abcam

Product datasheet

Anti-HIP1 antibody [EPR10814] ab181238



Recombinant RabMAb

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Overview

Product name Anti-HIP1 antibody [EPR10814]

Description Rabbit monoclonal [EPR10814] to HIP1

Host species Rabbit

Suitable for: ICC/IF, WB, IP **Tested applications** Species reactivity Reacts with: Mouse, Human

Predicted to work with: Rat

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

(Peptide available as ab188140)

Positive control WB: HAP1, HCT-116, HeLa, Jurkat and A549 cell lysates. ICC/IF: HeLa 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® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb® patents.

Properties

Form Liquid

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

term. Avoid freeze / thaw cycle.

Storage buffer Preservative: 0.01% Sodium azide

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

Purity Tissue culture supernatant

Clonality Monoclonal Clone number EPR10814

Isotype lgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab181238 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		1/100 - 1/250.
WB		1/1000 - 1/10000. Detects a band of approximately 116 kDa (predicted molecular weight: 116 kDa).
IP		1/20 - 1/40.

Target

Function

Plays a role in clathrin-mediated endocytosis and trafficking. Involved in regulating AMPA receptor trafficking in the central nervous system in an NMDA-dependent manner. Enhances androgen receptor (AR)-mediated transcription. May act as a proapoptotic protein that induces cell death by acting through the intrinsic apoptosis pathway. Binds 3-phosphoinositides (via ENTH domain). May act through the ENTH domain to promote cell survival by stabilizing receptor tyrosine kinases following ligand-induced endocytosis. May play a functional role in the cell filament networks. May be required for differentiation, proliferation, and/or survival of somatic and germline progenitors.

Tissue specificity

Ubiquitously expressed with the highest level in brain. Expression is up-regulated in prostate and colon cancer.

Involvement in disease

Note=A chromosomal aberration involving HIP1 is found in a form of chronic myelomonocytic leukemia (CMML). Translocation t(5;7)(q33;q11.2) with PDGFRB. The chimeric HIP1-PDGFRB transcript results from an in-frame fusion of the two genes. The reciprocal PDGFRB-HIP1 transcript is not expressed.

Sequence similarities

Belongs to the SLA2 family.

Contains 1 ENTH (epsin N-terminal homology) domain.

Contains 1 I/LWEQ domain.

Domain

The pseudo DED region (pDED) mediates the interaction with IFT57.

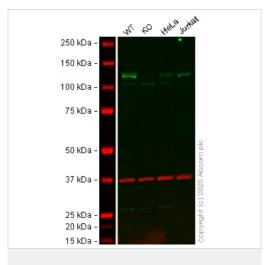
Binds F-actin via the talin-like I/LWEQ domain.

Cellular localization

Cytoplasm. Nucleus. Endomembrane system. Cytoplasmic vesicle > clathrin-coated vesicle membrane. Shuttles between cytoplasm and nucleus. Nuclear translocation can be induced by

AR.

Images



Western blot - Anti-HIP1 antibody [EPR10814] (ab181238)

All lanes : Anti-HIP1 antibody [EPR10814] (ab181238) at 1/1000 dilution

Lane 1: wild-type HAP1 cell lysate

Lane 2: HIP1 knockout HAP1 cell lysate

Lane 3 : HeLa cell lysate

Lane 4 : Jurkat cell lysate

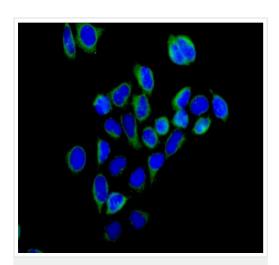
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 116 kDa **Observed band size:** 116 kDa

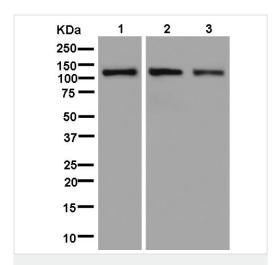
Lanes 1 - 3: Merged signal (red and green). Green - ab181238 observed at 116 kDa. Red - loading control <u>ab8245</u> (Mouse anti-GAPDH antibody [6C5]) observed at 37kDa.

ab181238 was shown to react with HIP1 in wild-type HAP1 cells in western blot with loss of signal observed in HIP1 knockout sample. Wild-type and HIP1 knockout HAP1 cell lysates were subjected to SDS-PAGE. Membranes were blocked in 3% milk in TBS-T (0.1% Tween®) before incubation with ab181238 and ab8245 (Mouse anti-GAPDH antibody [6C5]) overnight at 4°C at a 1 in 1000 Dilution and a 1 in 20000 dilution respectively. Blots were incubated 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 in 20000 dilution for 1 hour at room temperature before imaging.



Immunocytochemistry/ Immunofluorescence - Anti-HIP1 antibody [EPR10814] (ab181238)

Immunofluorescent analysis of HeLa cells (-20**l** Acetone-fixed) labeling HIP1 with ab181238 at 1/250 dilution, followed by Goat anti rabbit lgG (Alexa Fluor® 488) secondary at 1/200 dilution and counter-stained with DAPI (blue).



Western blot - Anti-HIP1 antibody [EPR10814] (ab181238)

All lanes : Anti-HIP1 antibody [EPR10814] (ab181238) at 1/2000 dilution

Lane 1 : HCT-116 cell lysate
Lane 2 : HeLa cell lysate

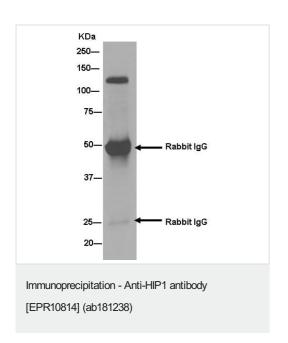
Lane 3 : Jurkat cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 116 kDa



Western blot analysis of immunoprecipitation pellet from A549 cell lysate immunoprecipitated using ab181238 at 1/50 dilution. Secondary: Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated at 1/1000 dilution.



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