abcam

Product datasheet

Anti-Syntenin antibody ab 19903





★★★★ 4 Abreviews 36 References 4 Images

Overview

Product name Anti-Syntenin antibody

Description Rabbit polyclonal to Syntenin

Host species Rabbit

Tested applications Suitable for: ICC/IF, WB

Species reactivity Reacts with: Mouse, Rat, Human

Predicted to work with: Xenopus laevis, Monkey, Zebrafish

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

General notes 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

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 or -

80°C. Avoid freeze / thaw cycle.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising

agent. If you would like information about the formulation of a specific lot, please contact our

scientific support team who will be happy to help.

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee

Our Abpromise guarantee covers the use of ab19903 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 |
|-------------|-------------------|--|
| ICC/IF | ★★★☆☆ (1) | Use a concentration of 1 µg/ml. |
| WB | ★★★★ ☆ (1) | Use a concentration of 1 µg/ml. Detects a band of approximately 32 kDa (predicted molecular weight: 32 kDa). |

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Function

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.

Tissue specificity

 $\label{thm:continuous} Widely\ expressed.\ Expressed\ in\ fetal\ kidney,\ liver,\ lung\ and\ brain.\ In\ adult\ highest\ expression\ in$

heart and placenta.

Sequence similarities

Contains 2 PDZ (DHR) domains.

Post-translational

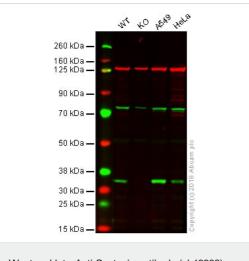
modifications

Phosphorylated on tyrosine residues.

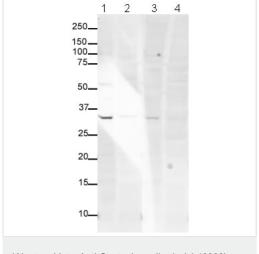
Cellular localization

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 spectrometry in melanosome fractions from stage I to stage IV.

Images



Western blot - Anti-Syntenin antibody (ab19903)



Western blot - Anti-Syntenin antibody (ab19903)

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 - ab19903 observed at 32 kDa. Red - loading control, ab130007, observed at 130 kDa.

ab19903 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. ab19903 and ab130007 (Mouse anti-Vinculin loading control) were incubated overnight at 4°C at 1 µg/ml and 1/20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG 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.

All lanes: Anti-Syntenin antibody (ab19903) at 1 µg/ml

Lane 1: HEK-293 cell lysate

Lane 2: Human heart tissue lysate

Lane 3: HEK-293 cell lysate with Mouse Syntenin peptide

(**ab20431**) at 1 µg/ml

Lane 4: Human heart tissue lysate with Mouse Syntenin peptide

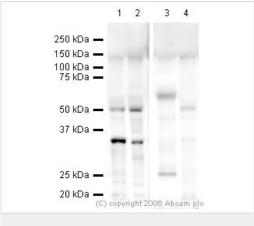
(ab20431) at 1 µg/ml

Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 32 kDa

ab19903 detects a band of the expected size (32kDa) in HEK-293 cell lysate. This band is partially quenched by the addition of ab20431. No band is observed in human heart lysate.



Western blot - Anti-Syntenin antibody (ab19903)

All lanes: Anti-Syntenin antibody (ab19903) at 1 µg/ml

Lane 1: Mouse heart lysate

Lane 2: Rat heart lysate

Lane 3: Mouse heart lysate with Mouse Syntenin peptide

(ab20431) at 1 µg/ml

Lane 4: Rat heart lysate with Mouse Syntenin peptide (ab20431)

at 1 µg/ml

Lysates/proteins at 20 µg per lane.

Secondary

All lanes: Goat Anti-Rabbit lgG H&L (HRP) (ab6721) at 1/50000

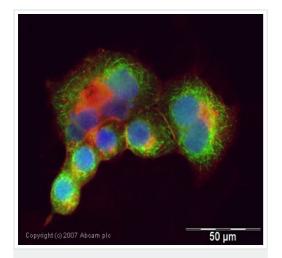
dilution

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

Additional bands at: 50 kDa. We are unsure as to the identity of

these extra bands.

ab19903 detects a band of the expected size (32kDa) in both mouse and rat heart lysate. This band is quenched completely by the addition of the immunizing peptide in the mouse lysate and is partially quenched by the addition of <u>ab20431</u> in the rat heart lysate.



Immunocytochemistry/ Immunofluorescence - Anti-Syntenin antibody (ab19903)

ICC/IF image of ab19903 stained human HEK 293 cells. The cells were PFA fixed (10 min), permabilised in TBS-T (20 min) and incubated with the antibody (ab19903, 1µg/ml) for 1h at room temperature. 1%BSA / 10% normal goat serum / 0.3M glycine was used to quench autofluorescence and block non-specific protein-protein interactions. The secondary antibody (green) was Alexa Fluor® 488 goat anti-rabbit IgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red). DAPI was used to stain the cell nuclei (blue).

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