abcam

Product datasheet

Anti-Rab5 antibody [EPR21801] - Early Endosome Marker ab218624





RabMAb

★★★★★ 1 Abreviews 14 References 13 图像

概述

产品名称 Anti-Rab5抗体[EPR21801] - Early Endosome Marker

宿主 Rabbit

特异性 This antibody could cross-react with RAB5C, but affinity is very low.

经测试应用 适用于: Flow Cyt (Intra), WB, IHC-P, ICC/IF, IP

种属反应性 与反应: Mouse, Rat, Human

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

阳性对照 WB: HAP1, HeLa, MCF7 and Neuro-2a whole cell lysates; Human fetal brain, fetal heart and fetal

spleen lysates; Rat and mouse brain and heart lysates. IHC-P: Human kidney and cerebrum tissues; Mouse kidney tissue; Rat liver tissue. ICC/IF: NIH/3T3, HeLa and HAP1 cells. Flow Cyt

(intra): MCF7 cells, Hap1 cells. IP: MCF7 whole cell lysate.

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: PBS, 40% Glycerol, 0.05% BSA

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纯**度** Protein A purified

克隆 单克隆

克隆编号 EPR21801

同种型 IgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab218624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
Flow Cyt (Intra)		1/50.
WB		1/1000. Detects a band of approximately 25 kDa (predicted molecular weight: 24 kDa).
IHC-P		1/4000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
ICC/IF		1/1000.
IP		1/30.

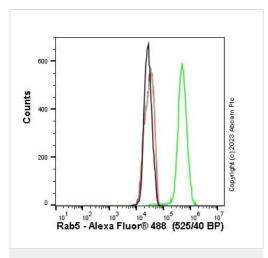
靶标

功能 Required for the fusion of plasma membranes and early endosomes.

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

细胞定位 Cell membrane. Early endosome membrane. Melanosome. Enriched in stage I melanosomes.

图片



Flow Cytometry (Intracellular) - Anti-Rab5 antibody [EPR21801] - Early Endosome Marker (ab218624)

Flow cytometry overlay histogram showing wild-type Hap1 (green line) and Rab5A knockout Hap1 stained with ab218624 (red line). The cells were fixed with 80% methanol (5 min) and then permeabilised with 0.1% PBS-Triton X-100 for 15 min. The cells were then incubated in 1x PBS containing 10% normal goat serum to block non-specific protein-protein interaction followed by the antibody (ab218624) (1x 10^6 in 100μ l at $0.2~\mu$ g/ml (1/10700)) for 30min at 22° C.

The secondary antibody Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed was incubated at 1/4000 for 30min at 22°C Isotype control antibody Recombinant Rabbit IgG, monoclonal [EPR25A] - Isotype Control was used at the same concentration and conditions as the primary antibody (wild-type Hap1 - black line, Rab5A knockout Hap1 - grey line). Unlabelled sample was also used as a control (this line is not shown for the purpose of simplicity).

Acquisition of >5000 events were collected using a 50 mW Blue laser (488nm) and 525/40 bandpass filter.

This antibody gave a positive signal in Hap1 Fixed with 4% formaldehyde (10 min) / permeabilised with 0.1% PBS-Triton X-100 for 15 min under the same conditions.



Western blot - Anti-Rab5 antibody [EPR21801] -Early Endosome Marker (ab218624)

All lanes : Anti-Rab5 antibody [EPR21801] - Early Endosome Marker (ab218624) at 1/1000 dilution

Lane 1: Wild type HAP1 whole cell lysate

Lane 2: Rab5 knockout HAP1 whole cell lysate

Lane 3: HeLa (Human epithelial cell line from cervix adenocarcinoma) whole cell lysate

Lane 4: MCF7 (Human breast adenocarcinoma cell line) whole cell lysate

Lysates/proteins at 20 µg per lane.

Predicted band size: 24 kDa **Observed band size:** 25 kDa

Exposure time: 15 seconds

HAP1 cells as signal was lost in Rab5 knockout cells. Wild-type and Rab5 knockout samples were subjected to SDS-PAGE. Ab218624 and **ab181602** (Human anti-GAPDH loading control) were incubated overnight at 4°C at 1/1000 dilution and 1/200000 dilution respectively. Blots were developed with Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated (**ab97051**) secondary antibody at 1/100,000 dilution for 1 hour at room temperature before imaging. The blot was developed on a BIO-RAD® ChemiDoc™ MP instrumentusing the ECL technique.

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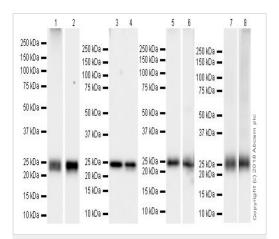
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Immunocytochemistry/ Immunofluorescence - Anti-Rab5 antibody [EPR21801] - Early Endosome Marker (ab218624)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilzed wild-type and RAB5A KO HAP1 cells labeling Rab5 with ab218624 at 1/1000 dilution, followed by Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) (ab150077) secondary antibody at 1/1000 dilution (green). Confocal image showing no staining in RAB5A KO HAP1 cell line and granular cytoplasmic staining in parental HAP1 cell line.

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.



Western blot - Anti-Rab5 antibody [EPR21801] -Early Endosome Marker (ab218624)

All lanes : Anti-Rab5 antibody [EPR21801] - Early Endosome Marker (ab218624) at 1/1000 dilution

Lane 1 : Neuro-2a (mouse neuroblastoma cell line) whole cell lysate

Lane 2: Human fetal brain lysate

Lane 3: Human fetal heart lysate

Lane 4: Human fetal spleen lysate

Lane 5: Rat brain lysate

Lane 6: Rat heart lysate

Lane 7: Mouse brain lysate

Lane 8: Mouse heart lysate

Lysates/proteins at 10 µg per lane.

Secondary

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

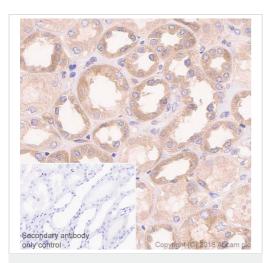
Predicted band size: 24 kDa **Observed band size:** 25 kDa

Exposure time: Lane 1:5 seconds; Lanes 2/8:15 seconds; Lanes 3/4:6 seconds; Lanes 5/7:3 seconds; Lane 6:26 seconds.

Blocking/Dilution buffer: 5% NFDM/TBST.

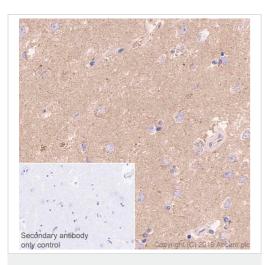
Immunohistochemical analysis of paraffin-embedded human kidney tissue labeling Rab5 with ab218624 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Cytoplasmic staining in human kidney is observed (PMID 7789520). Counterstained with hematoxylin. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) Ready to use.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Rab5 antibody

[EPR21801] - Early Endosome Marker (ab218624)

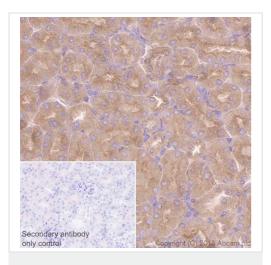


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Rab5 antibody

[EPR21801] - Early Endosome Marker (ab218624)

Immunohistochemical analysis of paraffin-embedded human cerebrum tissue labeling Rab5 with ab218624 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Cytoplasmic staining in human cerebrum is observed (PMID 7789520). Counterstained with hematoxylin. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) Ready to use.

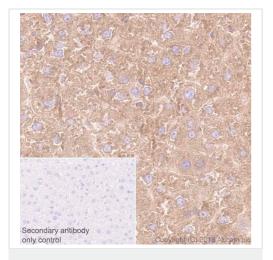


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Rab5 antibody

[EPR21801] - Early Endosome Marker (ab218624)

Immunohistochemical analysis of paraffin-embedded mouse kidney tissue labeling Rab5 with ab218624 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Cytoplasmic staining in mouse kidney is observed (PMID 7789520). Counterstained with hematoxylin. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) Ready to use.

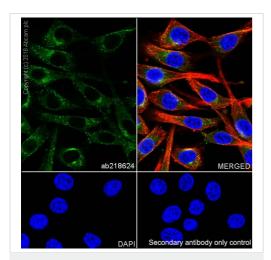


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Rab5 antibody

[EPR21801] - Early Endosome Marker (ab218624)

Immunohistochemical analysis of paraffin-embedded rat liver tissue labeling Rab5 with ab218624 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Cytoplasmic staining in rat liver is observed (PMID 7789520). Counterstained with hematoxylin. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) Ready to use.

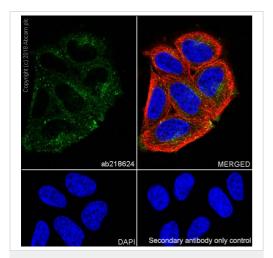


Immunocytochemistry/ Immunofluorescence - Anti-Rab5 antibody [EPR21801] - Early Endosome Marker (ab218624)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilzed NIH/3T3 (mouse embryo fibroblast cell line) cells labeling Rab5 with ab218624 at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (Alexa Fluor[®] 488) (**ab150077**) secondary antibody at 1/1000 dilution (green). Confocal image showing granular cytoplasmic staining in NIH/3T3 cell line.

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.

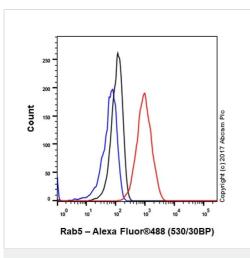


Immunocytochemistry/ Immunofluorescence - Anti-Rab5 antibody [EPR21801] - Early Endosome Marker (ab218624)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilzed HeLa (human epithelial cell line from cervix adenocarcinoma) cells labeling Rab5 with ab218624 at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (Alexa Fluor[®] 488) (ab150077) secondary antibody at 1/1000 dilution (green). Confocal image showing granular cytoplasmic staining in HeLa cell line.

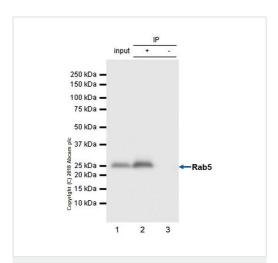
The nuclear counter stain is DAPI (blue). Tubulin is detected with Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor[®] 594) (ab195889) (red) at 1/200 dilution.

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-Rab5 antibody [EPR21801] - Early Endosome Marker (ab218624)

Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed, 90% methanol permeabilized MCF7 (human breast adenocarcinoma cell line) cell line labelling Rab5 with ab218624 at 1/50 dilution (red) compared with a Rabbit lgG, monoclonal [EPR25A] - Isotype Control (ab172730) (black) and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (blue). Goat anti rabbit lgG (Alexa Fluor® 488, ab150077) at 1/2000 dilution was used as the secondary antibody.



Immunoprecipitation - Anti-Rab5 antibody
[EPR21801] - Early Endosome Marker (ab218624)

Rab5 was immunoprecipitated from 0.35 mg MCF7 (human breast adenocarcinoma cell line) whole cell lysate with ab218624 at 1/30 dilution. Western blot was performed from the immunoprecipitate using ab218624 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (ab131366) was used for detection at 1/5000 dilution.

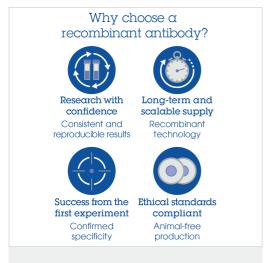
Lane 1: MCF7 whole cell lysate 10 µg (input).

Lane 2: ab218624 IP in MCF7 whole cell lysate.

Lane 3: Rabbit monoclonal IgG (<u>ab172730</u>) instead of ab218624 in MCF7 whole cell lysate.

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

Exposure time: 1 second.



Anti-Rab5 antibody [EPR21801] - Early Endosome Marker (ab218624) Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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