abcam

Product datasheet

Anti-PAX8 antibody [EPR18715] ab191870

Recombinant RabMAb

7 References 11 Images

Overview

Product name Anti-PAX8 antibody [EPR18715]

Description Rabbit monoclonal [EPR18715] to PAX8

Host species Rabbit

Tested applications Suitable for: IHC-Fr, ChIC/CUT&RUN-seq, WB, IHC-P, ICC/IF

Species reactivity Reacts with: Mouse, Human

Immunogen Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: Human thyroid cancer lysate; Human fetal kidney lysate; NIH:OVCAR-3 and SK-OV-3 whole

cell lysates. IHC-P: Human thyroid carcinoma and endometrium carcinoma tissues. ICC/IF:

NIH:OVCAR-3 and SK-OV-3 cells.

General notes This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

Improved sensitivity and specificityLong-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**.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

Purity Protein A purified

Clonality Monoclonal
Clone number EPR18715

1

Isotype IgG

Applications

The Abpromise guarantee

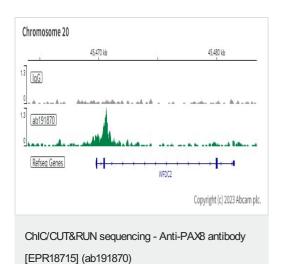
Our <u>Abpromise guarantee</u> covers the use of ab191870 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

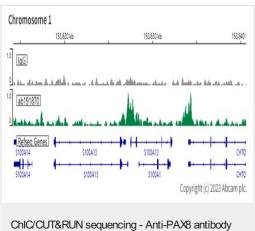
Application	Abreviews	Notes
IHC-Fr		1/500.
ChIC/CUT&RUN-seq		Use at an assay dependent concentration.
WB		1/1000. Detects a band of approximately 48 kDa (predicted molecular weight: 48 kDa).
IHC-P		1/1000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
ICC/IF		1/1000.

Target		
Function	Transcription factor for the thyroid-specific expression of the genes exclusively expressed in the thyroid cell type, maintaining the functional differentiation of such cells.	
Tissue specificity	Expressed in the excretory system, thyroid gland and Wilms tumors.	
Involvement in disease	Defects in PAX8 are the cause of congenital hypothyroidism non-goitrous type 2 (CHNG2) [MlM:218700]. CHNG2 is a disease characterized by thyroid dysgenesis, the most frequent cause of congenital hypothyroidism, accounting for 85% of case. The thyroid gland can be completely absent (athyreosis), ectopically located and/or severely hypoplastic. Ectopic thyroid gland is the most frequent malformation, with thyroid tissue being found most often at the base of the tongue.	
Sequence similarities	Contains 1 paired domain.	
Developmental stage	In developing excretory system, during thyroid differentiation and in adult thyroid.	
Cellular localization	Nucleus.	

Images



ChIC/CUT&RUN was performed using a pAG-MNase at a final concentration of 700 ng/ μ L, 2.5 x 10^5 NIH:OVCAR-3 (Human ovary adenocarcinoma epithelial cell) cells and 5 μ g of ab191870 [EPR18715]. The resulting DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 10 million reads. The negative IgG control **ab172730** is also shown. The University of Geneva owns patents relevant to ChIC (Chromatin Immuno-Cleavage) methods.

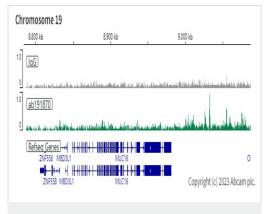


adenocarcinoma epithelial cell) cells and 5 μg of ab191870 [EPR18715]. The resulting DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 10 million reads. The negative IgG control <u>ab172730</u> is also shown. The University of Geneva owns patents relevant to ChIC (Chromatin Immuno-Cleavage) methods.

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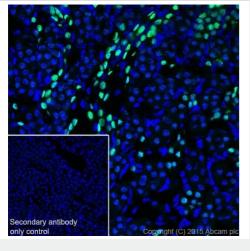
ChIC/CUT&RUN was performed using a pAG-MNase at a final

ChIC/CUT&RUN sequencing - Anti-PAX8 antibody [EPR18715] (ab191870)

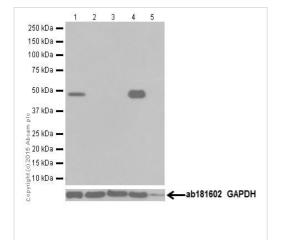


ChIC/CUT&RUN sequencing - Anti-PAX8 antibody [EPR18715] (ab191870)

ChIC/CUT&RUN was performed using a pAG-MNase at a final concentration of 700 ng/ μ L, 2.5 x 10^5 NIH:OVCAR-3 (Human ovary adenocarcinoma epithelial cell) cells and 5 μ g of ab191870 [EPR18715]. The resulting DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 10 million reads. The negative IgG control **ab172730** is also shown. The University of Geneva owns patents relevant to ChIC (Chromatin Immuno-Cleavage) methods.



Immunohistochemistry (Frozen sections) - Anti-PAX8 antibody [EPR18715] (ab191870) Immunohistochemistry (Frozen sections) analysis of mouse kidney tissue labelling PAX8 with ab191870 at a dilution of 1/500. **ab150077**, an Alexa Fluor[®] 488-conjugated goat anti-rabbit lgG was used as the secondary antibody (1/1000). Nuclear staining is visible in the mouse kidney.



Western blot - Anti-PAX8 antibody [EPR18715] (ab191870)

All lanes : Anti-PAX8 antibody [EPR18715] (ab191870) at 1/1000 dilution

Lane 1: Human thyroid cancer lysate

Lane 2: Human fetal liver lysate

Lane 3: Human fetal heart lysate

Lane 4: Human fetal kidney lysate

Lane 5: Human fetal spleen lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Anti-Rabbit $\lg G$ (HRP), specific to the non-reduced form of $\lg G$ at 1/10000 dilution

Predicted band size: 48 kDa

Observed band size: 48 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

PAX8 is expressed in the excretory system and thyroid gland.

(PMID: 1723950, 9590297, 1069301)

All lanes : Anti-PAX8 antibody [EPR18715] (ab191870) at 1/1000 dilution

Lane 1 : NIH:OVCAR-3 (Human ovary adenocarcinoma cell line) whole cell lysate

Lane 2: SK-OV-3 (Human ovarian cancer cell line) whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes: Goat Anti-Rabbit lgG, (H+L),Peroxidase conjugated at

1/10000 dilution

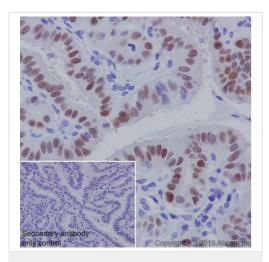
Predicted band size: 48 kDa **Observed band size:** 48 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

1 2
250 kDa —
150 kDa —
100 kDa —
75 kDa —
50 kDa —
37 kDa —
25 kDa —
20 kDa —
15 kDa —
10 kDa —
10 kDa —

Western blot - Anti-PAX8 antibody [EPR18715] (ab191870)

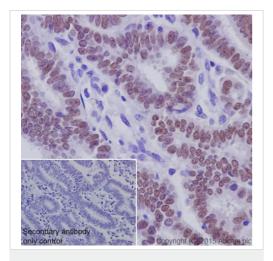


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PAX8 antibody
[EPR18715] (ab191870)

Immunohistochemical analysis of paraffin-embedded Human thyroid carcinoma tissue labeling PAX8 with ab191870 at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nucleus staining on tumor cells of thyroid carcinoma 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.

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

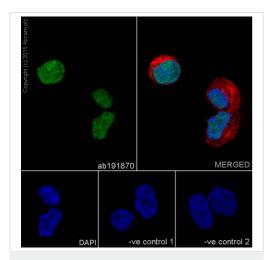


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PAX8 antibody
[EPR18715] (ab191870)

Immunohistochemical analysis of paraffin-embedded Human endometrium carcinoma tissue labeling PAX8 with ab191870 at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nucleus staining on tumor cells of the endometrium carcinoma 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.

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



Immunocytochemistry/ Immunofluorescence - Anti-PAX8 antibody [EPR18715] (ab191870)

ab191870 MERGED

DAPI -ve control 1 -ve control 2

Immunocytochemistry/ Immunofluorescence - Anti-PAX8 antibody [EPR18715] (ab191870)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized NIH:OVCAR-3 (Human ovary adenocarcinoma cell line) cells labeling PAX8 with ab191870 at 1/1000 dilution, followed by Goat anti-rabbit lgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green). Confocal image showing nucleus staining on NIH:OVCAR-3 cell line. The nuclear counter stain is DAPI (blue). Tubulin is detected with ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution and ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution (red).

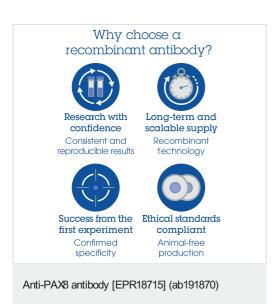
The negative controls are as follows:

-ve control 1: ab191870 at 1/1000 dilution followed by <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution. -ve control 2: <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> (Alexa Fluor®488 Goat Anti-Rabbit lgG H&L) at 1/1000 dilution.

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized SK-OV-3 (Human ovarian cancer cell line) cells labeling PAX8 with ab191870 at 1/1000 dilution, followed by Goat anti-rabbit lgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green). Confocal image showing nucleus staining on SK-OV-3 cell line. The nuclear counter stain is DAPI (blue). Tubulin is detected with ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution and ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution (red).

The negative controls are as follows:

-ve control 1: ab191870 at 1/1000 dilution followed by <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution. -ve control 2: <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> (Alexa Fluor®488 Goat Anti-Rabbit lgG H&L) at 1/1000 dilution.



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