abcam

Product datasheet

Anti-CD10 antibody [EPR22865-73] ab255609





重组 RabMAb

2 References 14 图像

概述

产品名称 Anti-CD10抗体[EPR22865-73]

描述 兔单克隆抗体[EPR22865-73] to CD10

宿主 Rabbit

特异性 IHC application is recommended for human only.

经测试应用 适用于: WB, IHC-P, ICC/IF, IP, mIHC

不适用于: Flow Cyt

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: Human placenta tisssue lysate; rat kidney and liver lysates; Raji, Ramos and HAP1 whole

> cell lysates; rat kidney and liver lysates; mouse kidney lysate. IHC-P: Human kidney, breast, liver and tonsil tissue, human diffuse large B-cell lymphoma ICC/IF: Raji and Ramos cells. IP: Raji

whole cell lysate. mIHC: Human breast and endometrium tissues.

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 (glycerin, glycerine), 0.05% BSA

纯度 Protein A purified

克隆 单克隆

克隆编号 EPR22865-73

同种型 IgG

应用

The Abpromise guarantee

Abpromise™承诺保证使用ab255609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
WB		1/1000. Detects a band of approximately 100 kDa (predicted molecular weight: 85 kDa).
IHC-P		1/1000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. IHC application is recommended for human only.
ICC/IF		1/50.
IP		1/30.
mIHC		1/1000.

应用说明 Is unsuitable for Flow Cyt.

靶标

功能 Thermolysin-like specificity, but is almost confined on acting on polypeptides of up to 30 amino

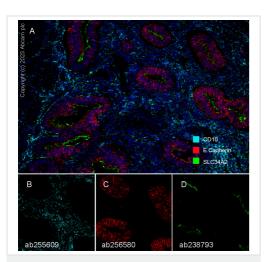
acids. Biologically important in the destruction of opioid peptides such as Met- and Leuenkephalins by cleavage of a Gly-Phe bond. Able to cleave angiotensin-1, angiotensin-2 and angiotensin 1-9. Involved in the degradation of atrial natriuretic factor (ANF). Displays UV-

inducible elastase activity toward skin preelastic and elastic fibers.

序列相似性 Belongs to the peptidase M13 family.

细**胞定位** Cell membrane.

图片



Multiplex immunohistochemistry - Anti-CD10 antibody [EPR22865-73] (ab255609)

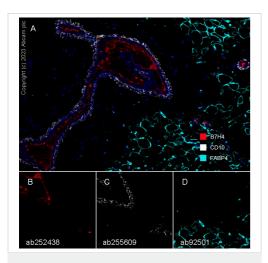
Fluorescence multiplex immunohistochemical analysis of the human endometrium (Formalin/PFA-fixed paraffin-embedded sections).

Panel A: merged staining of anti-E Cadherin (<u>ab256580</u>, red; Opal™690), anti-SLC34A2 (<u>ab238793</u>, green; Opal™520) and anti-CD10 (ab255609, cyan; Opal™570) on human endometrium. Panel B: anti-CD10 stained on stromal cells. Panel C: anti-E Cadherin stained on glandular cells. Panel D: anti-SLC34A2 stained on apical membrane of glandular cells. Opal Polymer HRP Ms + Rb was used as a secondary antibody.

The section was incubated in three rounds of staining: in the order of <u>ab256580</u> at 1/3000 dilution (0.324 μ g/ml) for 30mins, <u>ab238793</u> at 1/1000 dilution (2.26 μ g/ml) for 10mins and ab255609 at 1/1000 dilution (0.615 μ g/ml) for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

The immunostaining was performed on a Leica Biosystems $BOND^{\circledR} RX \ instrument \ with \ an \ Opal^{\intercal\intercal} \ 4-color \ kit. \ lmage \ acquisition$ was performed with Leica SP8 confocal microscope.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins. DAPI (blue) was used as a nuclear counter stain.



Multiplex immunohistochemistry - Anti-CD10 antibody [EPR22865-73] (ab255609)

1 2 3 4 5 6

250 kDa —
150 kDa —
100 kDa —
100 kDa —
50 kDa —
4 50 kDa —
20 kDa —
4 6 kDa —
20 kDa —
4 6 kDa —
4 6 APDH

Western blot - Anti-CD10 antibody [EPR22865-73] (ab255609)

Fluorescence multiplex immunohistochemical analysis of the human breast (Formalin/PFA-fixed paraffin-embedded sections).

Panel A: merged staining of anti-B7H4 (<u>ab252438</u>, red;

Opal[™]690), anti-CD10 (ab255609, gray; Opal[™]520) and antiFABP4 (<u>ab92501</u>, cyan; Opal[™]570) on human breast. Panel B:
anti-B7H4 stained on glandular lumens. Panel C: anti-CD10 stained on myoepithelial cells. Panel D: anti-FABP4 stained on adipocytes.

Opal Polymer HRP Ms + Rb was used as a secondary antibody.

The section was incubated in three rounds of staining: in the order of $\underline{ab252438}$ at 1/100 dilution (4.69 μ g/ml), ab255609 at 1/1000 dilution (0.615 μ g/ml) and $\underline{ab92501}$ at 1/10000 dilution (0.047 μ g/ml) for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

The immunostaining was performed on a Leica Biosystems ${\sf BOND}^{\circledR} \, {\sf RX} \ \text{instrument with an Opal}^{\textmd{\tiny TM}} \ 4\text{-color kit. Image acquisition}$ was performed with Leica SP8 confocal microscope.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins. DAPI (blue) was used as a nuclear counter stain.

All lanes : Anti-CD10 antibody [EPR22865-73] (ab255609) at 1/1000 dilution

Lane 1 : Human placenta tissue lysate

Lane 2: Raji (human Burkitt's lymphoma cell line) whole cell lysate

Lane 3 : Ramos (human Burkitt's lymphoma cell line) whole cell lysate

Lane 4: HT-29 (human colorectal adenocarcinoma cell line) whole cell lysate

Lane 5: Wild-type HAP1 whole cell lysate

Lane 6: CD10 knockout HAP1 whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

Lane 1 : VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) at 1/1000 dilution

Lanes 2-4: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Predicted band size: 85 kDa **Observed band size:** 100 kDa

ab255609 was shown to specifically react with CD10 in wild-type HAP1 cells as signal was lost in CD10 knockout cells. Wild-type and CD10 knockout samples were subjected to SDS-PAGE. ab255609 and <u>ab181602</u> (Rabbit anti-GAPDH loading control) were incubated 1 hour at room temperature at 1/1000 dilution and 1/200,000 dilution respectively. Blots were developed with Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated (<u>ab97051</u>) 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 instrument using the ECL technique.

Lanes 5-6 in this blot were developed using a higher sensitivity ECL substrate.

Blocking and dilution buffer: 5% NFDM/TBST.

Exposure times.

Lane 1: 10 seconds; Lanes 2-4: 15 seconds; Lanes 5-6: 70 seconds.

The molecular weight observed is consistent with what has been described in the literature (PMID:15286660).

Negative control: HT-29 (PMID:19828468).

Immunoprecipitation - Anti-CD10 antibody [EPR22865-73] (ab255609)

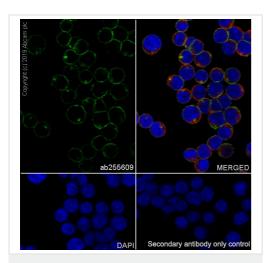
CD10 was immunoprecipitated from 0.35 mg of Raji (human Burkitt's lymphoma cell line) whole cell lysate with ab255609 at 1/30 dilution. Western blot was performed from the immunoprecipitate using ab255609 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (ab131366), was used as secondary antibody at 1/5000 dilution.

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

Lane 2: ab255609 IP in Raji whole cell lysate.

Lane 3: Rabbit monoclonal lgG (<u>ab172730</u>) instead of ab255609 in Raji whole cell lysate.

Blocking and dilution buffer and concentration: 5% NFDM/TBST. Exposure time: 30 seconds.

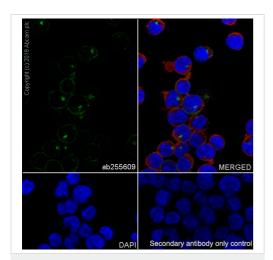


Immunocytochemistry/ Immunofluorescence - Anti-CD10 antibody [EPR22865-73] (ab255609)

Immunofluorescent analysis of 100% methanol-fixed Ramos (human Burkitt's lymphoma cell line) cells labeling CD10 with ab255609 at 1/50 dilution, followed by Goat Anti-Rabbit lgG H&L (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green). Confocal image showing cytoplasmic and membranous staining in Ramos cell line. The nuclear counter stain is DAPI (blue). Tubulin is detected with Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor® 594) (ab195889) at 1/200 dilution (red).

Secondary antibody only control: Used PBS instead of primary antibody, followed by Goat Anti-Rabbit lgG H&L (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution.

100% methanol was preferred as fixative.



Immunocytochemistry/ Immunofluorescence - Anti-CD10 antibody [EPR22865-73] (ab255609)

Immunofluorescent analysis of 100% methanol-fixed Raji (human Burkitt's lymphoma cell line) cells labeling CD10 with ab255609 at 1/50 dilution, followed by Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) (**ab150077**) secondary antibody at 1/1000 dilution (green). Confocal image showing cytoplasmic and membranous staining in Raji cell line. The nuclear counter stain is DAPI (blue). Tubulin is detected with Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor[®] 594) (**ab195889**) at 1/200 dilution (red).

Secondary antibody only control: Used PBS instead of primary antibody, followed by Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution.

100% methanol was preferred as fixative.



Western blot - Anti-CD10 antibody [EPR22865-73] (ab255609)

Anti-CD10 antibody [EPR22865-73] (ab255609) at 1/1000 dilution + Mouse kidney tissue lysate at 20 µg

Secondary

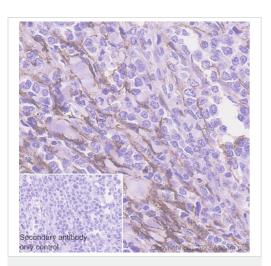
Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Predicted band size: 85 kDa **Observed band size:** 100 kDa

Exposure time: 3 minutes

Blocking and dilution buffer: 5% NFDM/TBST.

The molecular weight observed is consistent with what has been described in the literature (PMID:15286660).

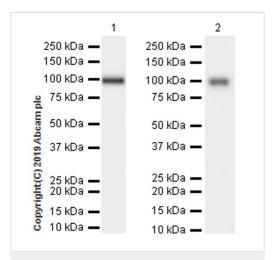


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD10 antibody
[EPR22865-73] (ab255609)

Immunohistochemical analysis of paraffin-embedded Human diffuse large B-cell lymphoma labelling CD10 with ab255609 at 1/1000 dilution, followed by a Goat Anti-Rabbit IgG H&L (HRP polymer) ready to use (ab214880). Positive staining on human diffuse large B-cell lymphoma is observed. Counter stained with hematoxylin.

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

Heat mediated antigen retrieval using ab93684 (Tris/EDTA buffer, pH 9.0).



Western blot - Anti-CD10 antibody [EPR22865-73] (ab255609)

All lanes : Anti-CD10 antibody [EPR22865-73] (ab255609) at 1/1000 dilution

Lane 1: Rat kidney tissue lysate

Lane 2: Rat liver tissue lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) ($\underline{ab97051}$) at

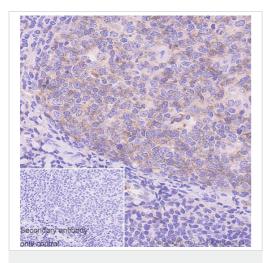
1/100000 dilution

Predicted band size: 85 kDa **Observed band size:** 100 kDa

Exposure time: 15 seconds

Blocking and dilution buffer: 5% NFDM/TBST.

The molecular weight observed is consistent with what has been described in the literature (PMID:15286660).

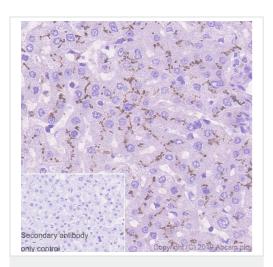


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD10 antibody
[EPR22865-73] (ab255609)

Immunohistochemical analysis of paraffin-embedded human tonsil tissue labeling CD10 with ab255609 at 1/1000 dilution, followed by a Goat Anti-Rabbit lgG H&L (HRP) ready to use. Membranous staining on germinal center of human tonsil (PMID:10843287) is observed. Counter stained with hematoxylin.

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

Heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).

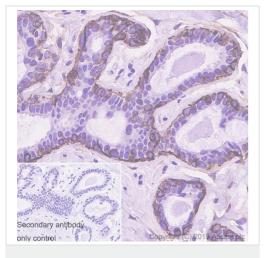


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD10 antibody
[EPR22865-73] (ab255609)

Immunohistochemical analysis of paraffin-embedded human liver tissue labeling CD10 with ab255609 at 1/1000 dilution, followed by a Goat Anti-Rabbit IgG H&L (HRP) ready to use. Membranous staining on intrahepatic canaliculi of human liver (PMID:10705818) is observed. Counter stained with hematoxylin.

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

Heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).



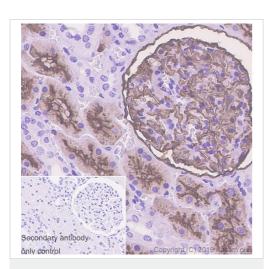
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD10 antibody
[EPR22865-73] (ab255609)

Immunohistochemical analysis of paraffin-embedded human breast tissue labeling CD10 with ab255609 at 1/1000 dilution, followed by a ready to use Goat Anti-Rabbit IgG H&L (HRP) ready to use.

Membranous staining on myoepithelial cells of human breast (PMID:10705818, 17143263) is observed. Counter stained with hematoxylin.

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

Heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).

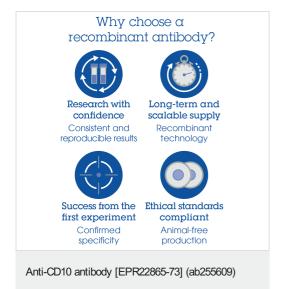


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD10 antibody
[EPR22865-73] (ab255609)

Immunohistochemical analysis of paraffin-embedded human kidney tissue labeling CD10 with ab255609 at 1/1000 dilution, followed by a Goat Anti-Rabbit IgG H&L (HRP) ready to use. Positive staining on proximal convoluted tubules and glomerular epithelial cells of human kidney (PMID:10705818) is observed. Counter stained with hematoxylin.

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

Heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).



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