abcam

Product datasheet

Anti-Syntenin antibody [EPR8102] ab133267





重组 RabMAb

★★★★★ 1 Abreviews 68 References 14 图像

概述

产品名称 Anti-Syntenin抗体[EPR8102]

描述 兔单克隆抗体[EPR8102] to Syntenin

宿主 Rabbit

经测试应用 适用于: Flow Cyt (Intra), WB, IP, IHC-P, ICC/IF

种属反应性 与反应: Human

免疫原 Synthetic peptide within Human Syntenin aa 1-100. The exact sequence is proprietary.

Database link: **000560**

阳性对照 Human fetal brain lysate, Human fetal heart lysate, Human placenta lysate, HeLa, 293T and A549

(Human lung carcinoma cell line) cell lysates; Human brain tissue

This product is a recombinant monoclonal antibody, which offers several advantages including: 常规说明

- 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.

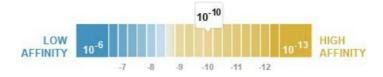
性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle.

 $K_D = 1.24 \times 10^{-10} M$ 解离常数(Kn)



Learn more about K_D

存储溶液 pH: 7.20

Preservative: 0.01% Sodium azide

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

纯**度** Protein A purified

克隆 单克隆

克隆编号 EPR8102

同种型 lgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab133267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 (Intra)		1/50. ab172730 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody. For unpurified use at 1/1000 - 1/10000.
WB	★★★☆☆ (1)	1/1000 - 1/10000. Detects a band of approximately 32 kDa (predicted molecular weight: 32 kDa).
IP		1/10 - 1/100.
IHC-P		1/50 - 1/100. Perform heat mediated antigen retrieval before commencing with IHC staining protocol. See IHC antigen retrieval protocols.
ICC/IF		1/50 - 1/200.

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功能 Seems to function as an adapter protein. In adherens junctions may function to couple syndecans

to cytoskeletal proteins or signaling components. Seems to couple transcription factor SOX4 to the IL-5 receptor (IL5RA). May also play a role in vesicular trafficking. Seems to be required for

the targeting of TGFA to the cell surface in the early secretory pathway.

组织特异性 Widely expressed. Expressed in fetal kidney, liver, lung and brain. In adult highest expression in

heart and placenta.

序列相似性 Contains 2 PDZ (DHR) domains.

翻译**后修**饰 Phosphorylated on tyrosine residues.

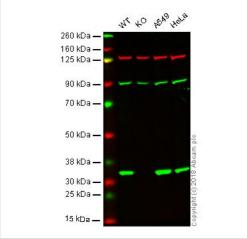
细胞定位 Cell junction > focal adhesion. Cell junction > adherens junction. Cell membrane. Endoplasmic

reticulum membrane. Nucleus. Melanosome. Cytoplasm > cytosol. Cytoplasm > cytoskeleton.

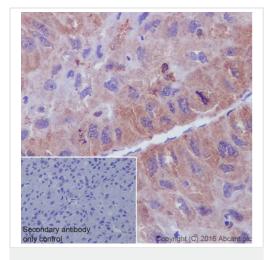
Mainly membrane-associated. Localized to adherens junctions, focal adhesions and endoplasmic

reticulum. Colocalized with actin stress fibers. Also found in the nucleus. Identified by mass

图片



Western blot - Anti-Syntenin antibody [EPR8102] (ab133267)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Syntenin antibody [EPR8102] (ab133267)

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

Lane 2: Syntenin knockout HAP1 whole cell lysate (20 µg)

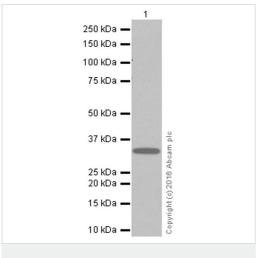
Lane 3: A549 whole cell lysate (20 µg)

Lane 4: HeLa whole cell lysate (20 µg)

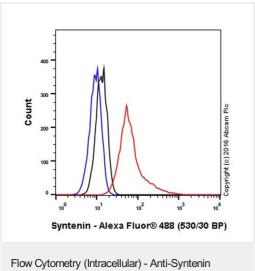
Lanes 1 - 4: Merged signal (red and green). Green - ab133267 observed at 32 kDa. Red - loading control, ab130007, observed at 130 kDa.

ab133267 was shown to recognize Syntenin in wild-type HAP1 cells as signal was lost at the expected MW in Syntenin knockout cells. Additional cross-reactive bands were observed in the wild-type and knockout cells. Wild-type and Syntenin knockout samples were subjected to SDS-PAGE. ab133267 and ab130007 (Mouse anti-Vinculin loading control) were incubated overnight at 4°C at 1/1000 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.

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Human hepatocellular carcinoma tissue labeling Syntenin with purified ab133267 at 1/50 dilution. Heat mediated antigen retrieval was performed using EDTA buffer pH 9. ab97051, a HRP-conjugated goat anti-rabbit lgG (H+L) was used as the secondary antibody (1/500). Negative control using PBS instead of primary antibody. Counterstained with hematoxylin



Western blot - Anti-Syntenin antibody [EPR8102] (ab133267)



Flow Cytometry (Intracellular) - Anti-Syntenin antibody [EPR8102] (ab133267)

Anti-Syntenin antibody [EPR8102] (ab133267) at 1/2000 dilution (purified) + A549 (Human lung carcinoma cell line) whole cell lysate at $15 \, \mu g$

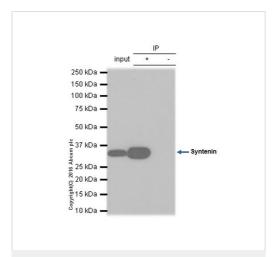
Secondary

Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution (Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated)

Predicted band size: 32 kDa **Observed band size:** 32 kDa

Blocking and dilution buffer: 5% NFDM/TBST

Intracellular Flow Cytometry analysis ofHeLa (Human epithelial cell line from cervix adenocarcinoma)labeling Syntenin with purifiedab133267 at 1/50 (red). Cells were fixed with 4% paraformaldehyde. A goat anti rabbit lgG (Alexa Fluor® 488) 1/2000 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-Syntenin antibody [EPR8102] (ab133267)

ab133267 (purified) at 1/40 immunoprecipitating Syntenin in HeLa whole cell lysate.

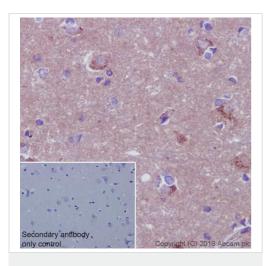
Lane 1 (input): HeLa whole cell lysate (10µg)

Lane 2 (+): ab133267 + HeLa whole cell lysate.

Lane 3 (-): Rabbit monoclonal $\lg G$ (ab172730) instead of ab133267 in HeLa whole cell lysate.

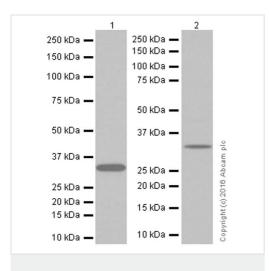
Blocking buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM /TBST.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Syntenin antibody
[EPR8102] (ab133267)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Human cerebral cortex tissue labeling Syntenin with purified ab133267 at 1/50 dilution. Heat mediated antigen retrieval was performed using EDTA buffer pH 9. 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.



Western blot - Anti-Syntenin antibody [EPR8102] (ab133267)

All lanes : Anti-Syntenin antibody [EPR8102] (ab133267) at 1/2000 dilution (purified)

Lane 1 : Human placenta lysate

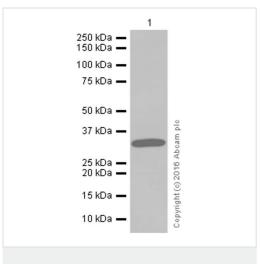
Lane 2 : Human fetal brain lysate

Lysates/proteins at 15 µg per lane.

Secondary

All lanes : Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/2000 dilution

Predicted band size: 32 kDa Observed band size: 32 kDa



Western blot - Anti-Syntenin antibody [EPR8102] (ab133267)

Blocking and dilution buffer: 5% NFDM/TBST

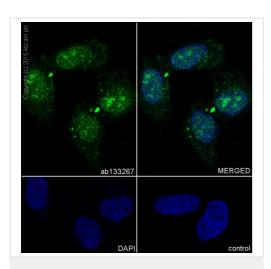
Anti-Syntenin antibody [EPR8102] (ab133267) at 1/10000 dilution (purified) + HeLa (Human epithelial cell line from cervix adenocarcinoma) whole cell lysate at 20 µg

Secondary

Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/2000 dilution

Predicted band size: 32 kDa Observed band size: 32 kDa

Blocking and dilution buffer: 5% NFDM/TBST

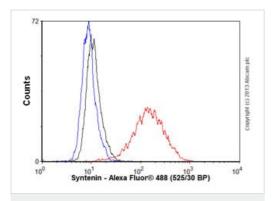


Immunocytochemistry/ Immunofluorescence - Anti-Syntenin antibody [EPR8102] (ab133267)

ab133267 staining Syntenin in HeLa (human cervix adenocarcinoma) cells by ICC/IF

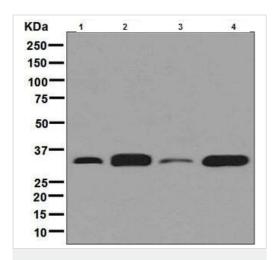
(Immunocytochemistry/immunofluorescence). Cells were fixed with 4% Paraformaldehyde and permeabilized with 0.1% Triton X-100. Samples were incubated with primary antibody at a dilution of 1/500. A goat anti rabbit IgG (Alexa Fluor® 488) (ab150077) was used as the secondary antibody at a dilution of 1/1000. DAPI was used as a nuclear counterstain.

Negative control 1: PBS only.



Flow Cytometry (Intracellular) - Anti-Syntenin antibody [EPR8102] (ab133267)

Overlay histogram showing SHSY-5Y cells stained with unpurified ab133267 (red line). The cells were fixed with 4% paraformaldehyde (10 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 proteinprotein interactions followed by the antibody (ab133267, 1/10000 dilution) for 30 min at 22°C. The secondary antibody used was Alexa Fluor[®] 488 goat anti-rabbit lgG (H&L) (ab150077) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit lgG (monoclonal) (0.1µg/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. This antibody gave a positive signal in SHSY-5Y cells fixed with 80% methanol (5 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.



Western blot - Anti-Syntenin antibody [EPR8102] (ab133267)

All lanes : Anti-Syntenin antibody [EPR8102] (ab133267) at 1/1000 dilution (unpurified)

Lane 1: Human fetal brain lysate

Lane 2: 293T cell lysate

Lane 3: Human fetal heart lysate

Lane 4: HeLa cell lysate

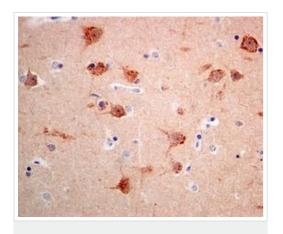
Lysates/proteins at 10 µg per lane.

Secondary

All lanes: Goat anti-rabbit HRP conjugated antibody at 1/2000

dilution

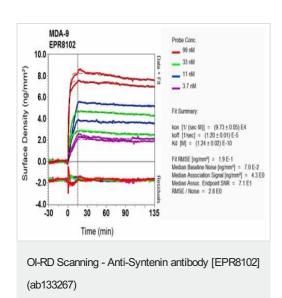
Predicted band size: 32 kDa **Observed band size:** 32 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Syntenin antibody
[EPR8102] (ab133267)

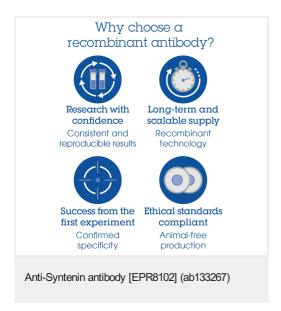
Immunohistochemical analysis of Syntenin in paraffin embedded Human brain tissue, using unpurified ab133267 at a 1/50 dilution.

Perform heat mediated antigen retrieval before commencing with IHC staining protocol.



Equilibrium disassociation constant (K_D) Learn more about K_D

Click here to learn more about K_D



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