abcam

Product datasheet

Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] ab200829



10 Images

Overview

Product name Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504]

Description Rabbit monoclonal [EPR14504] to NFIB / NF1B2 + NFIC/CTF

Host species Rabbit

Tested applications Suitable for: IP, Flow Cyt (Intra), WB, IHC-P, ICC/IF

Species reactivity Reacts with: Mouse, Rat, Human

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

Positive control WB: HeLa, HepG2, MCF7, C6, RAW 264.7, PC-12 and NIH/3T3 whole cell lysates; Mouse heart,

> kidney and spleen lysates; Rat brain, heart and spleen lysates. IHC-P: Human cervix carcinoma and breast carcinoma, and mouse cardiac muscle tissues. ICC/IF: HeLa cells. Flow Cyt (intra):

HeLa cells. IP: HeLa whole cell lysate.

General notes 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**.

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 long

term. 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

ClonalityMonoclonalClone numberEPR14504

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab200829 in the following tested applications.

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

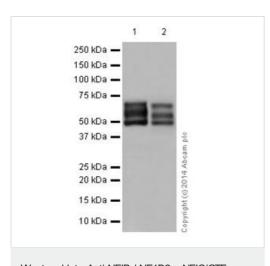
| Application | Abreviews | Notes |
|------------------|-----------|---|
| IP | | 1/70. |
| Flow Cyt (Intra) | | 1/200. |
| WB | | 1/2000. Detects a band of approximately 45-65 kDa (predicted molecular weight: 56, 47 kDa). |
| IHC-P | | 1/500. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. |
| ICC/IF | | 1/1000. |

Target

Cellular localization

NFIB / NF1B2: Nucleus. NFIC/CTF: Nucleus.

Images



Western blot - Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] (ab200829)

All lanes: Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] (ab200829) at 1/10000 dilution

Lane 1 : HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate

Lane 2 : MCF7 (Human breast adenocarcinoma cell line) whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 56, 47 kDa **Observed band size:** 45-65 kDa

Exposure time: 30 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

Based on the sequence analysis, ab200829 recognizes six isoforms within human with predicted Mw's of 56KDa, 55KDa, 48KDa, 45KDa, 48KDa and 49KDa, respectively.

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cells from cervix adenocarcinoma) cells labeling NFIB / NF1B2 + NFIC with ab200829 at 1/1000 dilution, followed by Goat anti-rabbit lgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/500 dilution (green). Nuclear staining on HeLa cell line is observed. The nuclear counterstain 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 1/500 dilution (red).

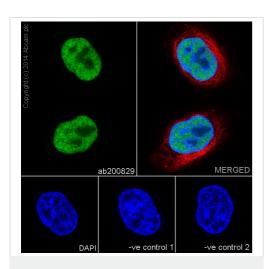
The negative controls are as follows:-

-ve control 1: ab200829 at 1/1000 dilution followed by <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 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 1/500 dilution.

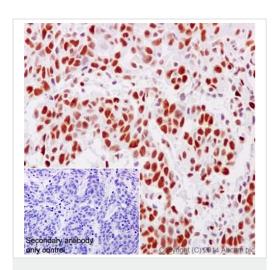
Immunohistochemical analysis of paraffin-embedded human cervix carcinoma tissue labeling NFIB / NF1B2 + NFIC with ab200829 at 1/1000 dilution followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear staining on Human cervix carcinoma tissue is observed. Counter stained with Hematoxylin.

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.

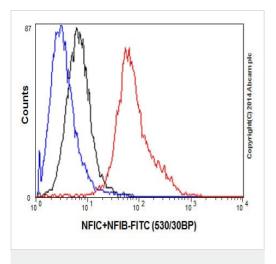
Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunocytochemistry/ Immunofluorescence - Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] (ab200829)

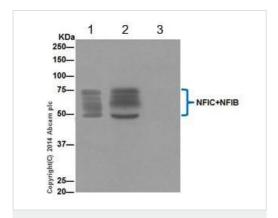


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] (ab200829)



Flow Cytometry (Intracellular) - Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] (ab200829)

Intracellular flow cytometric analysis of 2% paraformaldehyde-fixed HeLa (Human epithelial cells from cervix adenocarcinoma)cells labeling NFIB / NF1B2 + NFICwith ab200829 at 1/200 dilution (red) compared with a rabbit monoclonal IgG isotype control (ab172730; black) and an unlabelled control (cells without incubation with primary antibody and secondary antibody; blue). Goat anti rabbit IgG (FITC) at 1/150 dilution was used as the secondary antibody.



Immunoprecipitation - Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] (ab200829)

NFIB/NF1B2 + NFIC was immunoprecipitated from 1mg of HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate with ab200829 at 1/70 dilution. Western blot was performed from the immunoprecipitate using ab200829 at 1/5000 dilution. Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG, was used as secondary antibody at 1/1500 dilution.

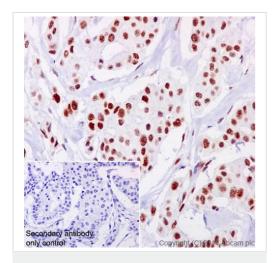
Lane 1: HeLa whole cell lysate, 10 μg (Input).

Lane 2: ab200829 IP in HeLa whole cell lysate.

Lane 3: Rabbit monoclonal $\lg G$ ($\underline{ab172730}$) instead of ab200829 in HeLa whole cell lysate.

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

Exposure time: 10 seconds.

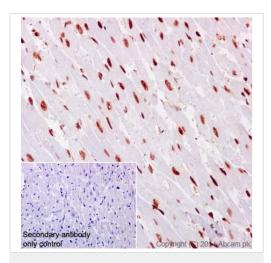


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] (ab200829)

Immunohistochemical analysis of paraffin-embedded Human breast carcinoma tissue labeling NFIB / NF1B2 + NFIC with ab200829 at 1/1000 dilution followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear staining on Human breast carcinoma tissue 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.

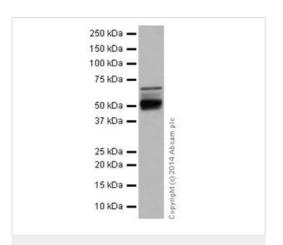


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] (ab200829)

Immunohistochemical analysis of paraffin-embedded Mouse cardiac muscle tissue labeling NFIB / NF1B2 + NFIC with ab200829 at 1/1000 dilution followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear staining on mouse cardiac muscle tissue 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.



Western blot - Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] (ab200829)

Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] (ab200829) at 1/10000 dilution + HepG2 (Human liver hepatocellular carcinoma cell line) cell lysate at 20 µg

Secondary

Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 56, 47 kDa **Observed band size:** 45-65 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer:5% NFDM /TBST

Based on the sequence analysis, ab200829 recognizes six isoforms within human with predicted Mw's of 56KDa, 55KDa, 48KDa, 45KDa, 48KDa and 49KDa, respectively.

All lanes : Anti-NFIB / NF1B2 + NFIC/CTF antibody [EPR14504] (ab200829) at 1/2000 dilution

Lane 1: Mouse heart lysate

Lane 2 : Mouse kidney lysate

Lane 3: Mouse spleen lysate

Lane 4: Rat brain lysate

Lane 5: Rat heart lysate

Lane 6: Rat spleen lysate

Lane 7: C6 (Rat glial tumor cells) whole cell lysate

Lane 8: RAW 264.7 (Mouse macrophage cells transformed with

Abelson murine leukemia virus) whole cell lysate

Lane 9: PC-12 (Rat adrenal gland pheochromocytoma) whole cell

lysate

Lane 10: NIH/3T3 (Mouse embyro fibroblast cells) whole cell lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 56, 47 kDa **Observed band size:** 45-65 kDa

Exposure time: 1 minute

Blocking/Dilution buffer: 5% NFDM/TBST.

Based on the sequence analysis, ab200829 recognizes six isoforms within human with predicted Mw's of 56KDa, 55KDa, 48KDa, 45KDa, 48KDa and 49KDa, respectively.

3

250 kDa -

150 kDa = 100 kDa = 75 kDa =

50 kDa -

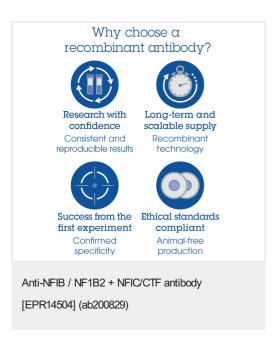
37 kDa -

25 kDa -

20 kDa -

5 6

7 8 9 10



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