abcam

Product datasheet

Anti-FHL2 antibody [EPR17860-20] ab202584





重组 RabMAb

4 References 11 图像

概述

产品名称 Anti-FHL2抗体[EPR17860-20]

描述 兔单克隆抗体[EPR17860-20] to FHL2

宿主 Rabbit

适用于: WB, ICC/IF, IP 经测试应用

种属反应性 与反应: Mouse, Rat, Human

免疫原 Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

阳性对照 WB: U-2 OS, K526, HeLa, SW480, PC-12 and HT1080 whole cell lysates; Human fetal heart

lysate; Mouse and rat heart lysates. ICC/IF: A-673 and NIH/3T3 cells. IP: SW480 whole cell lysate.

ICC/IF: U-2 OS cells.

常规说明 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**.

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 long

term. Avoid freeze / thaw cycle.

存储溶液 pH: 7.2

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 EPR17860-20

应用

The Abpromise guarantee Abpromise™承诺保证使用ab202584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
WB		1/1000. Detects a band of approximately 32 kDa (predicted molecular weight: 32 kDa).
ICC/IF		1/400 - 1/200. ab202584 works both with PFA and methanol fixation. Fixation with PFA gives the strongest signal.
IP		1/30.

靶标

功能 May function as a molecular transmitter linking various signaling pathways to transcriptional

regulation. Negatively regulates the transcriptional repressor E4F1 and may function in cell

growth.

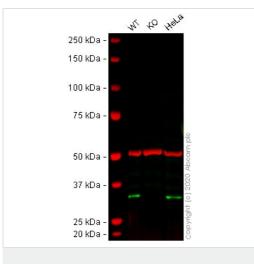
组织**特异性** Expressed in skeletal muscle and heart.

序列相似性 Contains 4 LIM zinc-binding domains.

结**构域** The third LIM zinc-binding mediates interaction with E4F1.

细胞定位 Cytoplasm. Nucleus.

图片



Western blot - Anti-FHL2 antibody [EPR17860-20] (ab202584)

All lanes : Anti-FHL2 antibody [EPR17860-20] (ab202584) at 1/1000 dilution

Lane 1 : Wild-type U-2 OS cell lysate

Lane 2: FHL2 knockout U-2 OS cell lysate

Lane 3: HeLa (Human epithelial cell line from cervix

adenocarcinoma) whole cell lysate

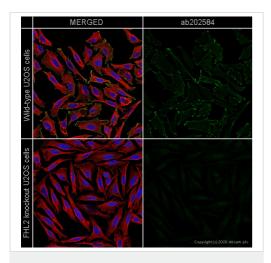
Lysates/proteins at 40 µg per lane.

Performed under reducing conditions.

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

Lanes 1 - 3: Merged signal (red and green). Green - ab202584 observed at 32 kDa. Red - loading control <u>ab7291</u> (Mouse anti-Alpha Tubulin [DM1A] observed at 55kDa.

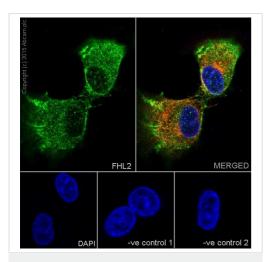
ab202584 was shown to react with FHL2 in wild-type U-2 OS cells in western blot with loss of signal observed in FHL2 knockout sample.Wild-type and FHL2 knockout U-2 OS cell lysates were subjected to SDS-PAGE. Membranes were blocked in 3% milk in TBS-T (0.1% Tween[®]) before incubation with ab202584 and ab7291 (Mouse anti-Alpha Tubulin [DM1A] overnight at 4°C at a 1 in 1000 dilution and a 1 in 20000 dilution respectively. Blots were incubated with Goat anti-Rabbit lgG H&L (IRDye[®] 800CW) preabsorbed (ab216773) and Goat anti-Mouse lgG H&L (IRDye[®] 680RD) preabsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Immunocytochemistry/ Immunofluorescence - Anti-FHL2 antibody [EPR17860-20] (ab202584)

ab202584 staining FHL2 in wild-type U2OS cells (top panel) and FHL2 knockout U2OS cells (bottom panel). The cells were fixed with 4% PFA (10 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated with ab202584 at 1/200 dilution and ab7291 (Tubulin) at 1/1000 dilution overnight at +4°C, followed by a further incubation at room temperature for 1h with a goat secondary antibody to rabbit lgG (Alexa Fluor® 488) (ab150081) at 2 µg/ml (shown in green) and a goat secondary antibody to mouse lgG (Alexa Fluor® 594) (ab150120) at 2 µg/ml (shown in pseudo color red). Nuclear DNA was labelled in blue with DAPI.

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).



Immunocytochemistry/ Immunofluorescence - Anti-FHL2 antibody [EPR17860-20] (ab202584)

260 kDa160 kDa125 kDa70 kDa38 kDa30 kDa25 kDa15 kDa-

Western blot - Anti-FHL2 antibody [EPR17860-20] (ab202584)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized A-673 (Human muscle Ewing's Sarcoma cell line) cells labeling FHL2 with ab202584 at 1/400 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/500 dilution (green). Confocal image showing cytoplasmic and weakly nuclear staining on A-673 cell line. The nuclear counter stain is DAPI (blue). Tubulin is detected with ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution and ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution (red).

The negative controls are as follows:

-ve control 1: ab202584 at 1/400 dilution followed by <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 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 1/500 dilution.

All lanes : Anti-FHL2 antibody [EPR17860-20] (ab202584) at 1/1000 dilution

Lane 1: Wild-type HeLa lysate

Lane 2: FHL2 knockout HeLa lysate

Lane 3: K562 lysate

Lysates/proteins at 20 µg per lane.

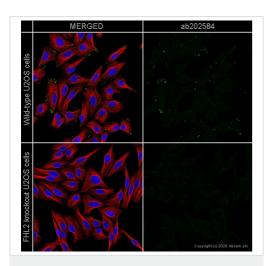
Performed under reducing conditions.

Predicted band size: 32 kDa

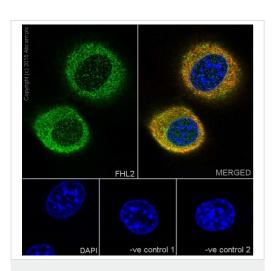
Lanes 1-3: Merged signal (red and green). Green - ab202584 observed at 32 kDa. Red - loading control <u>ab7291</u> observed at 50 kDa.

ab202584 Recombinant Anti-FHL2 antibody [EPR17860-20] was shown to specifically react with FHL2 in wild-type HeLa cells. Loss of signal was observed when knockout cell line <u>ab265475</u> (knockout cell lysate <u>ab257441</u>) was used. Wild-type and FHL2 knockout samples were subjected to SDS-PAGE. ab202584 and Anti-alpha Tubulin antibody [DM1A] - Loading Control?(<u>ab7291</u>) 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 (<u>ab216773</u>) and

Goat anti-Mouse IgG H&L (IRDye[®] 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Immunocytochemistry/ Immunofluorescence - Anti-FHL2 antibody [EPR17860-20] (ab202584)



Immunocytochemistry/ Immunofluorescence - Anti-FHL2 antibody [EPR17860-20] (ab202584)

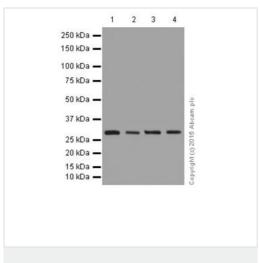
ab202584 staining FHL2 in wild-type U2OS cells (top panel) and FHL2 knockout U2OS cells (bottom panel). The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated with ab202584 at 1/200 dilution and ab7291 (Tubulin) at 1/1000 dilution overnight at +4°C, followed by a further incubation at room temperature for 1h with a goat secondary antibody to rabbit lgG (Alexa Fluor® 488) (ab150081) at 2 μ g/ml (shown in green) and a goat secondary antibody to mouse lgG (Alexa Fluor® 594) (ab150120) at 2 μ g/ml (shown in pseudo color red). Nuclear DNA was labelled in blue with DAPI.

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized NIH/3T3 (Mouse embyro fibroblast cells) cells labeling FHL2 with ab202584 at 1/400 dilution, followed by Goat anti-rabbit lgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/500 dilution (green). Confocal image showing cytoplasmic and weakly nuclear staining on NIH/3T3 cell line. The nuclear counter stain is DAPI (blue). Tubulin is detected with ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution and ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution (red).

The negative controls are as follows:

-ve control 1: ab202584 at 1/400 dilution followed by <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 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 1/500 dilution.



Western blot - Anti-FHL2 antibody [EPR17860-20] (ab202584)

All lanes : Anti-FHL2 antibody [EPR17860-20] (ab202584) at 1/1000 dilution

Lane 1 : SW480 (Human colorectal adenocarcinoma cell line) whole cell lysate

Lane 2: PC-12 (Rat adrenal gland pheochromocytoma) whole cell lysate

Lane 3: HT1080 (Human fibrosarcoma cells) whole cell lysate

Lane 4: HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate

Lysates/proteins at 10 µg per lane.

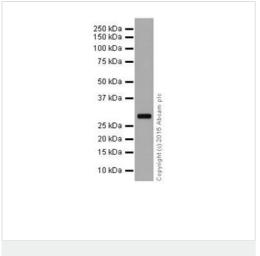
Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/1000 dilution (Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated)

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

Exposure time: 30 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.



Anti-FHL2 antibody [EPR17860-20] (ab202584) at 1/10000 dilution + Human fetal heart lysate at 10 μg

Secondary

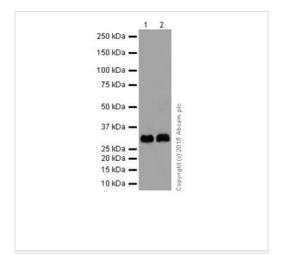
Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/1000 dilution

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

Exposure time: 15 seconds

Western blot - Anti-FHL2 antibody [EPR17860-20] (ab202584)

Blocking/Dilution buffer: 5% NFDM/TBST.



All lanes : Anti-FHL2 antibody [EPR17860-20] (ab202584) at 1/2000 dilution

Lane 1 : Mouse heart lysate

Lane 2 : Rat heart lysate

Lysates/proteins at 10 µg per lane.

Secondary

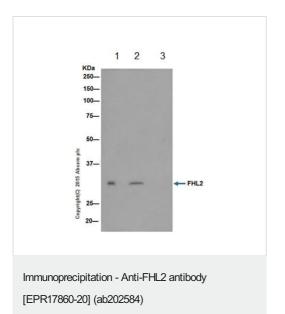
All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/1000 dilution (Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated)

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

Exposure time: 10 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

Western blot - Anti-FHL2 antibody [EPR17860-20] (ab202584)

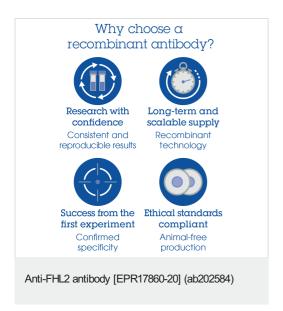


FHL2 was immunoprecipitated from 1mg of SW480 (Human colorectal adenocarcinoma cell line) whole cell lysate with ab202584 at 1/30 dilution. Western blot was performed from the immunoprecipitate using ab202584 at 1/1000 dilution. Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG, was used as secondary antibody at 1/1500 dilution.

Lane 1: SW480 whole cell lysate 10 μg (Input). Lane 2: ab202584 IP in SW480 whole cell lysate. Lane 3: Rabbit monoclonal IgG (ab172730) instead of ab202584 in SW480 whole cell lysate.

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

Exposure time: 10 seconds.



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