# abcam

# Product datasheet

# Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] ab269273

Recombinant RabMAb

1 References 12 Images

Overview

Product name Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147]

**Description**Rabbit monoclonal [EPR23235-147] to Acetyl Coenzyme A carboxylase alpha

Host species Rabbit

Tested applications Suitable for: IHC-P, Flow Cyt (Intra), IP, ICC/IF, WB

Unsuitable for: IHC-Fr

Species reactivity Reacts with: Mouse, Rat, Human

**Immunogen** Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: 293T, HeLa, PC-12 and C2C12 lysates. IHC-P: Human breast carcinoma and endometrial

carcinoma tissue. Mouse and rat lung tissue. ICC/IF: C2C12 and 293T cells. Flow Cyt (intra):

C2C12 and 293T cells. IP: C2C12 and 293T cells.

**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<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**<sup>®</sup> **patents**.

**Properties** 

**Form** Liquid

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

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

Purity Protein A purified

1

**Clonality** Monoclonal

Clone number EPR23235-147

**Isotype** IgG

#### **Applications**

The Abpromise guarantee Our Abpromise guarantee covers the use of ab269273 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
IHC-P		1/500. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
Flow Cyt (Intra)		1/500.
IP		1/30.
ICC/IF		1/50.
WB		1/2000. Predicted molecular weight: 265 kDa.

**Application notes** Is unsuitable for IHC-Fr.

**Target** 

Function 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.

Tissue specificity Expressed in brain, placental, skeletal muscle, renal, pancreatic and adipose tissues; expressed

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

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

Involvement in disease Defects in ACACA are a cause of acetyl-CoA carboxylase 1 deficiency (ACACAD)

[MIM:200350]; also known as ACAC deficiency or ACC deficiency. An inborn error of de novo fatty acid synthesis associated with severe brain damage, persistent myopathy and poor growth.

Sequence similarities Contains 1 ATP-grasp domain.

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

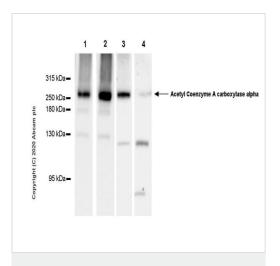
Post-translational

modifications

Phosphorylation on Ser-1263 is required for interaction with BRCA1.

Cellular localization Cytoplasm.

## Images



Western blot - Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)

**All lanes :** Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273) at 1/2000 dilution

**Lane 1**: 293T (human embryonic kidney epithelial cell), whole cell lysate, in the loading buffer containing double DTT

Lane 2: HeLa (human cervix adenocarcinoma epithelial cell), whole cell lysate, in the loading buffer containing double DTT

**Lane 3**: PC-12 (rat adrenal gland pheochromocytoma), whole cell lysate, in the loading buffer containing double DTT

**Lane 4**: C2C12 (mouse myoblasts myoblast), whole cell lysate, in the loading buffer containing double DTT

Lysates/proteins at 20 µg per lane.

### **Secondary**

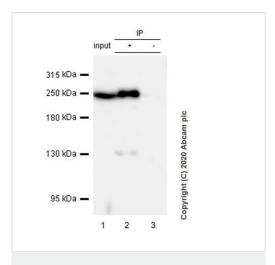
**All lanes :** Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated (**ab97051**) at 1/100000 dilution

**Predicted band size:** 265 kDa **Observed band size:** 265 kDa

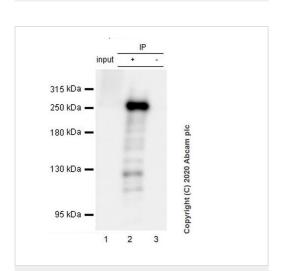
Blocking and diluting buffer and concentration: 5% NFDM/TBST.

This blot of lane 3, 4 was developed using a higher sensitivity ECL substrate.

Exposure time: Lane 1, 2: 26 secs; Lane 3, 4: 3 mins.



Immunoprecipitation - Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)



Immunoprecipitation - Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)

Acetyl Coenzyme A carboxylase alpha was immunoprecipitated from 0.35 mg C2C12 (mouse myoblasts myoblast), whole cell lysate 10 µg with ab269273 at 1/30 dilution (2µg in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab269273 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP)(ab131366) was used at 1/5000 dilution.

Lane 1: C2C12 whole cell lysate 10 µg.

Lane 2: ab269273 IP in C2C12 whole cell lysate 10 μg.

**Lane 3:** Rabbit monoclonal lgG (<u>ab172730</u>) instead of ab269273 in C2C12 whole cell lysate.

Blocking and dilution buffer and concentration: 5% NFDM/TBST.

Exposure time: 5 seconds.

Acetyl Coenzyme A carboxylase alpha was immunoprecipitated from 0.35 mg 293T (human embryonic kidney epithelial cell), whole cell lysate 10  $\mu$ g with ab269273 at 1/30 dilution (2 $\mu$ g in 0.35 mg lysates). Western blot was performed on the immunoprecipitate using ab269273 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP)(ab131366) was used at 1/5000 dilution.

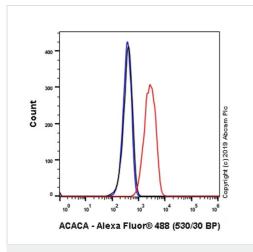
Lane 1: 293T whole cell lysate 10 µg.

Lane 2: ab269273 IP in 293T whole cell lysate.

**Lane 3:** Rabbit monoclonal lgG (<u>ab172730</u>) instead of ab269273 in 293T whole cell lysate.

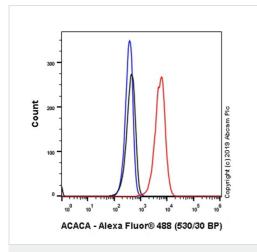
Blocking and dilution buffer and concentration: 5% NFDM/TBST.

Exposure time: 5 seconds.



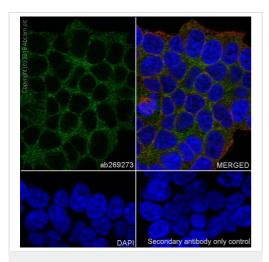
Flow Cytometry (Intracellular) - Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)

Intracellular flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized C2C12 (Mouse myoblasts myoblast) cells labelling Acetyl Coenzyme A carboxylase alpha with ab269273 at 1/500 dilution (0.1µg) (Red) compared with a Rabbit monoclonal IgG (ab172730) (Black) isotype control and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat anti rabbit IgG (Alexa Fluor®488, ab150077) at 1/2000 dilution was used as the secondary antibody.



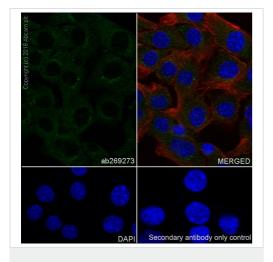
Flow Cytometry (Intracellular) - Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)

Intracellular flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized 293T (Human embryonic kidney epithelial cell) cells labelling Acetyl Coenzyme A carboxylase alpha with ab269273 at 1/500 dilution (0.1µg) (Red) compared with a Rabbit monoclonal IgG (ab172730) (Black) isotype control and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat anti rabbit IgG (Alexa Fluor®488, ab150077) at 1/2000 dilution was used as the secondary antibody.



Immunocytochemistry/ Immunofluorescence - Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)

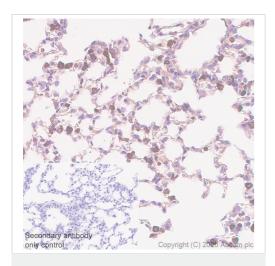
Secondary antibody only control: Secondary antibody is <u>ab150077</u> Goat Anti-Rabbit lgG H&L (Alexa Fluor<sup>®</sup>488) at 1/1000 dilution.



Immunocytochemistry/ Immunofluorescence - Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized C2C12 cells labelling Acetyl Coenzyme A carboxylase alpha with ab269273 at 1/50 dilution, followed by **ab150077** Goat Anti-Rabbit IgG H&L (Alexa Fluor<sup>®</sup>488) antibody at 1/1000 2 µg/ml dilution (Green). Confocal image showing cytoplasmic staining in C2C12 cell line. **ab195889** Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor<sup>®</sup>594) was used to counterstain tubulin at 1/200 2.5 µg/ml dilution (Red). The nuclear counterstain was DAPI (Blue).

Secondary antibody only control: Secondary antibody is <u>ab150077</u> Goat Anti-Rabbit lgG H&L (Alexa Fluor  $^{\tiny @}488$ ) at 1/1000 2  $\mu$ g/ml dilution.

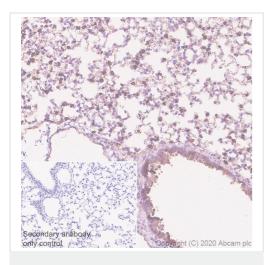


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)

Immunohistochemical analysis of paraffin-embedded rat lung tissue labeling Acetyl Coenzyme A carboxylase alpha with ab269273 at 1/500 dilution followed by ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101). Positive staining on rat lung (PMID: 17521700). The section was incubated with ab269273 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

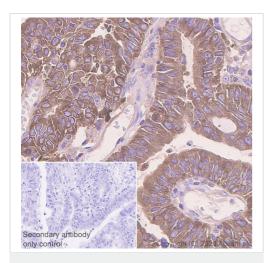


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)

Immunohistochemical analysis of paraffin-embedded mouse lung tissue labeling Acetyl Coenzyme A carboxylase alpha with ab269273 at 1/500 dilution followed by ready to use Rabbit specific IHC polymer detection kit HRP/DAB (<a href="mailto:ab209101">ab209101</a>). Positive staining on mouse lung (PMID: 17521700). The section was incubated with ab269273 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND<sup>®</sup>RX instrument. Counterstained with hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

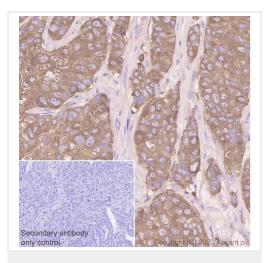


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)

Immunohistochemical analysis of paraffin-embedded human endometrial carcinoma tissue labeling Acetyl Coenzyme A carboxylase alpha with ab269273 at 1/500 dilution followed by ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101). Cytoplasmic staining on human endometrial carcinoma. The section was incubated with ab269273 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)

Immunohistochemical analysis of paraffin-embedded human breast carcinoma tissue labeling Acetyl Coenzyme A carboxylase alpha with ab269273 at 1/500 dilution followed by ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101). Cytoplasmic staining on human breast carcinoma (PMID: 21415164). The section was incubated with ab269273 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.



Anti-Acetyl Coenzyme A carboxylase alpha antibody [EPR23235-147] (ab269273)

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