abcam

Product datasheet

Anti-Pirh2 antibody [EPR18553] ab189907



ייבער RabMAb

2 References 画像数9

製品の概要

製品名 Anti-Pirh2 antibody [EPR18553]

製品の詳細 Rabbit monoclonal [EPR18553] to Pirh2

由来種 Rabbit

アプリケーション 適用あり: ICC/IF, IP, WB 種交差性 交差種: Mouse, Human

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

ポジティブ・コントロール WB: HeLa, HepG2, Daudi, HEK-293, HCT 116, LNCaP, RAW 264.7 and NIH/3T3 whole cell

lysates; Human fetal heart, fetal kidney and fetal spleen lysates; Mouse kidney and spleen lysates.

ICC/IF: HeLa and HEK-293 cells. IP: HeLa whole cell lysate.

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 long

term. Avoid freeze / thaw cycle.

バッファー pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 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 EPR18553

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ab189907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		1/500.
IP		1/50.
WB		1/1000. Detects a band of approximately 30 kDa (predicted molecular weight: 30 kDa).

ターゲット情報

機能 Mediates E3-dependent ubiquitination and proteasomal degradation of target proteins, including

p53/TP53, HDAC1 and CDKN1B. Preferentially acts on tetrameric p53/TP53. Contributes to the regulation of CDKN1B and p53/TP53 levels, and thereby contributes to the regulation of the cell

cycle progression. Increases AR transcription factor activity.

パスウェイ Protein modification; protein ubiquitination.

配列類似性 Contains 1 CHY-type zinc finger.

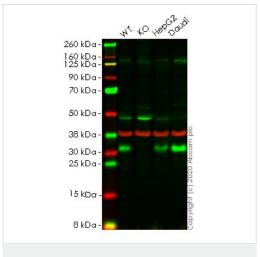
Contains 1 CTCHY-type zinc finger.
Contains 1 RING-type zinc finger.

翻訳後修飾 Subject to ubiquitination and proteasomal degradation. Interaction with PLAGL2 or KAT5

enhances protein stability.

細胞内局在 Nucleus. Nucleus speckle. Cytoplasm.

画像



Western blot - Anti-Pirh2 antibody [EPR18553] (ab189907)

All lanes : Anti-Pirh2 antibody [EPR18553] (ab189907) at 1/1000 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: RCHY1 knockout HeLa cell lysate

Lane 3 : HepG2 cell lysate

Lane 4 : Daudi cell lysate

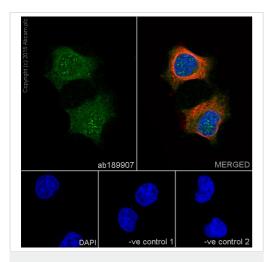
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 30 kDa

Lanes 1-4: Merged signal (red and green). Green - ab189907 observed at 30 kDa. Red - loading control **ab8245** observed at 37 kDa.

ab189907 Anti-Pirh2 antibody [EPR18553] was shown to specifically react with RCHY1 in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab265478 (knockout cell lysate ab258171) was used. Wild-type and RCHY1 knockout samples were subjected to SDS-PAGE. ab189907 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Immunocytochemistry/ Immunofluorescence - Anti-Pirh2 antibody [EPR18553] (ab189907)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cells from cervix adenocarcinoma) cells labeling Pirh2 with ab189907 at 1/500 dilution, followed by Goat anti-rabbit lgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green).

Confocal image showing nuclear and cytoplasmic staining on HeLa cell line.

The nuclear counterstain is DAPI (blue).

Tubulin is detected with <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution and <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution (red).

The negative controls are as follows:-

-ve control 1: ab189907 at 1/500 dilution followed by <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution.

-ve control 2: <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> (Alexa Fluor®488 Goat Anti-Rabbit lgG H&L) at 1/1000 dilution.

ab189907 MERGED

DAPI

-ve control 1

Immunocytochemistry/ Immunofluorescence - Anti-Pirh2 antibody [EPR18553] (ab189907)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HEK-293 (Human epithelial cells from embryonic kidney) cells labeling Pirh2 with ab189907 at 1/500 dilution, followed by Goat anti-rabbit lgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green).

Confocal image showing nuclear and cytoplasmic staining on HEK-293 cell line.

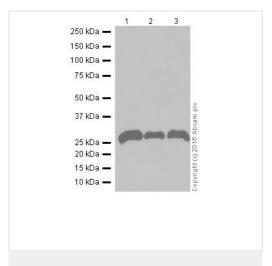
The nuclear counterstain is DAPI (blue).

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-ve control 2: <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> (Alexa Fluor®488 Goat Anti-Rabbit lgG H&L) at 1/1000 dilution.



Western blot - Anti-Pirh2 antibody [EPR18553] (ab189907)

All lanes : Anti-Pirh2 antibody [EPR18553] (ab189907) at 1/1000 dilution

Lane 1: HEK-293 (Human epithelial cells from embryonic kidney) whole cell lysate

Lane 2 : HCT 116 (Human colorectal carcinoma cell line) whole cell lysate

Lane 3: LNCaP (Human prostate cancer cell line) whole cell 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

Predicted band size: 30 kDa **Observed band size:** 30 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

1 2 3
250 kDa —
150 kDa —
100 kDa —
75 kDa —
37 kDa —
25 kDa —
25 kDa —
20 kDa —
15 kDa —
15 kDa —
10 kDa —

Western blot - Anti-Pirh2 antibody [EPR18553] (ab189907)

All lanes : Anti-Pirh2 antibody [EPR18553] (ab189907) at 1/1000 dilution

Lanes 1 & 3: Human fetal heart lysate

Lane 2: Human fetal kidney lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/10000 dilution

Predicted band size: 30 kDa
Observed band size: 30 kDa

Exposure time: 30 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

All lanes : Anti-Pirh2 antibody [EPR18553] (ab189907) at 1/1000 dilution

Lane 1 : Mouse kidney lysate
Lane 2 : Mouse spleen lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit $\lg G \ H\&L \ (HRP) \ (\underline{ab97051})$ at

1/100000 dilution

Predicted band size: 30 kDa **Observed band size:** 30 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

All lanes : Anti-Pirh2 antibody [EPR18553] (ab189907) at 1/1000 dilution

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Lane 1: RAW 264.7 (Mouse macrophage cells transformed with

Abelson murine leukemia virus) whole cell lysate

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

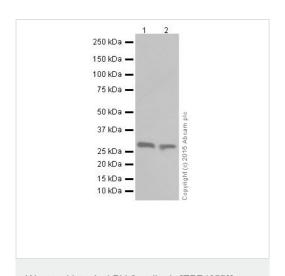
Lysates/proteins at 10 µg per lane.

Secondary

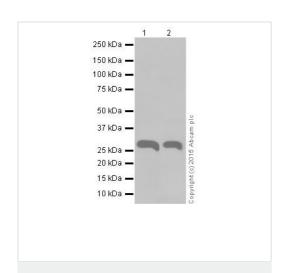
All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at

1/100000 dilution

Predicted band size: 30 kDa **Observed band size:** 30 kDa



Western blot - Anti-Pirh2 antibody [EPR18553] (ab189907)



Western blot - Anti-Pirh2 antibody [EPR18553] (ab189907)

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

Pirh2 was immunoprecipitated from 1mg of HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate with ab189907 at 1/50 dilution.

Western blot was performed from the immunoprecipitate using ab189907 at 1/1000 dilution.

VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>), was used for detection at 1/10000 dilution.

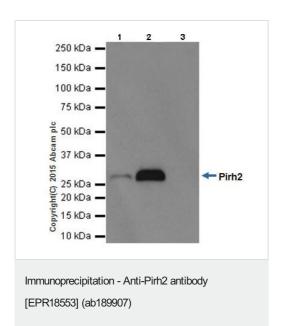
Lane 1: HeLa whole cell lysate 10ug (Input).

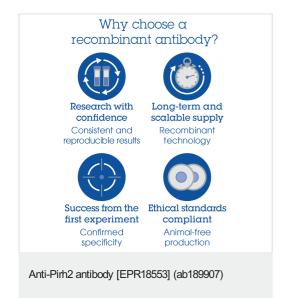
Lane 2: ab189907 IP in HeLa whole cell lysate.

Lane 3: Rabbit monoclonal $\lg G$ ($\underline{ab172730}$) instead of ab189907 in HeLa whole cell lysate.

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

Exposure time: 30 seconds.





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