

Anti-FANCA/FAA antibody [EPR16519] ab201457

敲除验证
重组
RabMAb

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概述

产品名称	Anti-FANCA/FAA抗体[EPR16519]
描述	兔单克隆抗体[EPR16519] to FANCA/FAA
宿主	Rabbit
经测试应用	适用于: IP, WB
种属反应性	与反应: Human
免疫原	Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.
阳性对照	WB: HeLa, HEK-293, A431, A549, HAP1 and Jurkat whole cell lysates; Human colon lysate. IP: HeLa whole cell lysate.
常规说明	<p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> - High batch-to-batch consistency and reproducibility - Improved sensitivity and specificity - Long-term security of supply - Animal-free production <p>For more information see here.</p> <p>Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb[®] patents.</p>

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	<p>pH: 7.2</p> <p>Preservative: 0.01% Sodium azide</p> <p>Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA</p>
纯度	Protein A purified
克隆	单克隆
克隆编号	EPR16519
同种型	IgG

应用

The Abpromise guarantee **Abpromise™** 承诺保证使用ab201457于以下的经测试应用

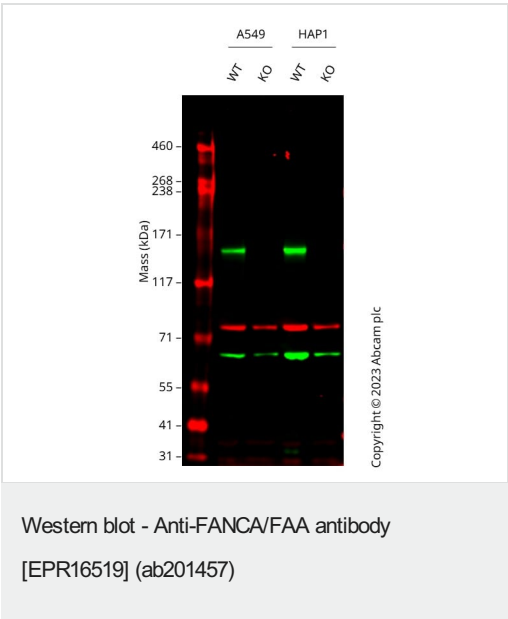
“应用说明”部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
IP		1/30.
WB		1/1000. Detects a band of approximately 163 kDa (predicted molecular weight: 163 kDa).

靶标

功能	DNA repair protein that may operate in a postreplication repair or a cell cycle checkpoint function. May be involved in interstrand DNA cross-link repair and in the maintenance of normal chromosome stability.
疾病相关	Defects in FANCA are a cause of Fanconi anemia (FA) [MIM:227650]. FA is a genetically heterogeneous, autosomal recessive disorder characterized by progressive pancytopenia, a diverse assortment of congenital malformations, and a predisposition to the development of malignancies. At the cellular level it is associated with hypersensitivity to DNA-damaging agents, chromosomal instability (increased chromosome breakage), and defective DNA repair.
翻译后修饰	Phosphorylated upon DNA damage, probably by ATM or ATR. Phosphorylation is required for the formation of the nuclear complex. Not phosphorylated in cells derived from groups A, B, C, E, F, G, and H.
细胞定位	Nucleus. Cytoplasm. The major form is nuclear. The minor form is cytoplasmic.

图片



All lanes : Anti-FANCA/FAA antibody [EPR16519] (ab201457) at 1/1000 dilution

- Lane 1 :** Wild-type A549 cell lysate
- Lane 2 :** FANCA knockout A549 cell lysate
- Lane 3 :** Wild-type HAP1 cell lysate
- Lane 4 :** FANCA knockout HAP1 cell lysate

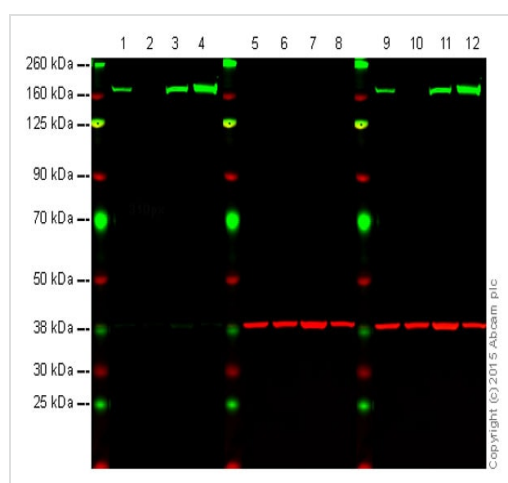
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 163 kDa

Observed band size: 163 kDa

Western blot: Anti-FANCA antibody [EPR16519] (ab201457) staining at 1/1000 dilution, shown in green; Mouse anti-CANX [CANX/1543] ([ab238078](#)) loading control staining at 1/20000 dilution, shown in red. In Western blot, ab201457 was shown to bind specifically to FANCA. A band was observed at 163 kDa in wild-type A549 cell lysates with no signal observed at this size in FANCA knockout cell line. To generate this image, wild-type and FANCA knockout A549 cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3% milk in TBS-0.1% Tween® 20 (TBS-T) before incubation with primary antibodies overnight at 4°C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L 800CW and Goat anti-Mouse IgG H&L 680RD at 1/20000 dilution.



Western blot - Anti-FANCA/FAA antibody
[EPR16519] (ab201457)

Lanes 1, 5 and 9: Wild-type HAP1 cell lysate (20 µg)

Lanes 2, 6 and 10: FANCA/FAA beta knockout HAP1 cell lysate (20 µg)

Lanes 3, 7 and 11: HeLa cell lysate (20 µg)

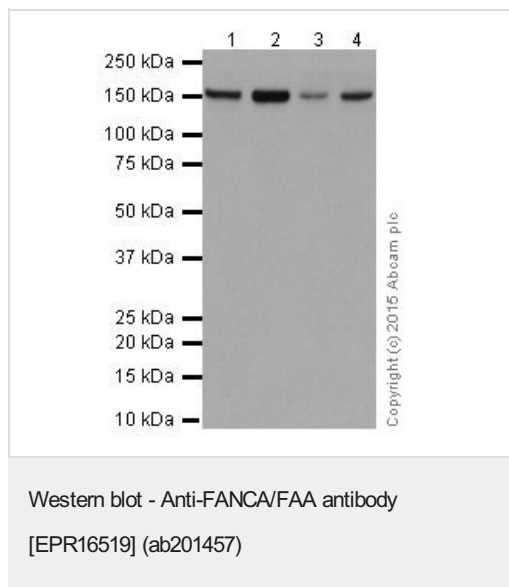
Lanes 4, 8 and 12: HEK293 cell lysate (20 µg)

Lanes 1, 2, 3 and 4: Green signal from target – ab201457 observed at 163 kDa

Lanes 5, 6, 7 and 8: Red signal from loading control – [ab8245](#) observed at 37 kDa

Lanes 9, 10, 11 and 12: Merged (red and green) signal

ab201457 was shown to specifically react with FANCA/FAA beta when FANCA/FAA beta knockout samples were used. Wild-type and FANCA/FAA beta knockout samples were subjected to SDS-PAGE. ab201457 and [ab8245](#) (loading control to GAPDH) were diluted 1/10000 and 1/2000 respectively and incubated overnight at 4°C. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed [ab216773](#) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed [ab216776](#) secondary antibodies at 1/10000 dilution for 1 h at room temperature before imaging.



All lanes : Anti-FANCA/FAA antibody [EPR16519] (ab201457) at 1/10000 dilution

Lane 1 : HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate

Lane 2 : HEK-293 (Human epithelial cells from embryonic kidney) whole cell lysate

Lane 3 : Jurkat (Human T cell leukemia cells from peripheral blood) whole cell lysate

Lane 4 : A431 (Human epidermoid carcinoma) whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

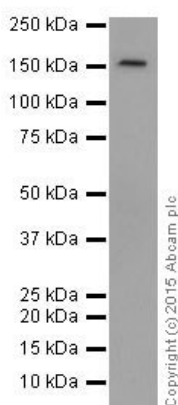
All lanes : Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 163 kDa

Observed band size: 163 kDa

Exposure time: 1 minute

Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot - Anti-FANCA/FAA antibody
[EPR16519] (ab201457)

Anti-FANCA/FAA antibody [EPR16519] (ab201457) at 1/1000 dilution + Human colon lysate at 10 µg

Secondary

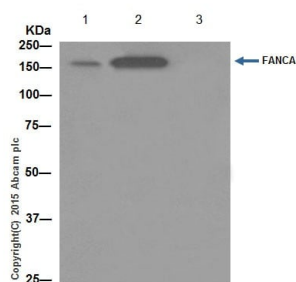
Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/1000 dilution

Predicted band size: 163 kDa

Observed band size: 163 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.



Immunoprecipitation - Anti-FANCA/FAA antibody
[EPR16519] (ab201457)

FANCA/FAA was immunoprecipitated from 1mg of HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate with ab201457 at 1/30 dilution. Western blot was performed from the immunoprecipitate using ab201457 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) ([ab131366](#)), was used for detection at 1/1500 dilution.

Lane 1: HeLa whole cell lysate 10 µg (Input). Lane 2: ab201457 IP in HeLa whole cell lysate. Lane 3: Rabbit monoclonal IgG ([ab172730](#)) instead of ab201457 in HeLa whole cell lysate.

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

Exposure time: 30 seconds.

Why choose a recombinant antibody?



Research with confidence
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Recombinant technology



Success from the first experiment
Confirmed specificity



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Animal-free production

Anti-FANCA/FAA antibody [EPR16519] (ab201457)

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