abcam

Product datasheet

Anti-Musashi 1 / MSI1 antibody [EPR26106-92] ab305170



Recombinant

RabMAb

13 Images

Overview

Product name Anti-Musashi 1 / MSI1 antibody [EPR26106-92]

Description Rabbit monoclonal [EPR26106-92] to Musashi 1 / Msi1

Host species Rabbit

Specificity Unsuitable for human IHC-P.

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

Unsuitable for: 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: Wild type and knockout HAP1 whole cell lysate; SH-SY5Y, PC-12, Neuro-2a, NIH/3T3 whole

cell lysates; Mouse P0 brainstem, kidney, and heart tissue lysates; mouse and rat embryonic brain tissue lysates; recombinant protein. IHC-P: Mouse testis, mouse and rat cerebellum and mouse and rat small intestine FFPE tissue sections. Flow Cyt (Intra): NIH/3T3 cells. IP: Mouse P0

brainstem tissue lysate.

General notesThis 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: 7.20

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Preservative: 0.01% Sodium azide

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

Purity Protein A purified

Clonality Monoclonal
Clone number EPR26106-92

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab305170 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		1/1000. Detects a band of approximately 39 kDa (predicted molecular weight: 39 kDa).
IHC-P		1/500. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. This antibody is not suitable for human species in IHC-P application.
Flow Cyt (Intra)		1/50.
IP		1/30.

Application notes Is unsuitable for ICC/IF.

Target

Function RNA binding protein that regulates the expression of target mRNAs at the translation level.

Regulates expression of the NOTCH1 antagonist NUMB. Binds RNA containing the sequence 5'-GUUAGUUAGUUAGUU-3' and other sequences containing the pattern 5'-[GA]U(1-3)AGU-3'. May

play a role in the proliferation and maintenance of stem cells in the central nervous system.

Tissue specificity Detected in fetal kidney, brain, liver and lung, and in adult brain and pancreas. Detected in

hepatoma cell lines.

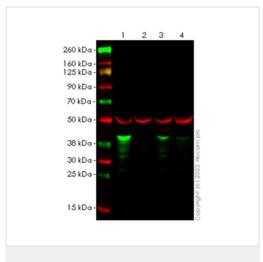
Sequence similarities Belongs to the Musashi family.

Contains 2 RRM (RNA recognition motif) domains.

Domain The first RNA recognition motif binds more strongly to RNA compared to the second one.

Cellular localization Cytoplasm. Nucleus.

Images



Western blot - Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170)

All lanes : Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170) at 1/1000 dilution

Lane 1: Wild-type HAP1 whole cell lysate

Lane 2: Musashi 1 / Msi1 knockout HAP1 whole cell lysate

Lane 3: SH-SY5Y (human neuroblastoma epithelial cell) whole cell

lysate

Lane 4: PC-12 (rat adrenal gland pheochromocytoma cell) whole

cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes: Goat Anti-Rabbit lgG H&L (IRDye® 800CW)
(ab216773) and Goat Anti-Mouse lgG H&L (IRDye® 680RD)
(ab216776) at 1/10000 dilution

Predicted band size: 39 kDa Observed band size: 39 kDa

Blocking / Diluting buffer and concentration: Intercept® (TBS) Blocking Buffer diluted with an equal volume of 0.1% TBS.

Lysates at 20 µg per lane.

Performed under reducing conditions.

False colour image of Western blot: Anti-Musashi 1 / Msi1 antibody [EPR26106-92] (ab305170) staining at 1/1000 dilution, shown in green; Mouse anti-GAPDH antibody [6C5] (ab8245) loading control staining at 1/20000 dilution, shown in red.

In Western blot, ab305170 was shown to bind specifically to Musashi 1 / Msi1. A band was observed at 39 kDa in wild-type HAP1 cell lysates with no signal observed at this size in Musashi 1 / Msi1 knockout cell line (knockout cell lysate). To generate this image, wild-type and Musashi 1 / Msi1 knockout HAP1 cell lysates were analyzed. First, samples were run on an SDS-PAGE gel then transferred onto an immobilon-FL PVDF membrane. Membranes were blocked in in Intercept[®] (TBS) Blocking Buffer diluted with an equal volume of 0.1% TBS before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-

1 2 3

250 kDa = 150 kDa = 150 kDa = 150 kDa = 75 kDa = 25 kDa = 25 kDa = 20 kDa = 15 kDa = 20 kDa = 15 kDa = 20 kDa = 2

Western blot - Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170) T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged.

All lanes : Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170) at 1/1000 dilution

Lane 1: Mouse P0 brainstem tissue lysate

Lane 2: Mouse kidney tissue lysate

Lane 3: Mouse heart tissue lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

Predicted band size: 39 kDa **Observed band size:** 39 kDa

Exposure time: 3 minutes

Blocking / Diluting buffer and concentration: 5% NFDM/TBST Low expression tissues: kidney, heart (PMID: 8660864).

All lanes : Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170) at 1/1000 dilution

Lane 1: Mouse embryonic brain tissue lysate

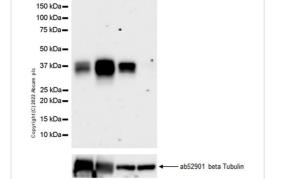
Lane 2: Rat embryonic brain tissue lysate

Lane 3 : Neuro-2a (mouse neuroblastoma neuroblast) whole cell

lysate

Lane 4: NIH/3T3 (mouse embryonic fibroblast) whole cell lysate

Lysates/proteins at 20 µg per lane.



2 3

250 kDa-

Western blot - Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170)

Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/50000 dilution

Predicted band size: 39 kDa **Observed band size:** 39 kDa

Exposure time: 59 seconds

Blocking / Diluting buffer and concentration: 5% NFDM/TBST Negative control: NIH/3T3 (PMID:11359897).

All lanes : Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170) at 1/1000 dilution

Lane 1 : His-tagged mouse Musashi 1 / Msi1 recombinant protein (aa52-251) 10 ng

Lane 2 : His-tagged mouse Musashi 2 / Msi2 recombinant protein (aa1-346) 10 ng

Secondary

All lanes : Goat Anti-Rabbit $\lg G \ H\&L \ (HRP) \ (\underline{ab97051})$ at 1/100000 dilution

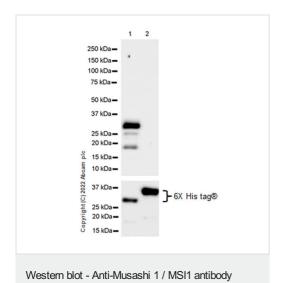
Predicted band size: 39 kDa

Exposure time: 8 seconds

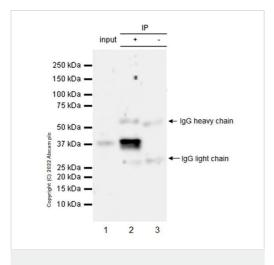
Blocking / Diluting buffer and concentration: 5% NFDM/TBST

Both recombinant proteins were made in-house and expressed from the *E. coli* expression systems.

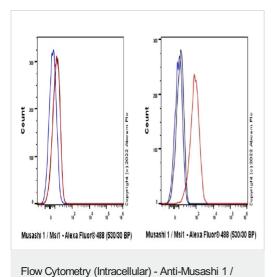
This antibody does not cross-react with mouse Musashi 2 / Msi2.



[EPR26106-92] (ab305170)



Immunoprecipitation - Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170)



MSI1 antibody [EPR26106-92] (ab305170)

Musashi 1 / Msi1 was immunoprecipitated from 0.35 mg mouse P0 brainstem tissue lysate 10 μ g with ab305170 at 1/30 dilution (2ug in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab305170 at 1/1000 dilution. VeriBlot for IP secondary antibody (HRP) (ab131366) was used at 1/5000 dilution.

Lane 1: mouse P0 brainstem tissue lysate 10 µg

Lane 2: ab305170 IP in mouse P0 brainstem tissue lysate

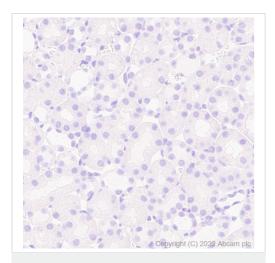
Lane 3: Rabbit monoclonal IgG (<u>ab172730</u>) instead of ab305170 in mouse P0 brainstem tissue lysate

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

Exposure time: 84 seconds.

Observed MW(KDa): 39 kDa

Flow cytometric analysis of 4% paraformaldehyde, fixed 90% methanol permeabilized NIH/3T3 (mouse embryonic fibroblast, Left) / Neuro-2a (mouse neuroblastoma neuroblast, Right) cells labeling Musashi 1 / Msi1 with AB305170 at 1/50 dilution (1 µg) (Red) compared with a Rabbit monoclonal IgG (ab172730) (Black) isotype control and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat Anti-Rabbit IgG (Alexa Fluor[®] 488, ab150081) at 1/2000 dilution was used as the secondary antibody. Negative control: NIH/3T3 (PMID: 11359897).

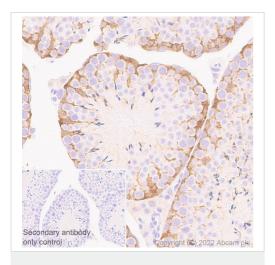


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170)

Immunohistochemical analysis of paraffin-embedded mouse kidney tissue labeling Musashi 1 / Msi1 with ab305170 at 1/500 (0.918 µg/ml) followed by a ready to use Leica DS9800 (Bond ™ Polymer Refine Detection) was used. **Negative control**: No staining on mouse kidney (PMID: 25717188). The section was incubated with ab305170 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use Leica DS9800 (Bond[™] Polymer Refine Detection) kit.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins

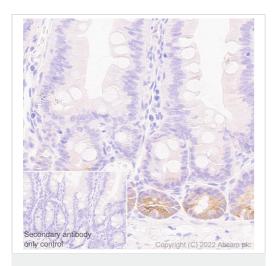


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170)

Immunohistochemical analysis of paraffin-embedded mouse testis tissue labeling Musashi 1 / Msi1 with ab305170 at 1/500 (0.918 µg/ml) followed by a ready to use Leica DS9800 (Bond ™ Polymer Refine Detection) was used. Positive staining on mouse testis (PMID:11804968). The section was incubated with ab305170 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use Leica DS9800 (Bond[™] Polymer Refine Detection) kit.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170)

Immunohistochemical analysis of paraffin-embedded rat small intestine tissue labeling Musashi 1 / Msi1 with ab305170 at 1/500 (0.918 µg/ml) followed by a ready to use Leica DS9800 (Bond [™] Polymer Refine Detection) was used. Positive staining on intestinal crypts of rat (PMID:12558601). The section was incubated with ab305170 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use Leica DS9800 (Bond[™] Polymer Refine Detection) kit.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins

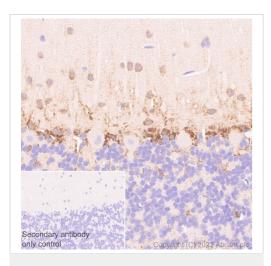


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170)

Immunohistochemical analysis of paraffin-embedded mouse small intestin tissue labeling Musashi 1 / Msi1 with ab305170 at 1/500 (0.918 µg/ml) followed by a ready to use Leica DS9800 (Bond ™ Polymer Refine Detection) was used. Positive staining on intestinal crypts of mouse (PMID:12558601). The section was incubated with ab305170 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use Leica DS9800 (Bond[™] Polymer Refine Detection) kit.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins

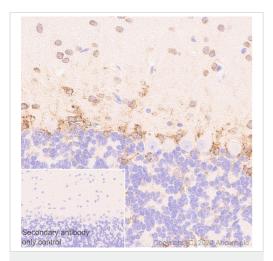


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170)

Immunohistochemical analysis of paraffin-embedded rat cerebellum tissue labeling Musashi 1 / Msi1 with ab305170 at 1/500 (0.918 µg/ml) followed by a ready to use Leica DS9800 (Bond ™ Polymer Refine Detection) was used. Positive staining on rat cerebellum (PMID: 11588182). The section was incubated with ab305170 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use Leica DS9800 (Bond[™] Polymer Refine Detection) kit.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Musashi 1 / MSI1 antibody [EPR26106-92] (ab305170)

Immunohistochemical analysis of paraffin-embedded mouse cerebellum tissue labeling Musashi 1 / Msi1 with ab305170 at 1/500 (0.918 µg/ml) followed by a ready to use Leica DS9800 (Bond[™] Polymer Refine Detection) was used. Positive staining on mouse cerebellum (PMID: 11588182). The section was incubated with ab305170 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins

use Leica DS9800 (Bond[™] Polymer Refine Detection) kit.



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