abcam

Product datasheet

Anti-MYSM1 antibody [EPR18657] ab193081



ייבער RabMAb

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製品の概要

製品名 Anti-MYSM1 antibody [EPR18657]

製品の詳細 Rabbit monoclonal [EPR18657] to MYSM1

由来種 Rabbit

アプリケーション **適用あり:** WB, IP

種交差性 交差種: Mouse, Rat, Human

免疫原 Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

ポジティブ・コントロール WB: HeLa, HEK-293, HepG2, NCCIT, K562, RAW 264.7, PC-12, NIH/3T3 and mESC whole cell

lysates. Wild-type HEK-293T cell lysate. IP: HeLa whole cell lysate.

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, 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 EPR18657

アイソタイプ ΙgG

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ab193081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
WB		1/1000. Detects a band of approximately 66, 95 kDa (predicted molecular weight: 66, 95 kDa).
IP		1/100.

ターゲット情報

機能 Metalloprotease that specifically deubiquitinates monoubiquitinated histone H2A, a specific tag

for epigenetic transcriptional repression, thereby acting as a coactivator. Preferentially deubiquitinates monoubiquitinated H2A in hyperacetylated nucleosomes. Deubiquitination of histone H2A leads to facilitate the phosphorylation and dissociation of histone H1 from the nucleosome. Acts as a coactivator by participating in the initiation and elongation steps of

androgen receptor (AR)-induced gene activation.

配列類似性 Belongs to the peptidase M67A family. MYSM1 subfamily.

Contains 1 MPN (JAB/Mov34) domain.

Contains 1 SANT domain.
Contains 1 SWIRM domain.

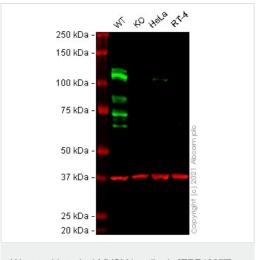
ドメイン Binds double-stranded DNA via the SANT domain. The SWIRM domain does not bind double-

stranded DNA.

翻訳後修飾 Phosphorylated upon DNA damage, probably by ATM or ATR.

細胞内局在 Nucleus.

画像



Western blot - Anti-MYSM1 antibody [EPR18657] (ab193081)

All lanes : Anti-MYSM1 antibody [EPR18657] (ab193081) at 1/1000 dilution

Lane 1: Wild-type HEK-293T cell lysate

Lane 2: MYSM1 knockout HEK-293T cell lysate

Lane 3 : HeLa cell lysate

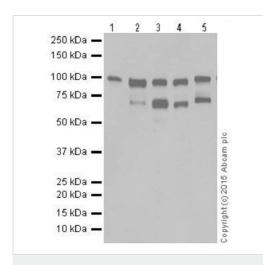
Lane 4 : RT-4 cell lysate

Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 66, 95 kDa **Observed band size:** 110 kDa

False colour image of Western blot: Anti-MYSM1 antibody [EPR18657] 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, ab193081 was shown to bind specifically to MYSM1. A band was observed at 110 kDa in wild-type HEK-293T cell lysates with no signal observed at this size in MYSM1 knockout cell line ab267297 (knockout cell lysate ab257548). To generate this image, wild-type and MYSM1 knockout HEK-293T cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3 % milk in TBS-0.1 % Tween® 20 (TBS-T) before incubation with primary antibodies overnight at 4°C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L (IRDye® 800CW) preabsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed (ab216776) at 1/20000 dilution.



Western blot - Anti-MYSM1 antibody [EPR18657] (ab193081)

All lanes : Anti-MYSM1 antibody [EPR18657] (ab193081) at 1/1000 dilution

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

Lane 2: HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate

Lane 3 : Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate

Lane 4 : NCCIT (Human pluripotent embryonic carcinoma cell line) whole cell lysate

Lane 5: K562 (Human chronic myelogenous leukemia cell line from bone marrow) whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

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

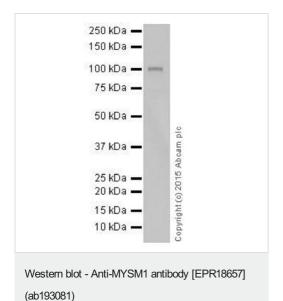
Developed using the ECL technique.

Predicted band size: 66, 95 kDa **Observed band size:** 66,95 kDa

Exposure time: 1 minute

Blocking/Dilution buffer: 5% NFDM/TBST.

Two bands detected represent two isoforms.



Anti-MYSM1 antibody [EPR18657] (ab193081) at 1/1000 dilution + mESC (Mouse embryonic stem cell line) whole cell lysate at 20 μg

Secondary

Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Developed using the ECL technique.

Predicted band size: 66, 95 kDa Observed band size: 95 kDa

Exposure time: 8 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

250 kDa -150 kDa -100 kDa -75 kDa -50 kDa -Copyright (c) 2015 Abcam plo 37 kDa -25 kDa -20 kDa -15 kDa -10 kDa -

Western blot - Anti-MYSM1 antibody [EPR18657]

(ab193081)

All lanes: Anti-MYSM1 antibody [EPR18657] (ab193081) at 1/1000 dilution

Lane 1: RAW 264.7 (Mouse macrophage cell line transformed with Abelson murine leukemia virus) whole cell lysate

Lane 2: PC-12 (Rat adrenal gland pheochromocytoma cell line) whole cell lysate

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

Lysates/proteins at 10 µg per lane.

Secondary

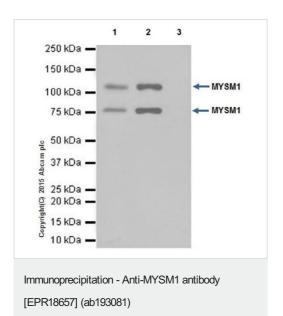
All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Developed using the ECL technique.

Predicted band size: 66, 95 kDa Observed band size: 95 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.



MYSM1 was immunoprecipitated from 1mg of HeLa (Human epithelial cell line from cervix adenocarcinoma) whole cell lysate with ab193081 at 1/100 dilution. Western blot was performed from the immunoprecipitate using ab193081 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (ab131366), was used for detection at 1/1000 dilution.

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

Lane 2: ab193081 IP in HeLa whole cell lysate.

Lane 3: Rabbit monoclonal $\lg G$ ($\underline{ab172730}$) instead of ab193081 in HeLa whole cell lysate.

Predicted band size: 66kDa, 95kDa.

Observed band size: 66kDa, 95kDa.

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

Exposure time: 3 minutes.

Two bands detected represent two isoforms.



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