abcam

Product datasheet

Anti-PAX8 antibody [PAX8R1] ab53490

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Overview

Product name Anti-PAX8 antibody [PAX8R1]

Description Mouse monoclonal [PAX8R1] to PAX8

Host species Mouse

Tested applications Suitable for: Flow Cyt, IHC-P, WB, IP, ICC

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Recombinant fragment corresponding to Human PAX8 (C terminal).

Database link: Q06710

Positive control HeLa, NIH3T3 or F2408 nuclear cell lysate HeLa cells RIPA lysate of HeLa cells

General notes IHC-P is not batch-tested and therefore variability between batches might be seen for these

applications.

This product was changed from ascites to tissue culture supernatant on 9th August 2018. Please note that the dilutions may need to be adjusted accordingly. If you have any questions, please do

not hesitate to contact our scientific support team.

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw

cycles.

Storage buffer pH: 7.40

Preservative: 0.05% Sodium azide Constituents: 1% BSA, PBS

Purity Protein G purified

Purification notes Purified from TCS

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ClonalityMonoclonalClone numberPAX8R1IsotypeIgG1

Applications

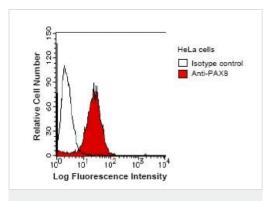
The Abpromise guarantee Our <u>Abpromise guarantee</u> covers the use of ab53490 in the following tested applications.

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

Application	Abreviews	Notes
Flow Cyt		Use at an assay dependent concentration. 0.5-2 µg/sample
IHC-P	★★★★ <u>(2)</u>	Use a concentration of 5 µg/ml. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
WB		Use a concentration of 0.2 - 2 µg/ml. Predicted molecular weight: 48 kDa.
IP		Use at an assay dependent concentration. 1-5 μg/sample
ICC		Use a concentration of 2 - 100 μg/ml.

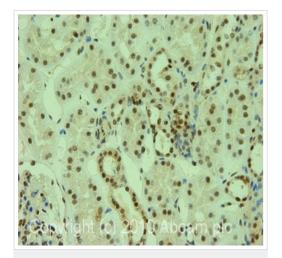
Target		
Function	Transcription factor for the thyroid-specific expression of the genes exclusively expressed in the thyroid cell type, maintaining the functional differentiation of such cells.	
Tissue specificity	Expressed in the excretory system, thyroid gland and Wilms tumors.	
Involvement in disease	Defects in PAX8 are the cause of congenital hypothyroidism non-goitrous type 2 (CHNG2) [MIM: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



Flow Cytometry - Anti-PAX8 antibody [PAX8R1] (ab53490)

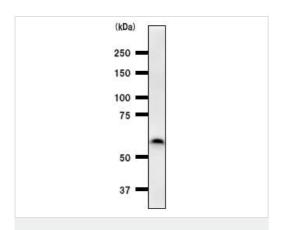
HeLa cells were fixed in 2% paraformaldehyde/PBS and then permeabilized in 90% methanol. Cells were stained with ab53490 at 0.5 μ g/sample (shaded) or isotype control (unshaded) followed by Alexa Fluor® 488-conjugated goat anti-mouse lgG (0.25 μ g)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PAX8 antibody [PAX8R1] (ab53490)

IHC image of ab53490 staining in human kidney formalin fixed paraffin embedded tissue section, performed on a Leica Bond TM system using the standard protocol F. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab53490, 5 μ g/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.



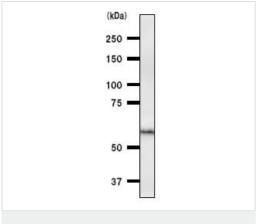
Western blot - Anti-PAX8 antibody [PAX8R1] (ab53490)

Anti-PAX8 antibody [PAX8R1] (ab53490) at 1/100 dilution + HeLa whole cell lysate at 50 μg

Secondary

Mouse IgG antibody at 1/2500 dilution

Predicted band size: 48 kDa **Observed band size:** 60 kDa



NIH3T3 whole cell lysate at 25 μg

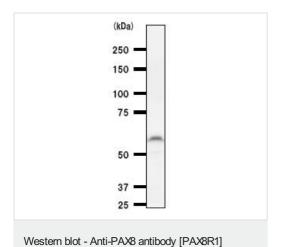
Secondary

Mouse IgG antibody at 1/2500 dilution

Anti-PAX8 antibody [PAX8R1] (ab53490) at 1/100 dilution +

Predicted band size: 48 kDa **Observed band size:** 60 kDa

Western blot - Anti-PAX8 antibody [PAX8R1] (ab53490)

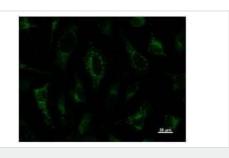


Anti-PAX8 antibody [PAX8R1] (ab53490) at 1/100 dilution + F2408 whole cell lysate at 25 µg

Secondary

Mouse IgG antibody at 1/2500 dilution

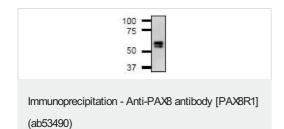
Predicted band size: 48 kDa **Observed band size:** 60 kDa



Immunocytochemistry - Anti-PAX8 antibody [PAX8R1] (ab53490)

(ab53490)

ab53490, at a 1/10 dilution, staining PAX8 in HeLa cells, using Immunocytochemistry, plus AlexaFluor®488 Goat Anti-mouse IgG secondary anitbody.



Immunoprecipitation: ab53490: $5\mu g$, RIPA lysate of HeLa cells: $500\mu g$. WB: ab53490 at a 1:100 dilution, Secondary antibody - antimouse lgG at a 1:1000 dilution

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