abcam

Product datasheet

Anti-Acetyl Coenzyme A Carboxylase antibody [EP687Y] - BSA and Azide free ab173584





重组 RabMAb

7 图像

概述

产品名称 Anti-Acetyl Coenzyme A Carboxylase抗体[EP687Y] - BSA and Azide free

描述 兔单克隆抗体[EP687Y] to Acetyl Coenzyme A Carboxylase - BSA and Azide free

宿主 Rabbit

特异性 Acetyl Coenzyme A Carboxylase is highly expressed in lipogenic tissues such as liver, adipose,

and lactating mammary gland, and its activities are regulated at various levels [Proc Natl Acad

Sci U S A. 2003 Jun 24;100(13):7515-20.].

经测试应用 适用于: WB. IHC-P

种属反应性 与反应: Mouse, Rat, Human

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

阳性对照 IHC-P: Human breast carcinoma, Mouse stomach and Rat kidney tissue.

常规说明 ab173584 is the carrier-free version of ab45174.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar[®] Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] is a trademark of Fluidigm Canada Inc.

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**.

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C. Do Not Freeze.

存储溶液 Constituent: PBS

无载体 是

纯**度** Protein A purified

 克隆
 单克隆

 克隆编号
 EP687Y

 同种型
 IgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab173584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 at an assay dependent concentration. Detects a band of approximately 265 kDa (predicted molecular weight: 265 kDa). Can be blocked with Acetyl Coenzyme A Carboxylase peptide (ab195232).
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. See IHC antigen retrieval protocols.

靶标

功能 Catalyzes the rate-limiting reaction in the biogenesis of long-chain fatty acids. Carries out three

functions: biotin carboxyl carrier protein, biotin carboxylase and carboxyltransferase.

组织特异性 Expressed in brain, placental, skeletal muscle, renal, pancreatic and adipose tissues; expressed

at low level in pulmonary tissue; not detected in the liver.

通路 Lipid metabolism; malonyl-CoA biosynthesis; malonyl-CoA from acetyl-CoA: step 1/1.

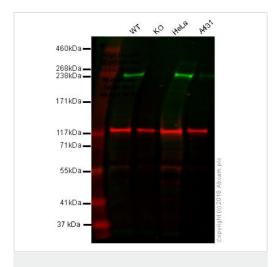
疾病相关 Acetyl-CoA carboxylase 1 deficiency

序列相似性 Contains 1 ATP-grasp domain.

Contains 1 biotin carboxylation domain.
Contains 1 biotinyl-binding domain.
Contains 1 carboxyltransferase domain.

翻译后修饰 Phosphorylation on Ser-1263 is required for interaction with BRCA1.

图片



Western blot - Anti-Acetyl Coenzyme A

Carboxylase antibody [EP687Y] - BSA and Azide
free (ab173584)

All lanes : Anti-Acetyl Coenzyme A Carboxylase antibody [EP687Y] (ab45174) at 1/2000 dilution (unpurified)

Lane 1: Wild-type HAP1 whole cell lysate

Lane 2: ACACA (Acetyl Coenzyme A Carboxylase) knockout

HAP1 whole cell lysate

Lane 3: HeLa whole cell lysate

Lane 4: A431 whole cell lysate

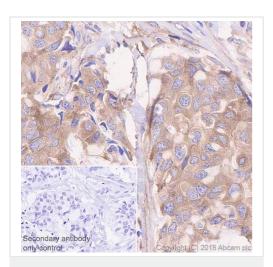
Lysates/proteins at 20 µg per lane.

Predicted band size: 265 kDa

Lanes 1 - 4: Merged signal (red and green). Green - <u>ab45174</u> observed at 265 kDa. Red - loading control, <u>ab130007</u>, observed at 130 kDa.

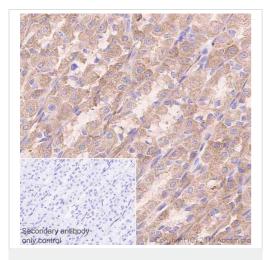
ab45174 was shown to specifically react with Acetyl Coenzyme A Carboxylase in wild-type HAP1 cells as signal was lost in ACACA (Acetyl Coenzyme A Carboxylase) knockout cells. Wild-type and ACACA (Acetyl Coenzyme A Carboxylase) knockout samples were subjected to SDS-PAGE. Ab45174 and ab130007 (Mouse anti-Vinculin loading control) were incubated overnight at 4°C at 1/2000 dilution and 1/20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preabsorbed ab216773 and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed ab216776 secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab45174).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Acetyl Coenzyme A
Carboxylase antibody [EP687Y] - BSA and Azide
free (ab173584)

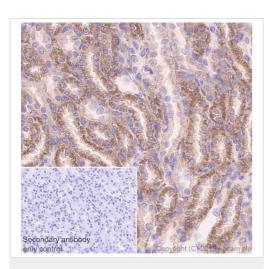
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human breast carcinoma tissue sections labeling Acetyl Coenzyme A Carboxylase with purified ab45174 at 1/400 dilution (2.06 µg/mL). Heat mediated antigen retrieval was performed using ab93684 (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain. This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab45174)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Acetyl Coenzyme A
Carboxylase antibody [EP687Y] - BSA and Azide
free (ab173584)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of rat stomcah tissue sections labeling Acetyl Coenzyme A Carboxylase with purified <u>ab45174</u> at 1/400 dilution (2.06 µg/mL). Heat mediated antigen retrieval was performed using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.

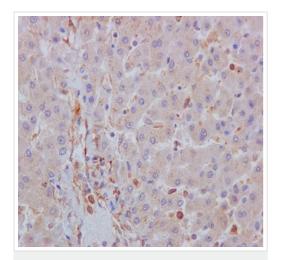
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab45174)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Acetyl Coenzyme A
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Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of mouse kidney tissue sections labeling Acetyl Coenzyme A Carboxylase with purified <u>ab45174</u> at 1/400 dilution (2.06 µg/mL). Heat mediated antigen retrieval was performed using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.

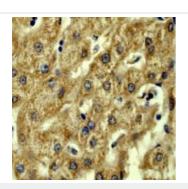
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab45174)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Acetyl Coenzyme A
Carboxylase antibody [EP687Y] - BSA and Azide
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Immunohistochemical analysis of paraffin-embedded human liver tissue labeling Acetyl Coenzyme A Carboxylase with ab45174 (unpurified) at 1/250 dilution followed by Goat Anti-Rabbit lgG H&L (HRP) (ab97051, 1/500). Counter stained with hematoxylin. Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.

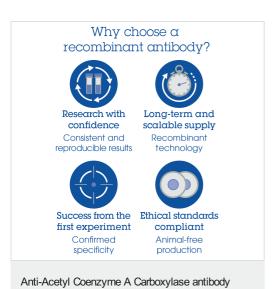
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab45174).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Acetyl Coenzyme A
Carboxylase antibody [EP687Y] - BSA and Azide
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<u>ab45174</u> (unpurified), at a dilution of 1/50, staining human Acetyl Coenzyme A Carboxylase in human liver by immunohistochemistry using paraffin embedded tissue. Heat mediated antigen retrieval was perfromed with citrate buffer pH 6 before commencing with IHC staining protocol.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab45174).



[EP687Y] - BSA and Azide free (ab173584)

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