyde PBS fixed for 15 minutes

Permeabilization: 0.3% Triton X-100 PBS for 30 minutes at room

temperature while shaking

Blocking: 2x Sigma Block 0.3% Triton X-100 H2O for 2 hours at

room temperature while shaking

Primary antibodies: All primaries diluted in 1x Sigma Block with 0.3% Triton X-100 incubated overnight at 4 °C.

- Columns 1-3: Anti-CPT1A antibody (ab128568) 10 ug/mL
- Columns 4-6: Anti-CPT1A antibody (ab128568) 1 ug/mL
- Columns 7-9: Anti-CPT1A antibody (ab128568) 0.1 ug/mL
- Columns 10-12: No Primary

Washing: Briefly 4x with 0.3% TWEEN-20 PBS

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