abcam

Product datasheet

Anti-MSI2 antibody [EP1305Y] ab76148



★★★★★ 1 Abreviews 45 References 画像数 12

製品の概要

製品名 Anti-MSI2 antibody [EP1305Y]

製品の詳細 Rabbit monoclonal [EP1305Y] to MSI2

由来種 Rabbit

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

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.

エピトープ Based on the immunogen sequence for this antibody, it is not predicted to detect the shorter

isoforms of MSI2.

ポジティブ・コントロール WB: HeLa, A549 MCF7, HAP1, Rat brain, Human brain, A549, SW480 and T47D cell lysates

IHC-P: Human placenta, Human bladder carcinoma Tissue IP: T-47D cell lysate. ICC/IF: PC12

and MCF7 cells Flow Cyt (intra): T-47D and 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.

Avoid freeze / thaw cycle.

バッファー pH: 7.20

Preservative: 0.01% Sodium azide

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

精製度 Protein A purified

ポリモノ モノクローナル

クローン名 EP1305Y

アイソタイプ IgG

アプリケーション

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab76148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
Flow Cyt (Intra)		1/100. ab172730 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody. For unpurified use at 1/30
IP		1/50.
ICC/IF		1/100.
WB		1/1000 - 1/2000. Detects a band of approximately 35 kDa (predicted molecular weight: 35 kDa).
IHC-P	**** (1)	1/500. Perform heat mediated antigen retrieval via the pressure cooker method before commencing with IHC staining protocol. The mouse and rat recommendation is based on the WB results. We do not guarantee IHC-P for mouse and rat. See IHC antigen retrieval protocols. For unpurified use at 1/100 - 1/250.

ターゲット情報

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

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

組織特異性 Ubiquitous; detected at low levels.

関連疾患 Note=Chromosomal aberrations involving MSI2 may contribute to disease progression in chronic

myeloid leukemia. Translocation t(7;17)(p15;q23) with HOXA9; translocation t(7;17)(q32-34;q23).

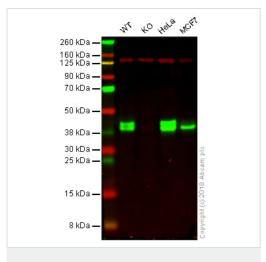
配列類似性 Belongs to the Musashi family.

Contains 2 RRM (RNA recognition motif) domains.

翻訳後修飾 Phosphorylated.

細胞内局在 Cytoplasm. Associated with polysomes.

画像



Western blot - Anti-MSI2 antibody [EP1305Y] (ab76148)

All lanes : Anti-MSI2 antibody [EP1305Y] (ab76148) at 1/1000 dilution

Lane 1: Wild-type HAP1 whole cell lysate

Lane 2: MSI2 knockout HAP1 whole cell lysate

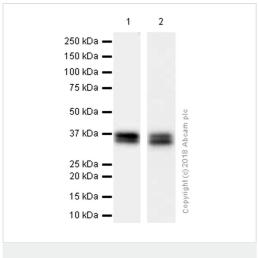
Lane 3 : HeLa whole cell lysate
Lane 4 : MCF7 whole cell lysate

Lysates/proteins at 20 µg per lane.

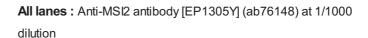
Predicted band size: 35 kDa

Lanes 1 - 4: Merged signal (red and green). Green - ab76148 observed at 40 kDa. Red - loading control, **ab18058**, observed at 130 kDa.

ab76148 was shown to specifically react with MSI2 in wild-type HAP1 cells as signal was lost in MSI2 knockout cells. Wild-type and MSI2 knockout samples were subjected to SDS-PAGE. Ab76148 and ab18058 (Mouse anti-Vinculin loading control) were incubated overnight at 4°C at 1/1000 dilution and 1/20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preabsorbed ab216773 and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed ab216776 secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-MSI2 antibody [EP1305Y] (ab76148)



Lane 1: Mouse brain lysates

Lane 2: Rat brain lysates

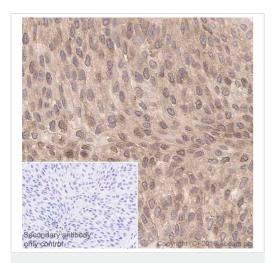
Lysates/proteins at 15 µg per lane.

Secondary

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

dilution

Predicted band size: 35 kDa
Observed band size: 35 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-MSI2 antibody
[EP1305Y] (ab76148)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Human bladder carcinoma tissue sections labeling MSI2 with purified ab76148 at 1:500 dilution (2.14 µg/ml). Heat mediated antigen retrieval was performed using ab93684 (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use)was used as the secondary antibody. Negative control:PBS instead of the primary antibody. Hematoxylin was used as a counterstain.



Western blot - Anti-MSI2 antibody [EP1305Y] (ab76148)

All lanes : Anti-MSI2 antibody [EP1305Y] (ab76148) at 1/1000 dilution

Lane 1 : T-47D (Human ductal breast epithelial tumor epithelial cell) whole cell lysates

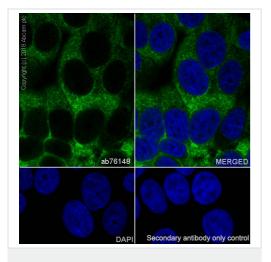
Lane 2: A549 (Human lung carcinoma epithelial cell) whole cell lysates

Lysates/proteins at 15 µg per lane.

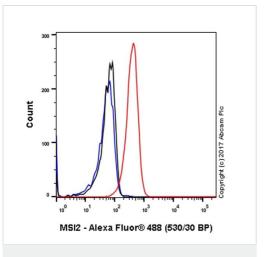
Secondary

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

Predicted band size: 35 kDa Observed band size: 35 kDa



Immunocytochemistry/ Immunofluorescence - Anti-MSI2 antibody [EP1305Y] (ab76148) Immunocytochemistry/ Immunofluorescence analysis of MCF7 (Human breast adenocarcinoma epithelial cell) cells labeling MSI2 with purified ab76148 at 1:100 dilution (10 μ g/ml). Cells were fixed in 4% Paraformaldehyde and permeabilized with 0.1% tritonX-100. Cells were counterstained with None. Goat anti rabbit lgG (Alexa Fluor® 488, **ab150077**) was used as the secondary antibody at 1:1000 (2 μ g/ml) dilution. DAPI nuclear counterstain. PBS instead of the primary antibody was used as the secondary antibody only control.



Flow Cytometry (Intracellular) - Anti-MSI2 antibody [EP1305Y] (ab76148)

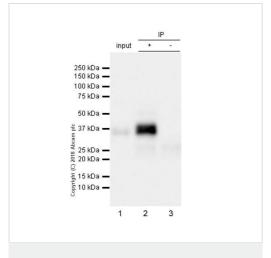
Intracellular Flow Cytometry analysis of T-47D (Human ductal breast epithelial tumor epithelial cell) cells labeling MSI2 with purified ab76148 at 1/100 dilution (10 μ g/ml) (red). Cells were fixed with 4% Paraformaldehyde. A Goat anti rabbit lgG (Alexa Fluor® 488, ab150077) secondary antibody was used at 1/2000. Isotype control - Rabbit monoclonal lgG (Black). Unlabeled control - Cell without incubation with primary antibody and secondary antibody (Blue).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-MSI2 antibody
[EP1305Y] (ab76148)

Immunohistochemical analysis of paraffin-embedded human placenta with ab76148 at 1/100-1/250 dilution.

Heat mediated antigen retrieval was performed via the pressure cooker method before commencing with IHC staining protocol.



Immunoprecipitation - Anti-MSI2 antibody [EP1305Y] (ab76148)

All lanes : Anti-MSI2 antibody [EP1305Y] (ab76148) at 1/1000 dilution

Lane 1 : T-47D (Human ductal breast epithelial tumor epithelial cell) whole cell lysate at 10 µg with 5% NFDM/TBST

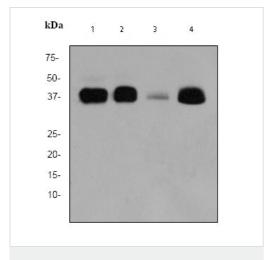
Lane 2: T-47D whole cell lysate. ab76148 use as the capture antibody at 1:50 dilution. with 5% NFDM/TBST

Lane 3 : T-47D whole cell lysate. <u>ab172730</u> use as the capture antibody at 1:50 dilution. with 5% NFDM/TBST

Secondary

All lanes : VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) at 1/5000 dilution (VeriBlot for IP secondary antibody (HRP))

Observed band size: 35 kDa



Western blot - Anti-MSI2 antibody [EP1305Y] (ab76148)

Immunocytochemistry/ Immunofluorescence - Anti-MSI2 antibody [EP1305Y] (ab76148)

Flow Cytometry (Intracellular) - Anti-MSI2 antibody [EP1305Y] (ab76148)

Exposure time: 5 seconds

All lanes : Anti-MSI2 antibody [EP1305Y] (ab76148) at 1/2000 dilution

Lane 1: Rat brain cell lysates

Lane 2: Human brain cell lysates

Lane 3: SW480 cell lysates

Lane 4: T47D cell lysates

Lysates/proteins at 10 µg per lane.

Secondary

All lanes: HRP labelled goat anti-rabbit at 1/2000 dilution

Predicted band size: 35 kDa Observed band size: 35 kDa

ICC/IF image of ab76148 stained PC12 cells. The cells were 4% formaldehyde fixed (10 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab76148, 1 μ g/ml) overnight at +4°C. The secondary antibody (green) was Alexa Fluor® 488 goat anti-rabbit lgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43 μ M.

Overlay histogram showing HeLa cells stained with ab76148 (red line). The cells were fixed with 4% paraformaldehyde (10 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab76148, 1/100 dilution) for 30 min at 22°C. The secondary antibody used was DyLight® 488 goat anti-rabbit IgG (H+L) (ab96899) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit IgG (monclonal) (1µg/1x10⁶ cells) used under the same conditions. Acquisition of >5,000 events was performed. This antibody gave a positive result in HeLa cells fixed with 80% methanol (5 min)/permeabilized in 0.1% PBS-Tween for 20 min used under the same conditions.



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