abcam

Product datasheet

Anti-SF3B3 antibody [EPR18441] ab209403

Recombinant RabMAb

1 References 13 Images

Overview

Product name Anti-SF3B3 antibody [EPR18441]

Description Rabbit monoclonal [EPR18441] to SF3B3

Host species Rabbit

Tested applications Suitable for: WB, IHC-P, ICC/IF, IP Species reactivity Reacts with: Mouse, Rat, Human

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

Positive control WB: HeLa, Jurkat, C6, RAW 264.7, PC-12 and NIH/3T3 whole cell lysates; Human fetal liver and

> fetal heart lysates; Mouse brain, kidney and spleen lysates; Rat kidney and spleen lysates. IHC-P: Human colon, Human breast cancer (ER+) and Human breast cancer (ER-) tissues; Mouse cerebral cortex tissue; Rat kidney tissue. ICC/IF: C6 and RAW 264.7 cells. IP: HeLa and C6 cell

lysates.

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 EPR18441

Isotype ΙgG

Applications

The Abpromise guarantee

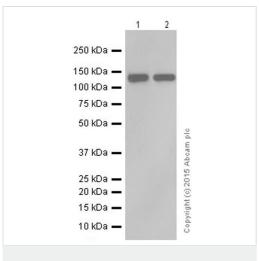
Our <u>Abpromise guarantee</u> covers the use of ab209403 in the following tested applications.

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

Application	Abreviews	Notes
WB		1/2000. Detects a band of approximately 136 kDa (predicted molecular weight: 136 kDa).
IHC-P		1/250. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
ICC/IF		1/500.
IP		1/60.

Target		
Function	Subunit of the splicing factor SF3B required for 'A' complex assembly formed by the stable binding of U2 snRNP to the branchpoint sequence (BPS) in pre-mRNA. Sequence independent binding of SF3A/SF3B complex upstream of the branch site is essential, it may anchor U2 snRNP to the pre-mRNA. May also be involved in the assembly of the 'E' complex. Belongs also to the minor U12-dependent spliceosome, which is involved in the splicing of rare class of nuclear pre-mRNA intron.	
Sequence similarities	Belongs to the RSE1 family.	
Cellular localization	Nucleus.	

Images



Western blot - Anti-SF3B3 antibody [EPR18441] (ab209403)

All lanes : Anti-SF3B3 antibody [EPR18441] (ab209403) at 1/5000 dilution

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

Lane 2: Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate

Lysates/proteins at 20 µg per lane.

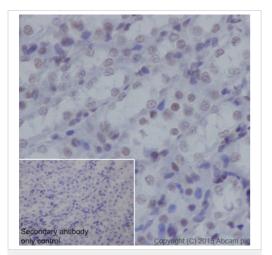
Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/100000 dilution

Predicted band size: 136 kDa **Observed band size:** 136 kDa

Exposure time: 8 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-SF3B3 antibody
[EPR18441] (ab209403)

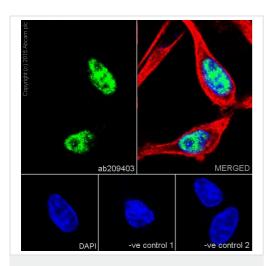
Immunohistochemical analysis of paraffin-embedded Rat kidney tissue labeling SF3B3 with ab209403 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Nucleus staining on rat kidney 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-SF3B3 antibody [EPR18441] (ab209403)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized C6 (Rat glial tumor cell line) cells labeling SF3B3 with ab209403 at 1/500 dilution, followed by Goat Anti-Rabbit lgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green).

Confocal image showing nuclear staining on C6 cell line.

The nuclear counterstain is DAPI (blue).

Tubulin is detected with Anti-alpha Tubulin antibody - Loading Control (<u>ab7291</u>) at 1/1000 dilution and Goat Anti-Mouse IgG (AlexaFluor®594) preadsorbed (<u>ab150120</u>) at 1/1000 dilution (red).

The negative controls are as follows:-

-ve control 1: ab209403 at 1/500 dilution followed by $\underline{ab150120}$ at 1/1000 dilution.

-ve control 2: <u>ab7291</u> at 1/1000 dilution followed by <u>ab150077</u> at 1/1000 dilution.

1 2 3

250 kDa —
150 kDa —
100 kDa —
75 kDa —
250 kDa —

Immunoprecipitation - Anti-SF3B3 antibody [EPR18441] (ab209403)

SF3B3 was immunoprecipitated from 1mg / 300μ l of HeLa (Human epithelial cell line from cervix adenocarcinoma) whole cell lysate with ab209403 at 1/60 dilution (5μ g/ 1mg of lysate).

Western blot was performed from the immunoprecipitate using ab209403 at 1/1000 dilution.

VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>), was used for detection at 1/10000 dilution.

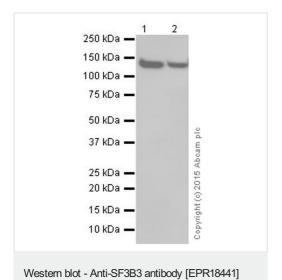
Lane 1: HeLa whole cell lysate, 10ug (Input).

Lane 2: ab209403 IP in HeLa whole cell lysate.

Lane 3: Rabbit lgG,monoclonal - Isotype Control (<u>ab172730</u>) instead of ab209403 in HeLa whole cell lysate.

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

Exposure time: 1 second.



(ab209403)

All lanes: Anti-SF3B3 antibody [EPR18441] (ab209403) at 1/2000 dilution

Lane 1: Human fetal liver lysate Lane 2: Human fetal heart lysate

Lysates/proteins at 10 µg per lane.

Secondary

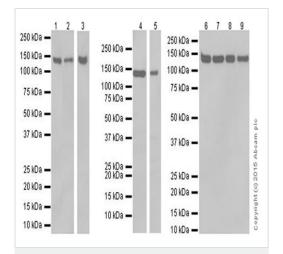
All lanes: Goat Anti-Rabbit IgG Peroxidase Conjugate, specific to the non-reduced form of IgG at 1/10000 dilution

Predicted band size: 136 kDa

Observed band size: 136 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot - Anti-SF3B3 antibody [EPR18441]

(ab209403)

All lanes: Anti-SF3B3 antibody [EPR18441] (ab209403) at 1/2000 dilution

Lane 1: Mouse brain lysate

Lane 2: Mouse kidney lysate

Lane 3: Mouse spleen lysate

Lane 4: Rat kidney lysate

Lane 5: Rat spleen lysate

Lane 6: C6 (Rat glial tumor cell line) whole cell lysate

Lane 7: RAW 264.7 (Mouse macrophage cell line transformed

with Abelson murine leukemia virus) whole cell lysate

Lane 8: PC-12 (Rat adrenal gland pheochromocytoma cell line)

whole cell lysate

Secondary

Lane 9: NIH/3T3 (Mouse embryonic fibroblast cell line) whole cell

lysate

Lysates/proteins at 10 µg per lane.

All lanes: Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at

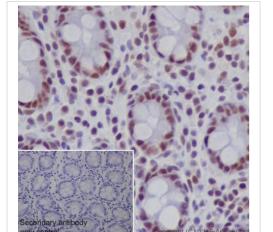
1/100000 dilution

Predicted band size: 136 kDa **Observed band size:** 136 kDa

Blocking/Dilution buffer: 5% NFDM/TBST.

Exposure time: Lane 1/2/4: 3 minutes; Lane 3/5: 30 seconds;

Lane 6/7/8/9: 8 seconds.



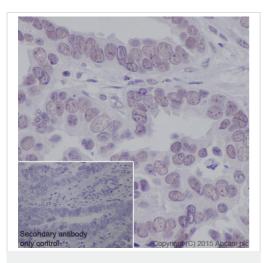
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-SF3B3 antibody
[EPR18441] (ab209403)

Immunohistochemical analysis of paraffin-embedded Human colon tissue labeling SF3B3 with ab209403 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Nucleus staining on Human colon 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.



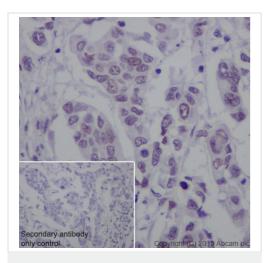
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-SF3B3 antibody
[EPR18441] (ab209403)

Immunohistochemical analysis of paraffin-embedded Human breast cancer (ER+) tissue labeling SF3B3 with ab209403 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Nucleus staining on ER+ breast cancer 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.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-SF3B3 antibody
[EPR18441] (ab209403)

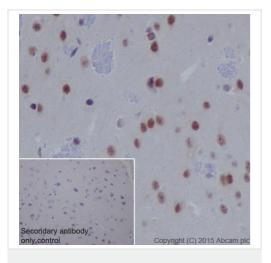
Immunohistochemical analysis of paraffin-embedded Human breast cancer (ER-) tissue labeling SF3B3 with ab209403 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Weak nucleus staining on ER- breast cancer 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-SF3B3 antibody
[EPR18441] (ab209403)

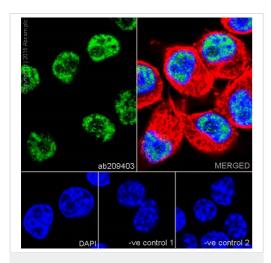
Immunohistochemical analysis of paraffin-embedded mouse cerebral cortex tissue labeling SF3B3 with ab209403 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Nucleus staining on mouse cerebral cortex 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-SF3B3 antibody [EPR18441] (ab209403)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized RAW 264.7 (Mouse macrophage cell line transformed with Abelson murine leukemia virus) cells labeling SF3B3 with ab209403 at 1/500 dilution, followed by Goat Anti-Rabbit IgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green).

Confocal image showing nuclear staining on RAW 264.7 cell line.

The nuclear counterstain is DAPI (blue).

Tubulin is detected with Anti-alpha Tubulin antibody- Loading Control (<u>ab7291</u>) at 1/1000 dilution and Goat Anti-Mouse lgG(AlexaFluor®594) preadsorbed (<u>ab150120</u>) at 1/1000 dilution (red).

The negative controls are as follows:-

-ve control 1: ab209403 at 1/500 dilution followed by $\underline{ab150120}$ at 1/1000 dilution.

-ve control 2: $\underline{ab7291}$ at 1/1000 dilution followed by $\underline{ab150077}$ at 1/1000 dilution.

1 2 3

250 kDa —

150 kDa —

100 kDa —

75 kDa —

50 kDa —

250 kDa —

251 kDa —

252 kDa —

253 kDa —

254 kDa —

255 kDa —

256 kDa —

257 kDa —

258 kD

Immunoprecipitation - Anti-SF3B3 antibody [EPR18441] (ab209403)

SF3B3 was immunoprecipitated from 1mg / 300μ l of C6 (Rat glial tumor cell line) whole cell lysate with ab209403 at 1/60 dilution (5 μ g / 1mg of lysate).

Western blot was performed from the immunoprecipitate using ab209403 at 1/1000 dilution.

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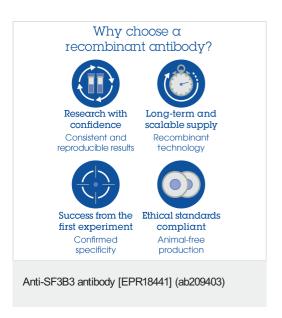
Lane 1: C6 whole cell lysate, 10ug (Input).

Lane 2: ab209403 IP in C6 whole cell lysate.

Lane 3: Rabbit lgG,monoclonal - Isotype Control (<u>ab172730</u>) instead of ab209403 in HeLa whole cell lysate.

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

Exposure time: 1 second.



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