abcam

Product datasheet

Rabbit IgG, monoclonal [EPR25A] - Isotype Control ab172730



重组 RabMAb

457 References 15 图像

概述

产品名称 兔lgG,单克隆抗体[EPR25A] -同型对照

经测试应用 适用于: ICC/IF, Flow Cyt, IHC-P, ChIP-sequencing, IP, ChIC/CUT&RUN-seq

免疫原 Chemical/ Small Molecule conjugated to keyhole limpet haemocyanin. KLH is a copper containing

oxygen carrier occurring freely dissolved in the hemolymph of many molluscs and arthropods. KLH

forms a large complex composed of ~50 kDa subunits.

常规说明 KLH is often used in molecular immunology as a carrier protein conjugated to low molecular

> weight molecules such as peptides, amino acids, nucleic acids, drugs or toxins to render them more immunogenic due to the size of the conjugate complex and the immunogenicity of KLH.

> 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 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 EPR25A

同种型 lgG

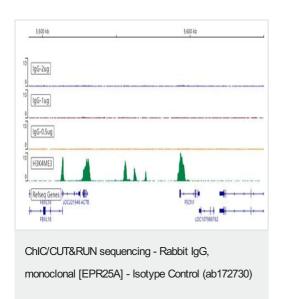
The Abpromise guarantee

Abpromise™承诺保证使用ab172730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。

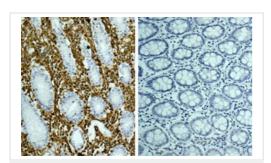
应用	Ab评论	说明
ICC/IF		Use at an assay dependent concentration. Please note: This product should be diluted to the same concentration (not dilution) of the primary antibody to be used.
Flow Cyt		Use at an assay dependent concentration. Please note: This product should be diluted to the same concentration (not dilution) of the primary antibody to be used.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. Please note: This product should be diluted to the same concentration (not dilution) of the primary antibody to be used.
ChIP-sequencing		Use at an assay dependent concentration. PubMed: 26455392
IP		Use at an assay dependent concentration.
ChIC/CUT&RUN-seq		Use at an assay dependent concentration. 0.5 µg - 2µg

图片



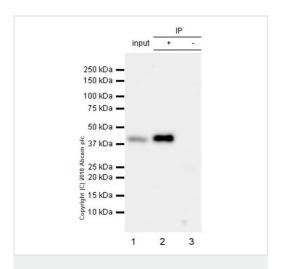
ChIC/CUT&RUN was performed using a pAG-MNAse at a final concentration of 700 ng/ μ L, 2.5 x 10^5 HeLa (Human cervix adenocarcinoma epithelial cell line) cells and 0.5 μ g, 1 μ g or 2 μ g of ab172730 [EPR25A]. The resulting DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 10 million reads. H3K4me3 (ab213224) used for comparison.

Additional screenshots of mapped reads can be downloaded here. The University of Geneva owns patents relevant to ChIC (Chromatin Immuno-Cleavage) methods.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

Immunohistochemical analysis of paraffin-embedded human colon tissue using anti-Vimentin RabMAb (<u>ab92547</u>, left panel) (brown) and Rabbit mAb IgG control (ab172730, right panel).



Immunoprecipitation - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

LINE-1 ORF1p was immunoprecipitated from 0.35 mg F9 (mouse embryonal carcinoma epithelial cell) whole cell lysate with ab216324 at 1/30 dilution. Western blot was performed from the immunoprecipitate using ab216324 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (ab131366), was used for detection at 1/5000 dilution.

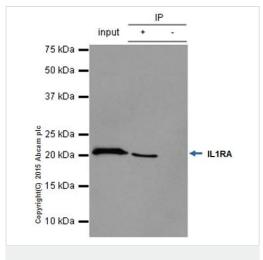
Lane 1: F9 whole cell lysate 10 µg (Input).

Lane 2: ab216324 IP in F9 whole cell lysate.

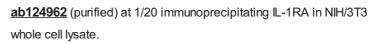
Lane 3: Rabbit monoclonal lgG (ab172730) instead of <u>ab216324</u> in F9 whole cell lysate.

Blocking/Dilution buffer: 5% NFDM/TBST.

Exposure time: 30 seconds.



Immunoprecipitation - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)



Lane 1 (input): NIH/3T3 whole cell lysate (10µg)

Lane 2 (+): <u>ab124962</u> + NIH/3T3 whole cell lysate.

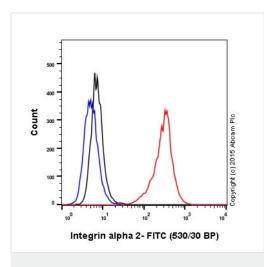
Lane 3 (-): Rabbit monoclonal IgG (ab172730) instead of ab124962 in NIH/3T3 whole cell lysate.

For western blotting, VeriBlot for IP Detection Reagent (HRP)

(ab131366), was used for detection at 1/10,000 dilution.

Blocking buffer and concentration: 5% NFDM/TBST.

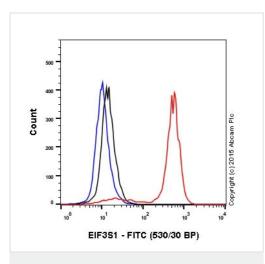
Diluting buffer and concentration: 5% NFDM /TBST.



Flow Cytometry - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

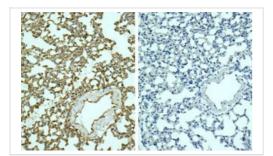
Overlay histogram showing A549 (human lung carcinoma) cells stained with <u>ab133557</u> (red line). The cells were fixed with 2% paraformaldehyde. The cells were then incubated with <u>ab133557</u> at 1/60 dilution. The secondary antibody used was goat anti-rabbit lgG (FITC) at 1/150 dilution. Isotype control antibody (black line) was rabbit lgG (monoclonal) (ab172730). Unlabelled control (blue line) was cells without incubation with primary antibody and secondary antibody. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 530/30 bandpass filter.

Conjugated versions are available for this clone: Alexa Fluor[®] 488 (<u>ab199091</u>), Alexa Fluor[®] 647 (<u>ab199093</u>), R-PE (<u>ab209478</u>), APC (<u>ab232814</u>).



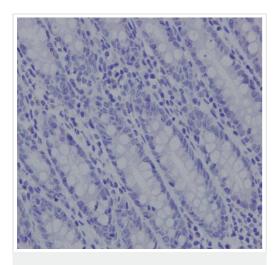
Flow Cytometry - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

Overlay histogram showing K562 (human chronic myelogenous leukemia) cells stained with ab196018 (red line). The cells were fixed with 2% paraformaldehyde. The cells were then incubated with ab196018 at 1/150 dilution. The secondary antibody used was goat anti-rabbit lgG (FITC) at 1/150 dilution. Isotype control antibody (black line) was rabbit lgG (monoclonal) (ab172730). Unlabelled control (blue line) was cells without incubation with primary antibody and secondary antibody. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 530/30 bandpass filter.



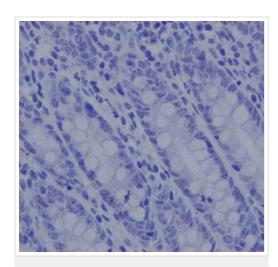
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

Immunohistochemical analysis of paraffin-embedded mouse lung tissue using anti-Vimentin RabMAb (<u>ab92547</u>, left panel) (brown) and Rabbit mAb lgG control (ab172730, right panel).



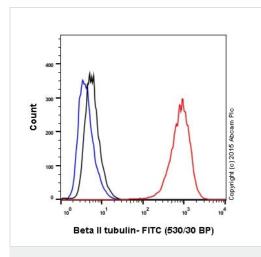
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human colon tissue with unpurified Rabbit IgG ab172730 at 1/10. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit IgG was used as the secondary antibody. Counterstained with Hematoxylin.



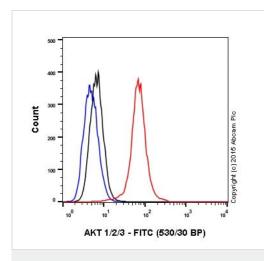
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human colon tissue with purified Rabbit IgG ab172730 at 1/100. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit IgG was used as the secondary antibody. Counterstained with Hematoxylin.



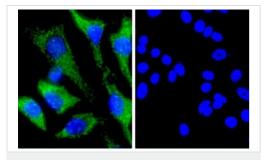
Flow Cytometry - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

Overlay histogram showing SH-SY5Y (human neuroblastoma) cells stained with <u>ab179513</u> (red line). The cells were fixed with 2% paraformaldehyde. The cells were then incubated with <u>ab179513</u> at 1/150 dilution. The secondary antibody used was goat anti-rabbit lgG (FITC) at 1/150 dilution. Isotype control antibody (black line) was rabbit lgG (monoclonal) (ab172730). Unlabelled control (blue line) was cells without incubation with primary antibody and secondary antibody. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 530/30 bandpass filter.



Flow Cytometry - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

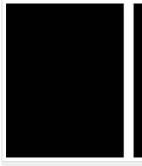
Overlay histogram showing A549 (human lung carcinoma) cells stained with <u>ab185633</u> (red line). The cells were fixed with 2% paraformaldehyde. The cells were then incubated with <u>ab185633</u> at 1/150 dilution. The secondary antibody used was goat anti-rabbit lgG (FITC) at 1/150 dilution. Isotype control antibody (black line) was rabbit lgG (monoclonal) (ab172730). Unlabelled control (blue line) was cells without incubation with primary antibody and secondary antibody. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 530/30 bandpass filter

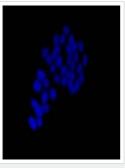


Immunocytochemistry/ Immunofluorescence - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

Immunofluorescent staining of HeLa cells using anti-AIF RabMAb (ab32516, left panel) (green) and Rabbit mAb lgG control (ab172730, right panel). DAPI nuclear staining (blue).

Conjugated versions are available for this clone: Alexa Fluor[®] 488 (<u>ab199091</u>), Alexa Fluor[®] 647 (<u>ab199093</u>), R-PE (<u>ab209478</u>), APC (<u>ab232814</u>).

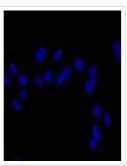




Immunocytochemistry/ Immunofluorescence - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

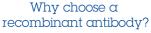
Immunocytochemistry/immunofluorescence analysis of HeLa cells with purified Rabbit lgG ab172730 at 1/100. Cells were fixed with 4% paraformaldehyde. An Alexa Fluor® 488-conjugated goat antirabbit IgG (1/200) was used as the secondary antibody. Counterstained with DAPI (blue).





Immunocytochemistry/ Immunofluorescence - Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

Immunocytochemistry/immunofluorescence analysis of HeLa cells with unpurified Rabbit IgG ab172730 at 1/10. Cells were fixed with 4% paraformaldehyde. An Alexa Fluor® 488-conjugated goat antirabbit IgG (1/200) was used as the secondary antibody. Counterstained with DAPI (blue).





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Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730)

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