abcam

Product datasheet

Anti-Pin1 antibody [EPR18546-317] ab192036



★★★★★ 4 Abreviews 3 References 8 Images

Overview

Product name Anti-Pin1 antibody [EPR18546-317]

Rabbit monoclonal [EPR18546-317] to Pin1 **Description**

Host species Rabbit

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

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Recombinant full length protein. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: HeLa, MCF7, C6 and NIH/3T3 cell lysates; Human fetal brain, fetal heart and fetal kidney

lysates; mouse and rat brain, heart, kidney and spleen lysates. ICC/IF: HeLa and NIH/3T3 cells.

Flow Cyt (intra).: HeLa cells. IP: NIH/3T3 cell lysate.

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

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

Purity Protein A purified

Clonality Monoclonal Clone number EPR18546-317

Isotype IgG

Applications

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab192036 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
WB	★★★★★(4)	1/2000. Detects a band of approximately 18 kDa (predicted molecular weight: 18 kDa).
ICC/IF		1/100.
IP		1/30.
Flow Cyt (Intra)		1/500.

Target	•
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Function Essential PPlase that regulates mitosis presumably by interacting with NIMA and attenuating its

mitosis-promoting activity. Displays a preference for an acidic residue N-terminal to the

isomerized proline bond. Catalyzing pSer/Thr-Pro cis/trans isomerizations.

Sequence similarities Contains 1 PpiC domain.

Contains 1 WW domain.

Domain The WW domain is required for the interaction with STIL and KIF20B.

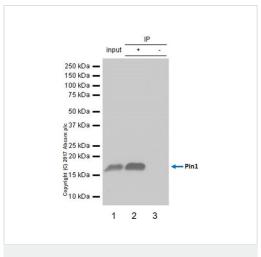
Post-translational

modifications

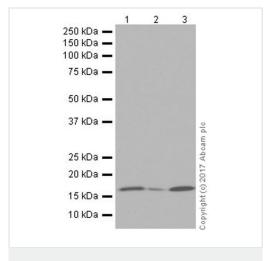
Phosphorylated upon DNA damage, probably by ATM or ATR.

Cellular localization Nucleus.

Images



Immunoprecipitation - Anti-Pin1 antibody [EPR18546-317] (ab192036)



Western blot - Anti-Pin1 antibody [EPR18546-317] (ab192036)

Pin1 was immunoprecipitated from 0.35 mg of NIH/3T3 (mouse embryonic fibroblast cell line) lysate with ab192036 at 1/30 dilution. Western blot was performed from the immunoprecipitate using ab192036 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (ab131366), was used for detection at 1/10000 dilution.

Lane 1: NIH/3T3 whole cell lysate 10 µg (Input).

Lane 2: ab192036 IP in NIH/3T3 whole cell lysate.

Lane 3: Rabbit monoclonal $\lg G$ ($\underline{ab172730}$) instead of ab192036 in NIH/3T3 whole cell lysate.

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

Exposure time: 2 seconds.

All lanes : Anti-Pin1 antibody [EPR18546-317] (ab192036) at 1/2000 dilution

Lane 1: Human fetal brain lysate

Lane 2: Human fetal heart lysate

Lane 3: Human fetal kidney lysate

Lysates/proteins at 10 µg per lane.

Secondary

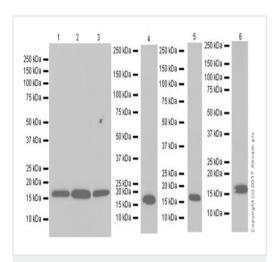
All lanes : VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) at 1/4000 dilution

Developed using the ECL technique.

Predicted band size: 18 kDa
Observed band size: 18 kDa

Exposure time: 5 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot - Anti-Pin1 antibody [EPR18546-317] (ab192036)

All lanes : Anti-Pin1 antibody [EPR18546-317] (ab192036) at 1/5000 dilution

Lane 1 : HeLa (human epithelial cell line from cervix adenocarcinoma) cell lysate

Lane 2: MCF7 (human breast adenocarcinoma cell line) cell lysate

Lane 3: C6 (rat glial tumor cell line) cell lysate

Lane 4: NIH/3T3 (mouse embryonic fibroblast cell line) cell lysate

Lane 5 : Mouse brain lysate

Lane 6 : Rat brain lysate

Lysates/proteins at 20 µg per lane.

Secondary

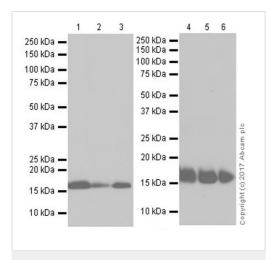
All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/100000 dilution

Developed using the ECL technique.

Predicted band size: 18 kDa **Observed band size:** 18 kDa

Blocking/Dilution buffer: 5% NFDM/TBST.

Exposure times: Lanes 1-3: 15 seconds; Lanes 4-6: 30 seconds.



Western blot - Anti-Pin1 antibody [EPR18546-317] (ab192036)

All lanes : Anti-Pin1 antibody [EPR18546-317] (ab192036) at 1/2000 dilution

Lane 1: Mouse heart lysate

Lane 2: Mouse kidney lysate

Lane 3: Mouse spleen lysate

Lane 4: Rat heart lysate

Lane 5: Rat kidney lysate

Lane 6: Rat spleen lysate

Lysates/proteins at 10 µg per lane.

Secondary

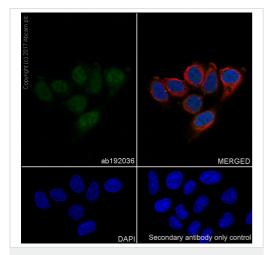
All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/100000 dilution

Developed using the ECL technique.

Predicted band size: 18 kDa **Observed band size:** 18 kDa

Blocking/Dilution buffer: 5% NFDM/TBST.

Exposure times: Lanes 1-3: 5 seconds; Lanes 4-6: 15 seconds.

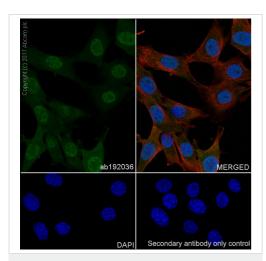


Immunocytochemistry/ Immunofluorescence - Anti-Pin1 antibody [EPR18546-317] (ab192036)

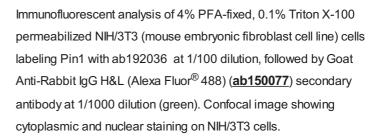
Immunofluorescent analysis of 4% PFA-fixed, 0.1% Triton X-100 permeabilized HeLa (human epithelial cell line from cervix adenocarcinoma) cells labeling Pin1 with ab192036 at 1/100 dilution, followed by Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) (ab150077) secondary antibody at 1/1000 dilution (green). Confocal image showing cytoplasmic and nuclear staining on HeLa cells.

The nuclear counter stain is DAPI (blue). Tubulin is detected with Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor[®] 594) (ab195889) (red) at 1/200 dilution.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) (ab150077) secondary antibody at 1/1000 dilution.

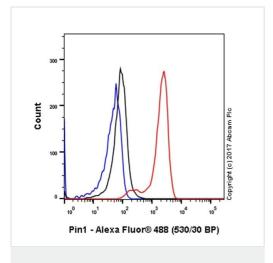


Immunocytochemistry/ Immunofluorescence - Anti-Pin1 antibody [EPR18546-317] (ab192036)



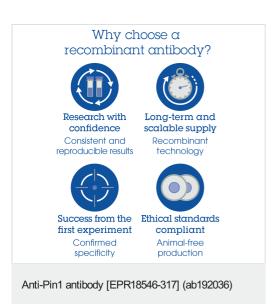
The nuclear counter stain is DAPI (blue). Tubulin is detected with Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor® 594) (ab195889) (red) at 1/200 dilution.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution.



Flow Cytometry (Intracellular) - Anti-Pin1 antibody [EPR18546-317] (ab192036)

Intracellular flow cytometric analysis of 4% PFA-fixed, 90% methanol permeabilized HeLa (human epithelial cell line from cervix adenocarcinoma) cell line labeling Pin1with ab192036 at 1/500 dilution (red) compared with a Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730) (black) and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (blue). Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) at 1/2000 dilution was used as the secondary antibody.



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