abcam

Product datasheet

Anti-AMPK alpha 1 antibody [Y365] ab32047



★★★★★ 6 Abreviews 165 References 画像数 12

製品の概要

製品名 Anti-AMPK alpha 1 antibody [Y365]

製品の詳細 Rabbit monoclonal [Y365] to AMPK alpha 1

由来種 Rabbit

特異性 This antibody is specific for human AMPK alpha 1. This antibody shows low affinity on mouse and

適用あり: ICC/IF, Flow Cyt (Intra), WB, IHC-P, IP アプリケーション

種交差性 交差種: Mouse, Rat, Human

交差が予測される動物種: African green monkey 🔷

免疫原 Synthetic peptide within Human AMPK alpha 1 aa 500 to the C-terminus (C terminal). The exact

> sequence is proprietary. Database link: Q13131

ポジティブ・コントロール WB: HeLa, HepG2, C6, NIH/3T3 and MCF-7 cell lysate. Mouse liver, brain, retina, and skeletal

> muscle tissue lysates, RAW 264.7 cell lysate, Neuro2A cell lysate. IHC-P: Human cervical carcinoma and lung carcinoma tissues. ICC/IF: MCF-7 cells. Flow Cyt (intra): HeLa cells. IP: HeLa

whole cell lysate (ab150035).

特記事項 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**.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle.

バッファー pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol, 0.05% BSA

精製度 Protein A purified

ポリ/モノ モノクローナル

クローン名 Y365 アイソタイプ IgG

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab32047の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
ICC/IF		1/250.
Flow Cyt (Intra)		1/100 - 1/150. ab172730 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.
WB	**** <u>(5)</u>	1/1000 - 1/5000. Predicted molecular weight: 63 kDa. This antibody shows low affinity on mouse and rat samples.
IHC-P		1/100 - 1/250. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. See IHC antigen retrieval protocols. Mouse and rat species are recommended by WB application.
IP		1/40 - 1/50.

ターゲット情報

機能 Responsible for the regulation of fatty acid synthesis by phosphorylation of acetyl-CoA

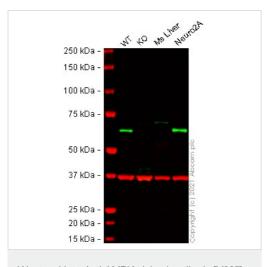
carboxylase. It also regulates cholesterol synthesis via phosphorylation and inactivation of hormone-sensitive lipase and hydroxymethylglutaryl-CoA reductase. Appears to act as a metabolic stress-sensing protein kinase switching off biosynthetic pathways when cellular ATP levels are depleted and when 5'-AMP rises in response to fuel limitation and/or hypoxia. This is a

catalytic subunit.

配列類似性 Belongs to the protein kinase superfamily. CAMK Ser/Thr protein kinase family. SNF1 subfamily.

Contains 1 protein kinase domain.

画像



Western blot - Anti-AMPK alpha 1 antibody [Y365] (ab32047)

All lanes : Anti-AMPK alpha 1 antibody [Y365] (ab32047) at 1/1000 dilution

Lane 1: Wild-type RAW 264.7 cell lysate

Lane 2: PRKAA1 knockout RAW 264.7 cell lysate

Lane 3: Mouse Liver cell lysate

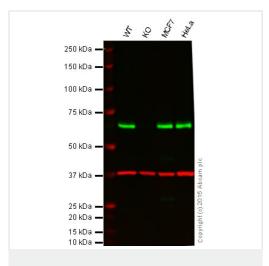
Lane 4: Neuro2A cell lysate

Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 63 kDa Observed band size: 64 kDa

False colour image of Western blot: Anti-AMPK alpha 1 antibody [Y365] staining at 1/1000 dilution, shown in green; Mouse anti-GAPDH antibody [6C5] (ab8245) loading control staining at 1/20000 dilution, shown in red. In Western blot, ab32047 was shown to bind specifically to AMPK alpha 1. A band was observed at 64 kDa in wild-type RAW 264.7 cell lysates with no signal observed at this size in PRKAA1 knockout cell line ab280055 (knockout cell lysate ab280114). To generate this image, wild-type and PRKAA1 knockout RAW 264.7 cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3 % milk in TBS-0.1 % Tween[®] 20 (TBS-T) before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L (IRDye® 800CW) preabsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed (ab216776) at 1/20000 dilution.



Western blot - Anti-AMPK alpha 1 antibody [Y365] (ab32047)

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

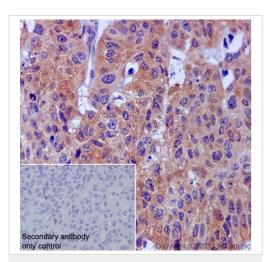
Lane 2: AMPK alpha knockout HAP1 cell lysate (20 µg)

Lane 3: MCF7 cell lysate (20 µg)

Lane 4: HeLa cell lysate (20 µg)

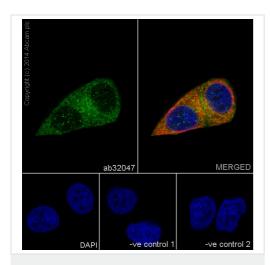
Lanes 1 - 4: Merged signal (red and green). Green - ab32047 observed at 63 kDa. Red - loading control, <u>ab8245</u>, observed at 37 kDa.

ab32047 was shown to specifically react with AMPK alpha in wild-type HAP1 cells. No band was observed when AMPK alpha knockout samples were examined. Wild-type and AMPK alpha knockout samples were subjected to SDS-PAGE. ab32047 and ab8245 (loading control to GAPDH) were diluted 1/5000 and 1/2000 respectively and incubated overnight at 4°C. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1/10,000 dilution for 1hr at room temperature before imaging.

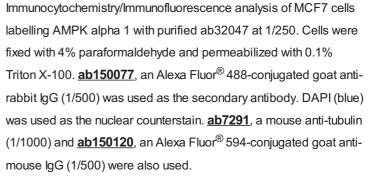


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-AMPK alpha 1 antibody [Y365] (ab32047)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human lung carcinoma tissue labelling AMPK alpha 1 with purified ab32047 at 1/100. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9 before commencing with IHC staining protocol. ab97051, a HRP-conjugated goat anti-rabbit IgG (H+L) was used as the secondary antibody (1/500). Negative control using PBS instead of primary antibody. Counterstained with hematoxylin.

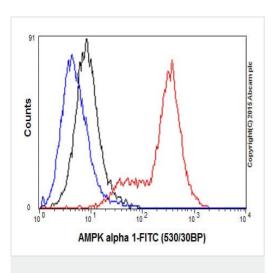


Immunocytochemistry/ Immunofluorescence - Anti-AMPK alpha 1 antibody [Y365] (ab32047)



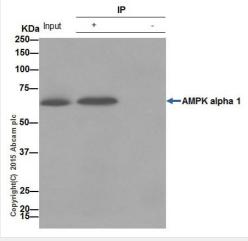
Control 1: primary antibody (1/250) and secondary antibody, **ab150120**, an Alexa Fluor[®] 594-conjugated goat anti-mouse IgG (1/500).

Control 2: <u>ab7291</u> (1/1000) and secondary antibody, <u>ab150077</u>, an Alexa Fluor[®] 488-conjugated goat anti-rabbit lgG (1/500).



Flow Cytometry (Intracellular) - Anti-AMPK alpha 1 antibody [Y365] (ab32047)

Intracellular Flow Cytometry analysis of HeLa cells labelling AMPK alpha 1 with purified ab32047 at 1/150 (red). Cells were fixed with 2% paraformaldehyde. A FITC-conjugated goat anti-rabbit lgG (1/150) was used as the secondary antibody. Black - Isotype control, rabbit monoclonal lgG. Blue - Unlabelled control, cells without incubation with primary and secondary antibodies.



Immunoprecipitation - Anti-AMPK alpha 1 antibody [Y365] (ab32047)

All lanes: Anti-AMPK alpha 1 antibody [Y365] (ab32047) at 1/500 dilution

ab32047 (purified) at 1/40 immunoprecipitating AMPK alpha 1 in

Lane 3 (-): Rabbit monoclonal lgG (ab172730) instead of ab32047

For western blotting, ab131366 VeriBlot for IP Detection Reagent

HeLa whole cell lysate.

in HeLa whole cell lysate.

Lane 1 (input): HeLa whole cell lysate (10µg)

(HRP) was used for detection (1/1500).

Lane 2 (+): ab32047 + HeLa whole cell lysate (10µg).

Blocking buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM /TBST.

Lane 1: HepG2 (human liver hepatocellular carcinoma cell line) whole cell lysate

Lane 2: NIH/3T3 (Mouse embryonic fibroblast) whole cell lysate

Lane 3: Mouse liver tissue lysate

Lane 4: Mouse brain tissue lysate Lane 5: Mouse kidney tissue lysate

Lane 6: Mouse retina tissue lysate

Lane 7: Mouse skeletal muscle tissue lysate

Lysates/proteins at 20 µg per lane.

250 kDa -150 kDa -100 kDa -75 kDa -37 kDa -20 kDa -15 kDa 🕳 (ab181602)

Western blot - Anti-AMPK alpha 1 antibody [Y365] (ab32047)

Secondary

All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

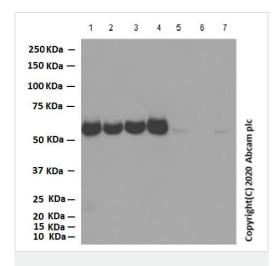
Predicted band size: 63 kDa Observed band size: 63 kDa

Additional bands at: 40 kDa (possible non-specific binding)

Exposure time: 3 minutes

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

This antibody shows low affinity on mouse samples.



Western blot - Anti-AMPK alpha 1 antibody [Y365] (ab32047)

All lanes : Anti-AMPK alpha 1 antibody [Y365] (ab32047) at 1/20000 dilution

Lane 1 : MCF7 (Human breast adenocarcinoma epithelial cell) whole cell lysate

Lane 2 : HepG2 (Human hepatocellular carcinoma epithelial cell) whole cell lysate

Lane 3 : HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate

Lane 4: K-562 (Human chronic myelogenous leukemia lymphoblast) whole cell lysate

Lane 5: C6 (Rat glial tumor glial cell) whole cell lysate

Lane 6 : Neuro-2a (Mouse neuroblastoma neuroblast) whole cell lysate

Lane 7: NIH/3T3 (Mouse embryonic fibroblast) whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

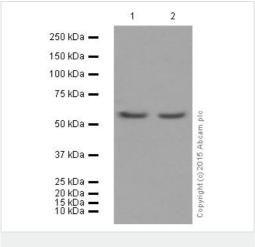
All lanes : Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

Predicted band size: 63 kDa **Observed band size:** 63 kDa

Exposure time: 180 seconds

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

This antibody shows low affinity on mouse and rat samples.



Western blot - Anti-AMPK alpha 1 antibody [Y365] (ab32047)

All lanes : Anti-AMPK alpha 1 antibody [Y365] (ab32047) at 1/2000 dilution (purified)

Lane 1: C6 cell lysate

Lane 2: NIH/3T3 cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

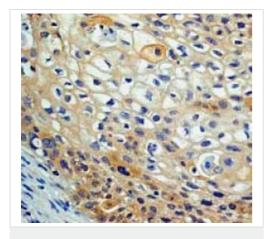
 $\textbf{All lanes:} \ Peroxidase-conjugated goat anti-rabbit \ lgG, \ (H+L) \ at$

1/1000 dilution

Predicted band size: 63 kDa **Observed band size:** 63 kDa

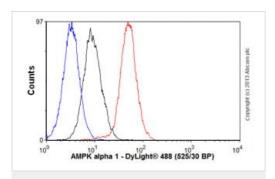
Blocking and dilution buffer: 5% NFDM/TBST.

This antibody shows low affinity on mouse and rat samples.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-AMPK alpha 1 antibody [Y365] (ab32047)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analaysis of human cervical carcinoma tissue labelling AMPK alpha 1 with unpurified ab32047 at a dilution of 1/100. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Flow Cytometry (Intracellular) - Anti-AMPK alpha 1 antibody [Y365] (ab32047)

Overlay histogram showing HeLa cells stained with unpurified ab32047 (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (unpurified ab32047, 1/100 dilution) for 30 min at 22°C. The secondary antibody used was DyLight[®] 488 goat antirabbit lgG (H+L) (ab96899) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit lgG (monoclonal) (1µg/1x10⁶ cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.



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