abcam

Product datasheet

Anti-RAP1A antibody [1D2-1C64] ab175329

5 References 5 图像

概述

产品名称	Anti-RAP1A抗体[1D2-1C64]
描述	小鼠单克隆抗体[1D2-1C64] to RAP1A
宿主	Mouse
经测试应用	**适用于:** IHC-P, WB, IP, ICC/IF
种属反应性	**与反应:** Mouse, Human **预测可用于:** Rat, Cow
免疫原	Recombinant full length protein corresponding to Human RAP1A aa 1 to the C-terminus. Database link: **P62834** Run BLAST with Run BLAST with
阳性对照	HeLa, Hek293T, U2OS and mouse NIH 3T3 cell lysates, HeLa cells, C2C12 cells.
常规说明	The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing. If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&As

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	Preservative: 0.05% Sodium azide Constituents: 30% Glycerol (glycerin, glycerine), 0.1% BSA, 69% PBS
纯度	Protein A purified
克隆	单克隆
克隆编号	1D2-1C64
同种型	IgG2a

应用

The Abpromise guarantee **Abpromise™承诺保证使用ab175329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
IHC-P		1/100 - 1/1000.
WB		1/500 - 1/1000. Predicted molecular weight: 21 kDa.
IP		Use at 2 µg/mg of lysate.
ICC/IF		1/50 - 1/200.

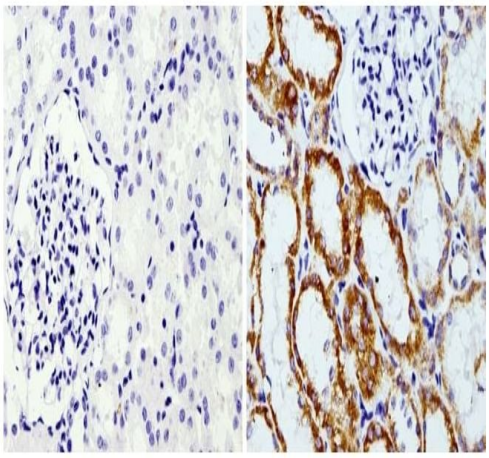
靶标

功能 Induces morphological reversion of a cell line transformed by a Ras oncogene. Counteracts the mitogenic function of Ras, at least partly because it can interact with Ras GAPs and RAF in a competitive manner.

序列相似性 Belongs to the small GTPase superfamily. Ras family.

细胞定位 Cell membrane.

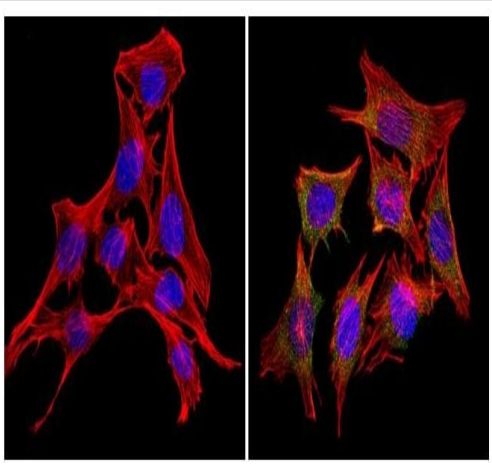
图片



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-RAP1A antibody [1D2-1C64] (ab175329)

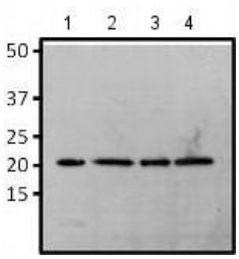
Immunohistochemistry analysis of RAP1A using ab175329 at 1/200 dilution showing staining in the membrane of paraffin-embedded human kidney tissue (right) compared with a negative control without primary antibody (left). Detection was performed using an HRP-conjugated secondary antibody followed by colorimetric detection using a DAB kit.

Antigen retrieval was performed using 10mM sodium citrate (pH 6.0), microwaved for 8-15 min.



Immunocytochemistry/ Immunofluorescence - Anti-RAP1A antibody [1D2-1C64] (ab175329)

Immunocytochemical analysis of RAP1A (green) showing staining in the in the cytoplasm of Formalin-fixed and 0.1% Triton X-100 permeabilized C2C12 cells (right) using ab175329 at 1/20 dilution compared to a negative control without primary antibody (left) followed by DyLight-conjugated secondary antibody. F-actin (red) was stained with a fluorescent red phalloidin and nuclei (blue) were stained with DAPI.



Western blot - Anti-RAP1A antibody [1D2-1C64] (ab175329)

All lanes : Anti-RAP1A antibody [1D2-1C64] (ab175329) at 1/500 dilution

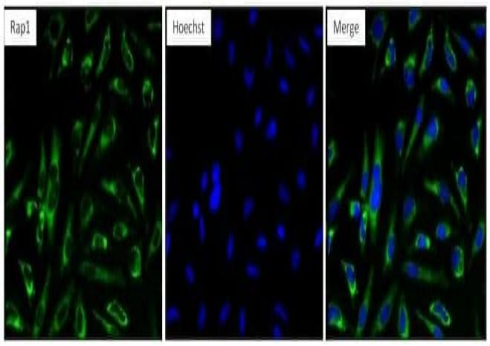
Lane 1 : HeLa cell lysate
Lane 2 : HEK293T cell lysate
Lane 3 : U2OS cell lysate
Lane 4 : Mouse NIH 3T3 cell lysate

Lysates/proteins at 25 µg per lane.

Secondary
All lanes : Goat anti-mouse IgG-HRP at 1/15000 dilution

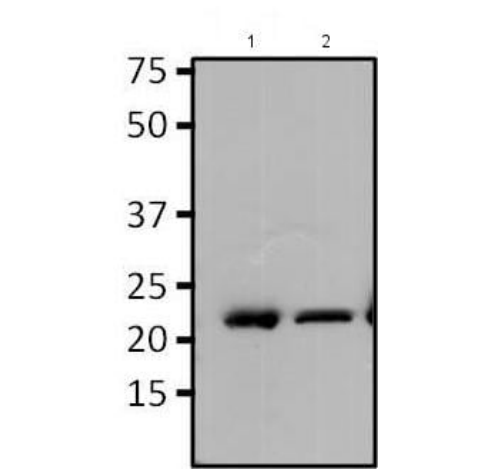
Developed using the ECL technique.

Predicted band size: 21 kDa



Immunocytochemistry/ Immunofluorescence - Anti-RAP1A antibody [1D2-1C64] (ab175329)

Immunofluorescence analysis of formalin-fixed permeabilized HeLa cells, labeling RAP1A (green, left panel) using ab175329 at a 1/100 dilution followed by DyLight 488-conjugated goat anti-mouse IgG secondary antibody at a 1/400 dilution. Nuclei (blue) were stained with Hoechst 33342 dye (central panel).



Immunoprecipitation - Anti-RAP1A antibody [1D2-1C64] (ab175329)

Western blot analysis on immunoprecipitation pellet from mouse NIH 3T3 cells. The antigen-antibody complex was formed by incubating 750 µg of NIH 3T3 whole cell lysate with 2 µg of ab175329 overnight at 4°C. The immune-complex was then captured on 50 µl Protein A/G Plus Agarose, washes extensively and eluted in sample buffer. 1) 25 µg of NIH 3T3 whole cell lysate, as a control, and 2) eluted sample were resolved on a SDS PAGE gel. The membrane was probed with ab175329 at a 1/500 dilution. Chemiluminescent detection was perfomed.

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