

## Product datasheet

### Anti-CD10 antibody [EPR22865-73] ab255609

KO VALIDATED

Recombinant

RabMAb

[2 References](#) [14 Images](#)

#### Overview

|                            |   |
|----------------------------|---|
| <b>Product name</b>        | Anti-CD10 antibody [EPR22865-73]  |
| <b>Description</b>         | Rabbit monoclonal [EPR22865-73] to CD10   |
| <b>Host species</b>        | Rabbit  |
| <b>Specificity</b>         | IHC application is recommended for human only.  |
| <b>Tested applications</b> | <b>Suitable for:</b> WB, IHC-P, ICC/IF, IP, mIHC<br><b>Unsuitable for:</b> Flow Cyt   |
| <b>Species reactivity</b>  | <b>Reacts with:</b> Mouse, Rat, Human   |
| <b>Immunogen</b>           | Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.   |
| <b>Positive control</b>    | WB: Human placenta tissssue 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.   |
| <b>General notes</b>       | <p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> <li>- High batch-to-batch consistency and reproducibility</li> <li>- Improved sensitivity and specificity</li> <li>- Long-term security of supply</li> <li>- Animal-free production</li> </ul> <p>For more information <a href="#">see here</a>.</p> <p>Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <a href="#">RabMAb<sup>®</sup> patents</a>.</p> |

#### Properties

|                             |   |
|-----------------------------|---|
| <b>Form</b>                 | Liquid  |
| <b>Storage instructions</b> | 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. |
| <b>Storage buffer</b>       | pH: 7.2<br>Preservative: 0.01% Sodium azide<br>Constituents: PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA                   |
| <b>Purity</b>               | Protein A purified  |

|                     |             |
|---------------------|-------------|
| <b>Clonality</b>    | Monoclonal  |
| <b>Clone number</b> | EPR22865-73 |
| <b>Isotype</b>      | IgG         |

## Applications

**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab255609 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

| Application   | Abreviews | Notes   |
|---------------|-----------|---|
| <b>WB</b>     |           | 1/1000. Detects a band of approximately 100 kDa (predicted molecular weight: 85 kDa).   |
| <b>IHC-P</b>  |           | 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. |
| <b>ICC/IF</b> |           | 1/50.   |
| <b>IP</b>     |           | 1/30.   |
| <b>mlHC</b>   |           | 1/1000.   |

**Application notes** Is unsuitable for Flow Cyt.

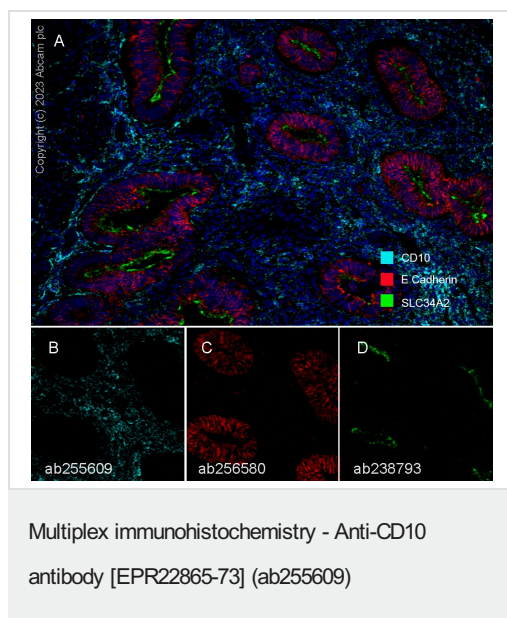
## Target

**Function** 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 Leu-enkephalins 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.

**Sequence similarities** Belongs to the peptidase M13 family.

**Cellular localization** Cell membrane.

## Images



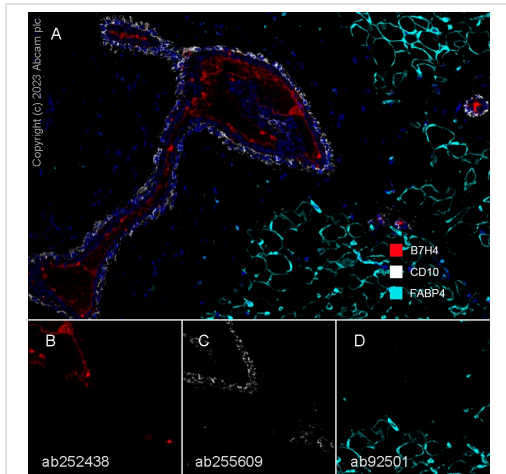
Fluorescence multiplex immunohistochemical analysis of the human endometrium (Formalin/PFA-fixed paraffin-embedded sections).

Panel A: merged staining of anti-E Cadherin (**ab256580**, red; Opal™690), anti-SLC34A2 (**ab238793**, 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 **ab256580** at 1/3000 dilution (0.324 µg/ml) for 30mins, **ab238793** at 1/1000 dilution (2.26 µg/ml) for 10mins and ab255609 at 1/1000 dilution (0.615 µg/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® RX instrument with an Opal™ 4-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.



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

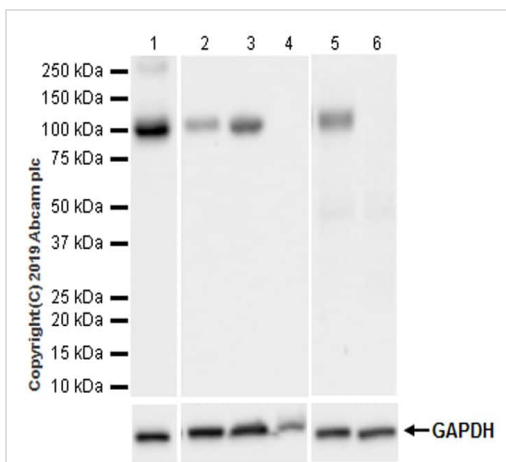
Fluorescence multiplex immunohistochemical analysis of the human breast (Formalin/PFA-fixed paraffin-embedded sections).

Panel A: merged staining of anti-B7H4 ([ab252438](#), red; Opal™690), anti-CD10 (ab255609, gray; Opal™520) and anti-FABP4 ([ab92501](#), 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 [ab252438](#) at 1/100 dilution (4.69 µg/ml), ab255609 at 1/1000 dilution (0.615 µg/ml) and [ab92501](#) at 1/10000 dilution (0.047 µg/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® RX instrument with an Opal™ 4-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.



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

**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) ([ab131366](#)) 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 **ab181602** (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 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 instrument using the ECL technique. Lanes 5-6 in this blot were developed using a higher sensitivity ECL substrate.

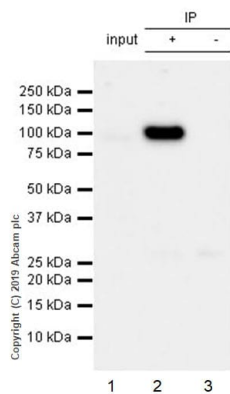
Blocking and dilution buffer: 5% NFDM/TBST.

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.

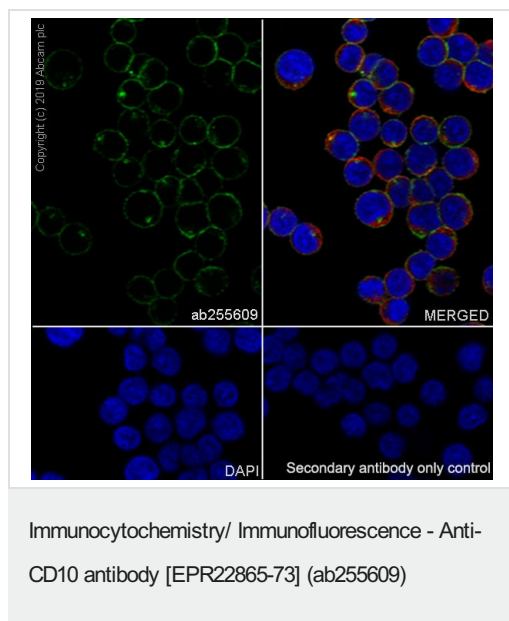
**Lane 1:** Raji whole cell lysate 10 µg (Input).

**Lane 2:** ab255609 IP in Raji whole cell lysate.

**Lane 3:** Rabbit monoclonal IgG (**ab172730**) instead of ab255609 in Raji whole cell lysate.

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

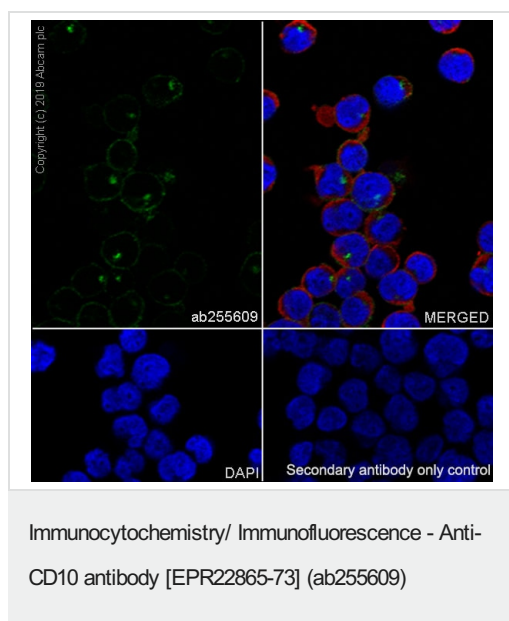
Exposure time: 30 seconds.



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 IgG 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 IgG H&L (Alexa Fluor® 488) (**ab150077**) secondary antibody at 1/1000 dilution.

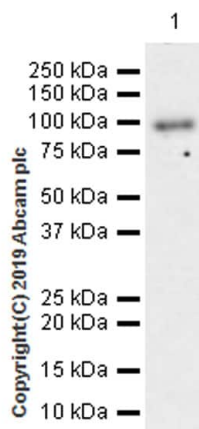
100% methanol was preferred as fixative.



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 IgG 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

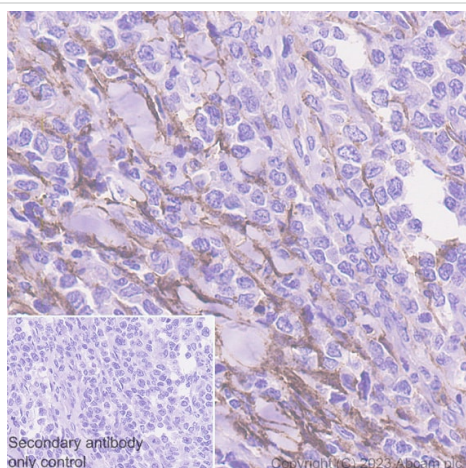
**Predicted band size:** 85 kDa

**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).



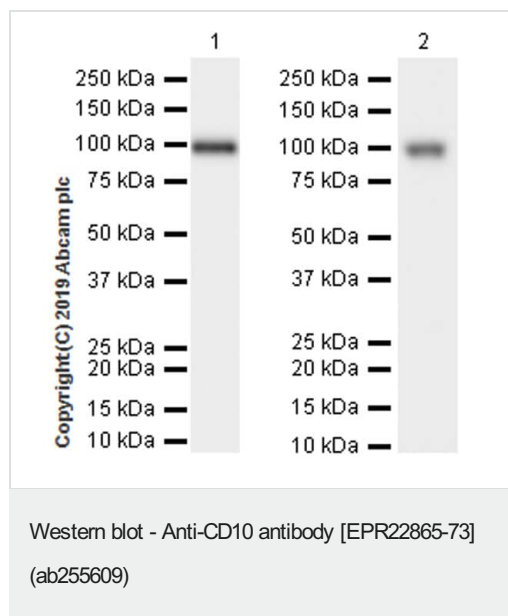
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-CD10 antibody  
[EPR22865-73] (ab255609)

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).





**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 IgG H&L (HRP) ([ab97051](#)) at 1/100000 dilution

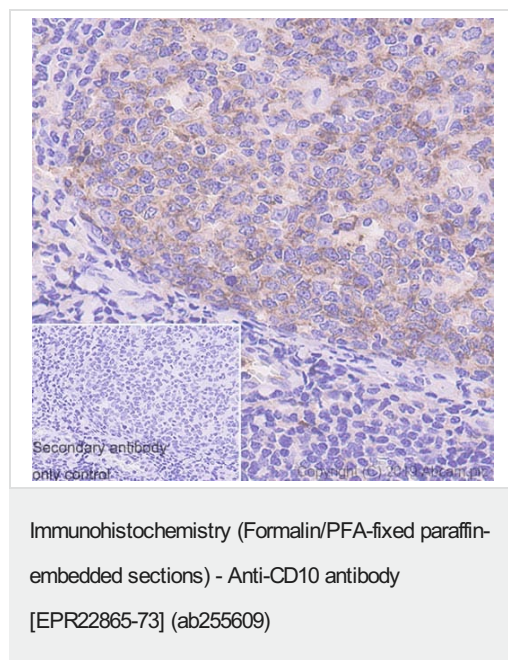
**Predicted band size:** 85 kDa

**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).

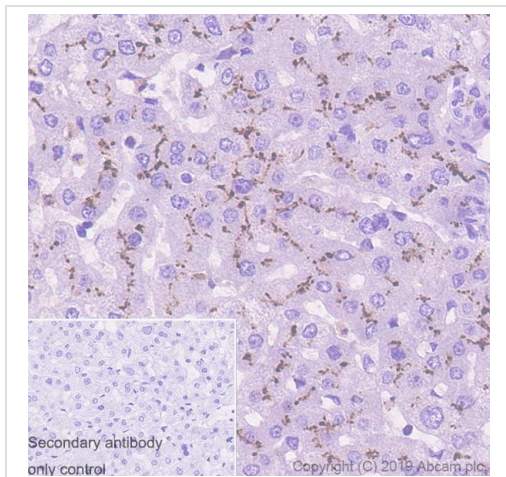


Immunohistochemical analysis of paraffin-embedded human tonsil tissue labeling CD10 with ab255609 at 1/1000 dilution, followed by a Goat Anti-Rabbit IgG H&L (HRP) ready to use. Membranous staining on germinal center of human tonsil (PMID:10843287) is observed. Counter stained with hematoxylin.

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

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



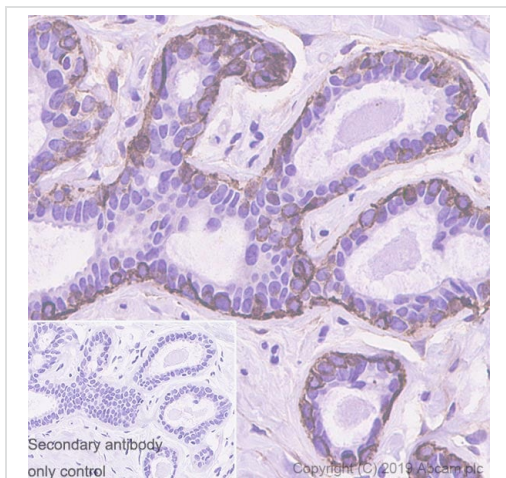


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-CD10 antibody [EPR22865-73] (ab255609)

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 IgG H&L (HRP) ready to use.

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

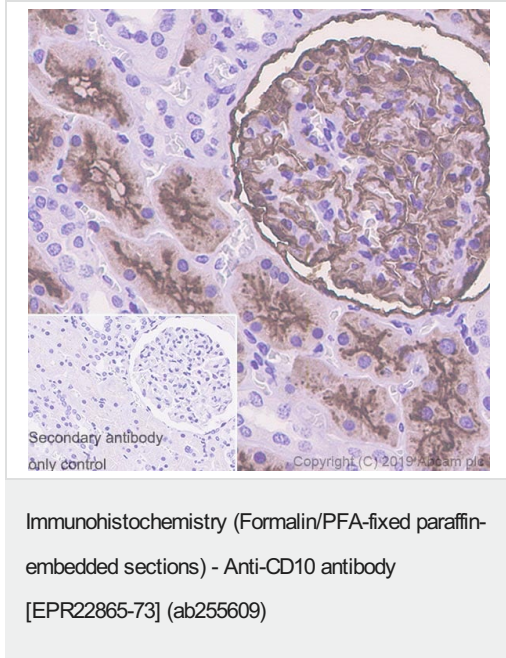


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-CD10 antibody [EPR22865-73] (ab255609)

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 IgG H&L (HRP) ready to use.

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



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 IgG H&L (HRP) ready to use.

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

Why choose a recombinant antibody?

|  |  |
|--|--|
|  <p><b>Research with confidence</b><br/>Consistent and reproducible results</p> |  <p><b>Long-term and scalable supply</b><br/>Recombinant technology</p> |
|  <p><b>Success from the first experiment</b><br/>Confirmed specificity</p>      |  <p><b>Ethical standards compliant</b><br/>Animal-free production</p>   |

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

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