abcam

Product datasheet

Anti-SF3B3 antibody [EPR18440] ab209402

יעלאעבע RabMAb

3 References 画像数 13

製品の概要

製品名 Anti-SF3B3 antibody [EPR18440]

製品の詳細 Rabbit monoclonal [EPR18440] to SF3B3

由来種 Rabbit

アプリケーション 適用あり: IP, Flow Cyt (Intra), WB, IHC-P, ICC/IF

種交差性 交差種: Mouse, Rat, Human

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

ポジティブ・コントロール WB: HeLa, Jurkat, C6 and RAW 264.7 whole cell lysates; Human fetal liver, fetal heart and fetal

> kidney lysates; Mouse brain, heart, kidney and spleen lysates; Rat brain, heart and kidney 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. Flow Cyt

(intra): HeLa cells.

特記事項 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**.

製品の特性

製品の状態 Liquid

保存方法 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.

バッファー pH: 7.2

Preservative: 0.01% Sodium azide

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

精製度 Protein A purified

モノクローナル ポリモノ

クローン名

EPR18440

アイソタイプ

lgG

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab209402の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
IP		1/60.
Flow Cyt (Intra)		1/200.
WB		1/2000. Detects a band of approximately 136 kDa (predicted molecular weight: 136 kDa).
IHC-P		1/4000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
ICC/IF		1/500.

ターゲット情報

機能

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.

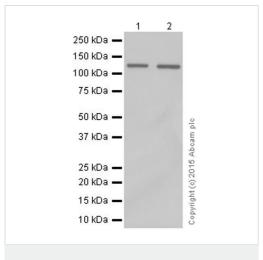
配列類似性

Belongs to the RSE1 family.

細胞内局在

Nucleus.

画像



Western blot - Anti-SF3B3 antibody [EPR18440] (ab209402)

All lanes : Anti-SF3B3 antibody [EPR18440] (ab209402) at 1/20000 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.

1 2 3
250 kDa —
150 kDa —
100 kDa —
75 kDa —
37 kDa —
25 kDa —
20 kDa —
15 kDa —
10 kDa —
10 kDa —
10 kDa —

Western blot - Anti-SF3B3 antibody [EPR18440] (ab209402)

All lanes : Anti-SF3B3 antibody [EPR18440] (ab209402) at 1/2000 dilution

Lane 1 : Human fetal liver lysate

Lane 2 : Human fetal heart lysate

Lane 3 : Human fetal kidney lysate

Lysates/proteins at 10 µg per lane.

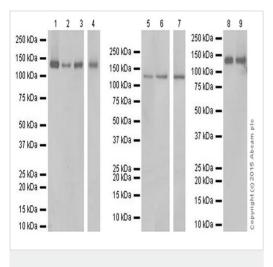
Secondary

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

Blocking/Dilution buffer: 5% NFDM/TBST.

Exposure time: Lane 1/2: 4 seconds; Lane 3: 1 second.



Western blot - Anti-SF3B3 antibody [EPR18440] (ab209402)

All lanes : Anti-SF3B3 antibody [EPR18440] (ab209402) at 1/2000 dilution

Lane 1: Mouse brain lysate

Lane 2: Mouse heart lysate

Lane 3: Mouse kidney lysate

Lane 4: Mouse spleen lysate

Lane 5: Rat brain lysate

Lane 6: Rat kidney lysate

Lane 7: Rat heart lysate

Lane 8: C6 (Rat glial tumor cell line) whole cell lysate

Lane 9: RAW 264.7 (Mouse macrophage cell line transformed

with Abelson murine leukemia virus) whole cell lysate

Lysates/proteins at 10 µg per lane.

Secondary

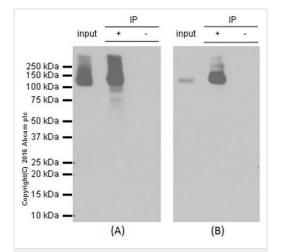
All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/100000 dilution

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

Blocking/Dilution buffer: 5% NFDM/TBST.

Exposure time: Lane 1/2/3: 4 seconds: Lane 4/5/6/8/9: 1

second; Lane 7: 8 seconds.



Immunoprecipitation - Anti-SF3B3 antibody [EPR18440] (ab209402)

ab209402 at 1/60 immunoprecipitating SF3B3 in HeLa (human cervix adenocarcinoma) whole cell lysate observed at 136 KDa (lanes 1 and 2).

Lane 1 (input): HeLa whole cell lysate 10µg

Lane 2 (+): ab209402 + HeLa whole cell lysate.

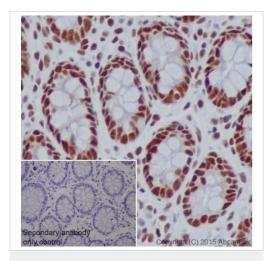
Lane 3 (-): Rabbit monoclonal IgG (ab172730) instead of ab209402 in HeLa whole cell lysate

For western blotting, Panel A: ab209402, 1:500; Panel B: ab209403, 1:500 and VeriBlot for IP Detection Reagent (HRP) (ab131366), was used for detection at 1/10,000 dilution.

Ab209402 was used to capture the target, and the WB was developed with ab209402 (A) <u>ab209403</u> (B).

Blocking buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM /TBST.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-SF3B3 antibody
[EPR18440] (ab209402)

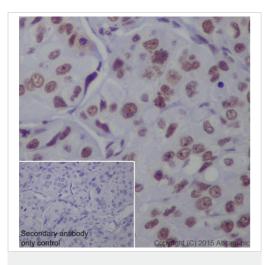
Immunohistochemical analysis of paraffin-embedded Human colon tissue labeling SF3B3 with ab209402 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

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.

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
[EPR18440] (ab209402)

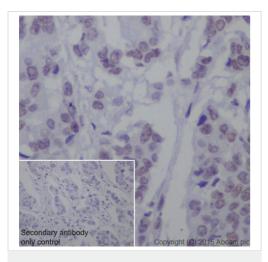
Immunohistochemical analysis of paraffin-embedded Human breast cancer (ER+) tissue labeling SF3B3 with ab209402 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

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.

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
[EPR18440] (ab209402)

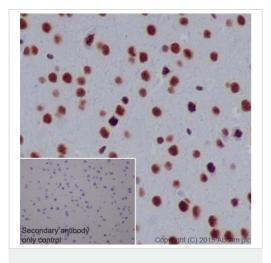
Immunohistochemical analysis of paraffin-embedded Human breast cancer (ER-) tissue labeling SF3B3 with ab209402 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

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.

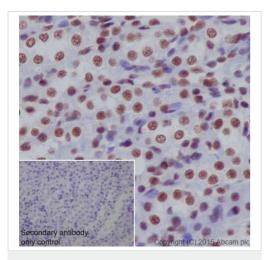


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-SF3B3 antibody
[EPR18440] (ab209402)

Immunohistochemical analysis of paraffin-embedded mouse cerebral cortex tissue labeling SF3B3 with ab209402 at 1/4000 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.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-SF3B3 antibody
[EPR18440] (ab209402)

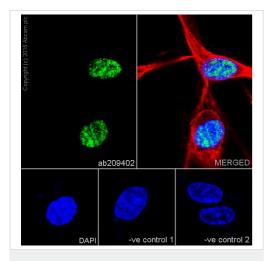
Immunohistochemical analysis of paraffin-embedded rat kidney tissue labeling SF3B3 with ab209402 at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

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 [EPR18440] (ab209402)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized C6 (Rat glial tumor cell line) cells labeling SF3B3 with ab209402 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.

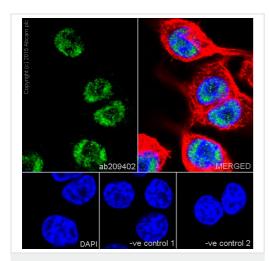
The nuclear counterstain is DAPI (blue).

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

The negative controls are as follows:-

-ve control 1: ab209402 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.



Immunocytochemistry/ Immunofluorescence - Anti-SF3B3 antibody [EPR18440] (ab209402)

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 ab209402 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.

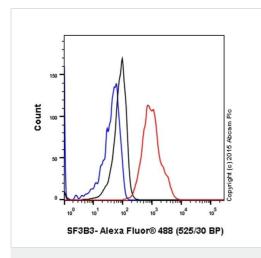
The nuclear counterstain is DAPI (blue).

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

The negative controls are as follows:-

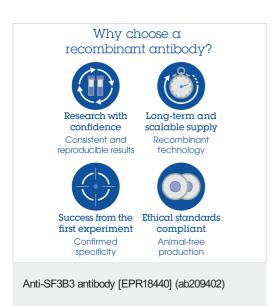
-ve control 1: ab209402 at 1/500 dilution followed by **ab150120** at 1/1000 dilution.

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



Flow Cytometry (Intracellular) - Anti-SF3B3 antibody [EPR18440] (ab209402)

Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed HeLa (Human epithelial cell line from cervix adenocarcinoma) cells labeling SF3B3with ab209402 at 1/200 dilution (red) compared with a Rabbit IgG,monoclonal - Isotype control (ab172730) (black) and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (blue). Goat anti Rabbit IgG (Alexa Fluor® 488) at 1/500 dilution was used as the secondary antibody.



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