abcam

Product datasheet

Anti-POMC antibody [EPR17571] - BSA and Azide free ab222486



重组 RabMAb

9 图像

概述

产品名称 Anti-POMC抗体[EPR17571] - BSA and Azide free

描述 兔单克隆抗体[EPR17571] to POMC - BSA and Azide free

宿主 Rabbit

经测试应用 适用于: IHC-P, IP, WB

种属反应性 与反应: Mouse, Rat, Human, Recombinant fragment

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

阳性对照 WB: HEK-293T transfected with POMC whole cell lysate; rat and mouse pituitary lysates; human

ACTH recombinant protein. IHC-P: Human, mouse and rat pituitary tissues. IP: Rat pituitary whole

cell lysate.

常规说明 ab222486 is the carrier-free version of ab210605.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar® is a trademark of Fluidigm Canada Inc.

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. Do Not Freeze.

存储溶液 pH: 7.2

Constituent: PBS

无载体 是

纯**度** Protein A purified

克隆 单克隆

克隆编号 EPR17571

同种型 IgG

应用

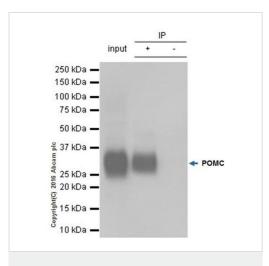
The Abpromise guarantee Abpromise™承诺保证使用ab222486于以下的经测试应用

"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		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
IP		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Detects a band of approximately 29 kDa (predicted molecular weight: 29 kDa).

靶标

图片



Immunoprecipitation - Anti-POMC antibody

[EPR17571] - BSA and Azide free (ab222486)

<u>ab210605</u> at 1/80 immunoprecipitating POMC in Rat pituitary whole cell lysate observed at 29 KDa (lanes 1 and 2).

Lane 1 (input): Rat pituitary whole cell lysate 10µg

Lane 2 (+): ab210605 + Rat pituitary whole cell lysate

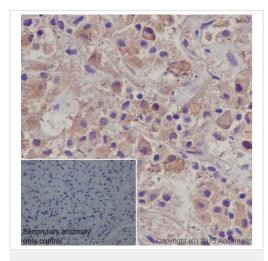
Lane 3 (-): Rabbit monoclonal lgG (ab172730) instead of

ab210605 in Rat pituitary whole cell lysate

For western blotting, <u>ab210605</u> at 1/1000 dilution and <u>ab131366</u> VeriBlot for IP (HRP) was used for detection at 1/10000 dilution.

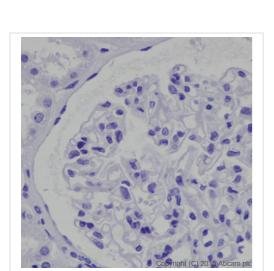
Blocking/Diluting buffer and concentration: 5% NFDM/TBST.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab210605).



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

[EPR17571] - BSA and Azide free (ab2222486)



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

[EPR17571] - BSA and Azide free (ab222486)

Immunohistochemical analysis of paraffin-embedded human pituitary tissue labeling POMC with <u>ab210605</u> at 1/16000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution.

Cytoplasmic staining on human pituitary is observed.

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab210605).

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

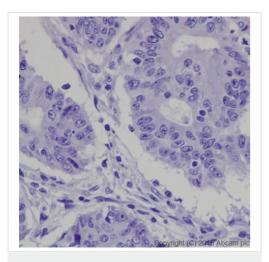
Immunohistochemical analysis of paraffin-embedded human kidney tissue labeling POMC with <u>ab210605</u> at 1/16000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution.

Negative staining on human kidney.

Counter stained with Hematoxylin.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab210605).

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



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

[EPR17571] - BSA and Azide free (ab222486)

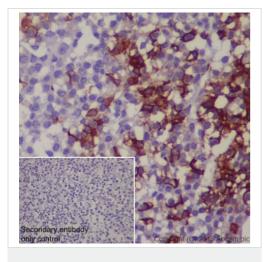
Immunohistochemical analysis of paraffin-embedded human colon cancer tissue labeling POMC with <u>ab210605</u> at 1/16000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution.

Negative staining on human colon cancer.

Counter stained with Hematoxylin.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab210605).

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



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

[EPR17571] - BSA and Azide free (ab2222486)

Immunohistochemical analysis of paraffin-embedded mouse pituitary tissue labeling POMC with <u>ab210605</u> at 1/16000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution.

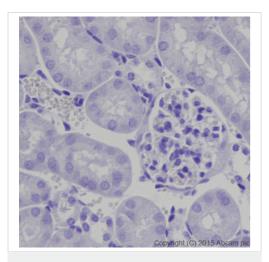
Cytoplasmic staining on mouse pituitary is observed.

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab210605).

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



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

[EPR17571] - BSA and Azide free (ab222486)

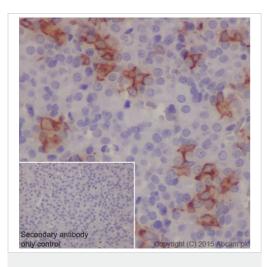
Immunohistochemical analysis of paraffin-embedded mouse kidney tissue labeling POMC with <u>ab210605</u> at 1/16000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution.

Negative staining on mouse kidney.

Counter stained with Hematoxylin.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab210605).

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



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

[EPR17571] - BSA and Azide free (ab2222486)

Immunohistochemical analysis of paraffin-embedded rat pituitary tissue labeling POMC with <u>ab210605</u> at 1/16000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution.

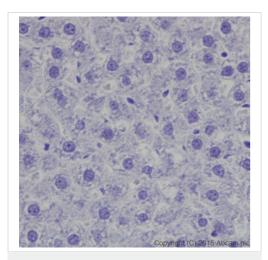
Cytoplasmic staining on rat pituitary is observed.

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab210605).

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



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

[EPR17571] - BSA and Azide free (ab222486)

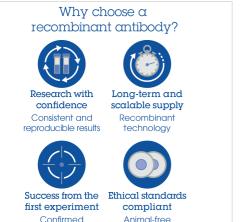
Immunohistochemical analysis of paraffin-embedded rat liver tissue labeling POMC with <u>ab210605</u> at 1/16000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution.

Negative staining on rat liver.

Counter stained with Hematoxylin.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab210605).

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



production

specificity

free (ab222486)

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Anti-POMC antibody [EPR17571] - BSA and Azide

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