abcam

Product datasheet

Anti-HNRNPA0 antibody [EP16085] ab197023





1 References 9 Images

Overview

Product name Anti-HNRNPA0 antibody [EP16085]

Rabbit monoclonal [EP16085] to HNRNPA0 **Description**

Host species Rabbit

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

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: HEK293T, HeLa, Jurkat, A549, 293 Mouse brain, Rat brain, C6, RAW 264.7, PC-12 and

NIH/3T3 lysates. IHC-P: Human cerebral cortex and Mouse spleen tissues. ICC/IF and Flow Cyt

(intra): HeLa 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 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

Clonality Monoclonal Clone number EP16085

Applications

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab197023 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 (Intra)		1/150.
ICC/IF		1/1000.
IHC-P		1/1000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
WB		1/1000. Detects a band of approximately 32, 34 kDa (predicted molecular weight: 31 kDa).

Target

Function This protein is a component of ribonucleosomes.

Sequence similarities Contains 2 RRM (RNA recognition motif) domains.

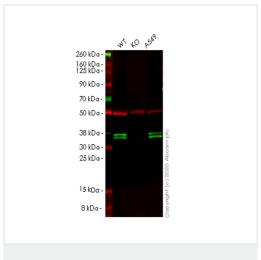
Post-translational modifications

Arg-291 is dimethylated, probably to asymmetric dimethylarginine.

Cellular localization

Nucleus. Component of ribonucleosomes.

Images



Western blot - Anti-HNRNPA0 antibody [EP16085] (ab197023)

All lanes : Anti-HNRNPA0 antibody [EP16085] (ab197023) at 1/1000 dilution

Lane 1: Wild-type HEK293T cell lysate

Lane 2: HNRNPA0 knockout HEK293T cell lysate

Lane 3: A549 cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

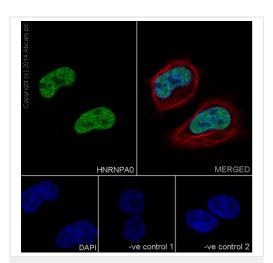
All lanes : Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (**ab216773**) at 1/10000 dilution

Predicted band size: 31 kDa

Observed band size: 32,34 kDa

Lanes 1-3: Merged signal (red and green). Green - ab197023 observed at 32,34 kDa. Red - loading control <u>ab8245</u> observed at 36 kDa.

ab197023 Anti-HNRNPA0 antibody [EP16085] was shown to specifically react with HNRNPA0 in wild-type HEK293T cells. Loss of signal was observed when knockout cell line ab266314 (knockout cell lysate ab257989) was used. Wild-type and HNRNPA0 knockout samples were subjected to SDS-PAGE. ab197023 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.

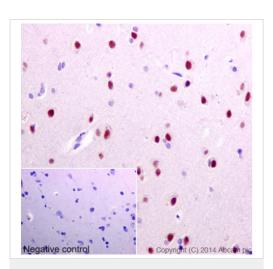


Immunocytochemistry/ Immunofluorescence - Anti-HNRNPA0 antibody [EP16085] (ab197023)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cells from cervix adenocarcinoma) cells labeling HNRNPA0 with ab197023 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 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 1/500 dilution (red).

The negative controls are as follows;

-ve control 1: ab197023 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 1/400 dilution.

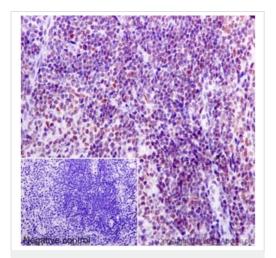


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-HNRNPA0 antibody
[EP16085] (ab197023)

Immunohistochemical analysis of paraffin-embedded
Human cerebral cortex labeling HNRNPA0 with ab197023 at
1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP)
(ab97051) at 1/500 dilution. Nuclear staining on Human cerebral cortex tissue is observed. Counter stained with Hematoxylin.

Negative control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution.

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

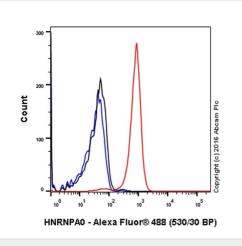


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-HNRNPA0 antibody
[EP16085] (ab197023)

Immunohistochemical analysis of paraffin-embedded Mouse spleen labeling HNRNPA0 with ab197023 at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear staining on Mouse spleen tissue is observed. Counter stained with Hematoxylin.

Negative control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution.

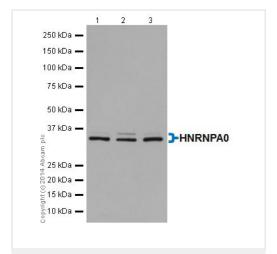
Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Flow Cytometry (Intracellular) - Anti-HNRNPA0

antibody [EP16085] (ab197023)

Intracellular Flow Cytometry analysis of HeLa cells labelling HNRNPA0 (red) with purified ab197023 at dilution of 1/150. The secondary antibody used was Alexa Fluorr® 488 goat-anti-rabbit IgG (1/2000). Cells were fixed with 4% paraformaldehyde and permeabilised with 90% methanol. Isotype control antibody was Rabbit monoclonal IgG (black). The blue line shows cells without incubation with primary antibody and secondary antibody.



Western blot - Anti-HNRNPA0 antibody [EP16085] (ab197023)

All lanes: Anti-HNRNPA0 antibody [EP16085] (ab197023) at 1/10000 dilution

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

Lane 2: Jurkat (Human T cell leukemia cells from peripheral blood)

Lane 3: A549 (Human lung carcinoma) lysate

Lysates/proteins at 20 µg per lane.

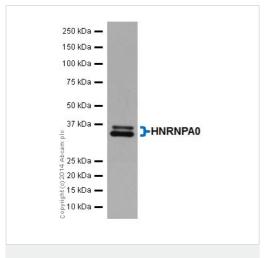
Secondary

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

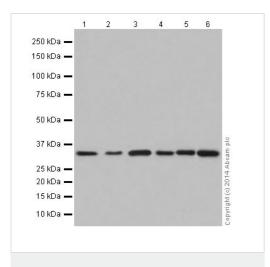
Predicted band size: 31 kDa Observed band size: 32,34 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot - Anti-HNRNPA0 antibody [EP16085] (ab197023)



Western blot - Anti-HNRNPA0 antibody [EP16085] (ab197023)

Anti-HNRNPA0 antibody [EP16085] (ab197023) at 1/1000 dilution + 293 (Human embryonic kidney) lysate at 20 µg

Secondary

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

Predicted band size: 31 kDa Observed band size: 32,34 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

All lanes: Anti-HNRNPA0 antibody [EP16085] (ab197023) at 1/1000 dilution

Lane 1 : Mouse brain lysate

Lane 2: Rat brain lysate

Lane 3: C6 (Rat glial tumor cells) lysate

Lane 4: RAW 264.7 (Mouse macrophage cells transformed with

Abelson murine leukemia virus) lysate

Lane 5: PC-12 (Rat adrenal gland pheochromocytoma) lysate

Lane 6: NIH/3T3 (Mouse embyro fibroblast cells) lysate

Lysates/proteins at 10 µg per lane.

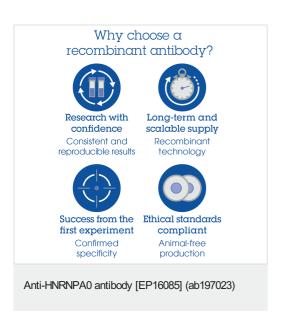
Secondary

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

Predicted band size: 31 kDa **Observed band size:** 32 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.



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