# abcam

### Product datasheet

## Anti-mSin3A antibody - ChIP Grade ab3479

★★★★★ 10 Abreviews 44 References 画像数 7

#### 製品の概要

製品名 Anti-mSin3A antibody - ChIP Grade

製品の詳細 Rabbit polyclonal to mSin3A - ChIP Grade

由来種 Rabbit

アプリケーション 適用あり: ChIP, ICC/IF, WB, IHC-P

種交差性 交差種: Mouse, Human

免疫原 Synthetic peptide corresponding to Mouse mSin3A aa 1-100.

■ Run BLAST with EXPASY ■ Run BLAST with S NCBI

特記事項

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

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 or -

80°C. Avoid freeze / thaw cycle.

パッファー Preservative: 0.05% Sodium azide

Constituents: 0.1% BSA, PBS

精製度 Immunogen affinity purified

ポリ/モノ ポリクローナル

アイソタイプ lgG

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ab3479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。

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アプリケーション	Abreviews	特記事項
ChIP	<b>★★★★</b> <u>(2)</u>	Use at an assay dependent concentration.
ICC/IF	<b>★★★★</b> <u>(1)</u>	1/50 - 1/500.
WB	★★★★☆ (5)	Use a concentration of 0.5 µg/ml. Detects a band of approximately 150 kDa (predicted molecular weight: 145 kDa).
IHC-P	<b>★★★★☆ (1)</b>	Use a concentration of 2 µg/ml.

#### ターゲット情報

機能 Acts as a transcriptional repressor. Interacts with MXI1 to repress MYC responsive genes and

antagonize MYC oncogenic activities. Also interacts with MAD-MAX heterodimers by binding to  $% \left( 1\right) =\left( 1\right) \left( 1\right) \left($ 

 $\mbox{MAD}.$  The heterodimer then represses transcription by tethering SIN3A to DNA. Acts as a

corepressor for REST.

**配列類似性** Contains 3 PAH (paired amphipathic helix) domains.

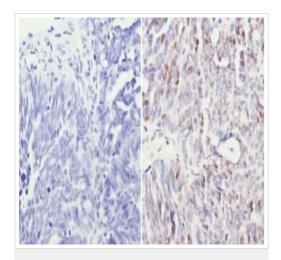
翻訳後修飾 SUMO1 sumoylated by TOPORS. Probably desumoylated by SENP2. 細胞内局在 Nucleus. Nucleus > nucleolus. Recruited to the nucleolus by SAP30L.

#### 画像



Western blot - Anti-mSin3A antibody - ChIP Grade (ab3479)

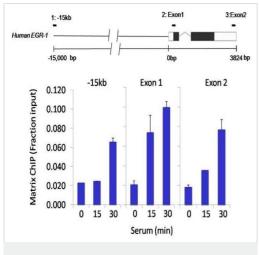
Western blot of mSin3A on K562 cell extract using ab3479.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-mSin3A antibody - ChIP Grade (ab3479)

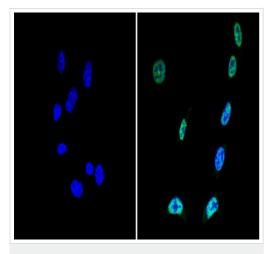
Paraffin-embedded human ovary carcinoma tissue (right panel) stained for mSin3A using ab3479 at 1/200 dilution compared to a negative control without primary antibody (left panel) in immunohistochemical analysis, followed by HRP-conjugated secondary antibody. Detection: DAB staining.

Antigen retrieval was performed using 10mM sodium citrate (pH 6.0), microwaved for 8-15 min.



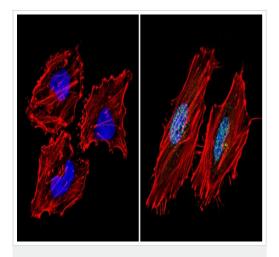
ChIP - Anti-mSin3A antibody - ChIP Grade (ab3479)

ChIP analysis of Sin3A was performed using cross-linked chromatin from 1x10<sup>6</sup> HCT 116 (human colorectal carcinoma cell line) cells treated with serum for 0, 15, and 30 minutes. IP was performed using a multiplex microplate Matrix ChIP assay with 1.0 µl/100 µl well volume of ab3479. Chromatin aliquots from ~1x10<sup>5</sup> cells were used per ChIP pull-down. Quantitative PCR data were done in quadruplicate using 1 µl of eluted DNA in 2 µl SYBR realtime PCR reactions containing primers to amplify -15kb upstream of the Egr1 gene or exon-1 or exon-2 of Egr1. Quantitation of immunoprecipitated chromatin is presented as signal relative to the total amount of input chromatin. A schematic representation of the Egr-1 locus is shown above the data where boxes represent exons (black boxes = translated regions, white boxes = untranslated regions), the zigzag line represents an intron, and the straight line represents upstream sequence. Regions amplified by Egr-1 primers are represented by black bars.



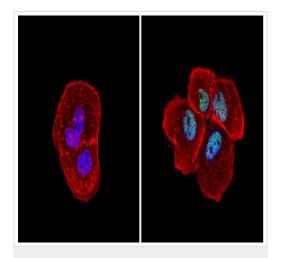
Immunocytochemistry/ Immunofluorescence - AntimSin3A antibody - ChIP Grade (ab3479)

Immunocytochemical immunoflurescence analysis of NIH-3T3 cells labelling mSin3A using ab3479. Formalin-fixed cells were permeablized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were then incubated with ab3479 in 3% BSA-PBS at a dilution of 1:100 overnight in a 4°C high humidity environment. Cells were then washed with PBST and incubated with a green DyLight-conjugated secondary antibody in PBS at room temperature in the dark. Cells were counterstained with DAPI or Hoechst labelling the nuclear DNA blue. The Left Image is a negative control without the prescence of ab3479. Image magnification is 60X.



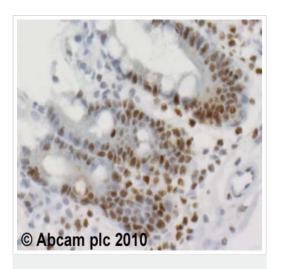
Immunocytochemistry/ Immunofluorescence - AntimSin3A antibody - ChIP Grade (ab3479)

Immunocytochemical immunoflurescence analysis of HeLa cells labelling mSin3A using ab3479. Formalin-fixed cells were permeablized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were then incubated with ab3479 in 3% BSA-PBS at a dilution of 1:200 overnight in a 4°C high humidity environment. Cells were then washed with PBST and incubated with a green DyLight-conjugated secondary antibody in PBS at room temperature in the dark. Cells were counterstained blue with DAPI or Hoechst labelling the nuclear DNA and red against actin using an Alexa Fluor® 554 conjugate. The Left Image is a negative control without the prescence of ab3479. Image magnification is 60X.



Immunocytochemistry/ Immunofluorescence - AntimSin3A antibody - ChIP Grade (ab3479)

Immunocytochemical immunoflurescence analysis of NIH-3T3 cells labelling mSin3A using ab3479. Formalin-fixed cells were permeablized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were then incubated with ab3479 in 3% BSA-PBS at a dilution of 1:100 overnight in a 4°C high humidity environment. Cells were then washed with PBST and incubated with a green DyLight-conjugated secondary antibody in PBS at room temperature in the dark. Cells were counterstained blue with DAPI or Hoechst labelling the nuclear DNA and red against actin using an Alexa Fluor® 554 conjugate. The Left Image is a negative control without the prescence of ab3479. Image magnification is 60X.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-mSin3A antibody - ChIP Grade (ab3479)

ab3479 (2µg/ml) staining mSin3A in human duodenum using an automated system (DAKO Autostainer Plus). Using this protocol there is strong staining of nuclei of epithelial cells.

Sections were rehydrated and antigen retrieved with the Dako 3 in 1 AR buffers citrate pH6.1/ EDTA pH 9.0 in a DAKO PT link. Slides were peroxidase blocked in 3% H2O2 in methanol for 10 mins.

They were then blocked with Dako Protein block for 10 minutes (containing casein 0.25% in PBS) then incubated with primary antibody for 20 min and detected with Dako envision flex amplification kit for 30 minutes. Colorimetric detection was completed with Diaminobenzidine for 5 minutes. Slides were counterstained with Haematoxylin and coverslipped under DePeX. Please note that, for manual staining, optimization of primary antibody concentration and incubation time is recommended.

Signal amplification may be required.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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