# abcam

# Product datasheet

# Anti-NAT10 antibody [EPR18663] ab194297

יולצעבע RabMAb

★★★★★ 2 Abreviews 11 References 画像数 10

#### 製品の概要

製品名 Anti-NAT10 antibody [EPR18663]

製品の詳細 Rabbit monoclonal [EPR18663] to NAT10

由来種 Rabbit

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

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: Human fetal brain and fetal heart lysates; Mouse brain, rat heart and rat spleen lysates. IHC-

P: Human colon, mouse stomach and rat colon tissues. ICC/IF: HeLa and NIH/3T3 cells. IP: HeLa

cell lysate. Flow Cyt (intra): NIH/3T3 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, 40% Glycerol (glycerin, glycerine), 0.05% BSA

精製度 Protein A purified

ポリモノ モノクローナル クローン名 EPR18663

### アプリケーション

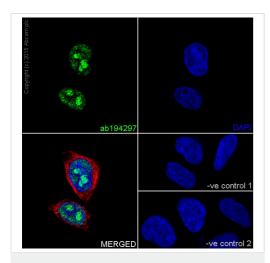
**The Abpromise guarantee Abpromise保証は、**次のテスト済みアプリケーションにおけるab194297の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
Flow Cyt (Intra)		1/600.
WB	**** (1)	1/2000. Detects a band of approximately 116 kDa (predicted molecular weight: 116 kDa).
ICC/IF	**** (1)	1/2000.
IP		1/80.
IHC-P		1/500. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

### ターゲット情報

Has protein acetyltransferase activity in vitro. Can acetylate both histone Histone acetylation may regulate transcription and mitotic chromosome Activates telomerase activity by stimulating the transcription of TERT, ar telomerase function by affecting the balance of telomerase subunit asse localization. Acetylates alpha-tubulin, which may affect microtubule stabi	e de-condensation. and may also regulate sembly, disassembly, and
Belongs to the UPF0202 family.  Contains 1 N-acetyltransferase domain.	
Nucleus > nucleolus. Nucleolar in interphase and redistributes to the per to the midbody during telophase.	erichromosomal layer and
telomerase function by affecting the balance of telomerase subunit asse localization. Acetylates alpha-tubulin, which may affect microtubule stabil Belongs to the UPF0202 family.  Contains 1 N-acetyltransferase domain.  Nucleus > nucleolus. Nucleolar in interphase and redistributes to the per	sembly, disassembly, and bility and cell division.

### 画像



Immunocytochemistry/ Immunofluorescence - Anti-NAT10 antibody [EPR18663] (ab194297)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cells from cervix adenocarcinoma) cells labeling NAT10 with ab194297 at 1/2000 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green).

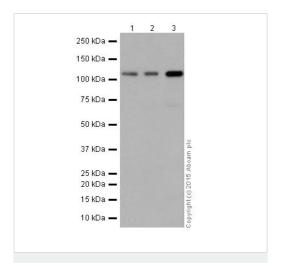
Confocal image showing nuclear staining on HeLa cell line.

The nuclear counterstain is DAPI (blue).

Tubulin is detected with <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution and <u>ab150120</u> (AlexaFluor<sup>®</sup>594 Goat anti-Mouse secondary) at 1/1000 dilution (red).

The negative controls are as follows:-

-ve control 1: ab194297 at 1/2000 dilution followed by <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution. -ve control 2: <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> (Alexa Fluor®488 Goat Anti-Rabbit lgG H&L) at 1/1000 dilution.



Western blot - Anti-NAT10 antibody [EPR18663] (ab194297)

**All lanes :** Anti-NAT10 antibody [EPR18663] (ab194297) at 1/2000 dilution

Lane 1: Mouse brain lysate

Lane 2: Rat heart lysate

Lane 3: Rat spleen lysate

Lysates/proteins at 10 µg per lane.

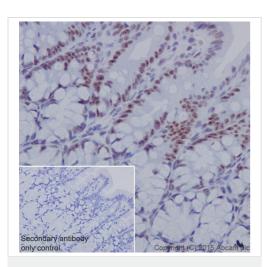
#### **Secondary**

All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/50000

dilution

**Predicted band size:** 116 kDa **Observed band size:** 116 kDa

Exposure time: 30 seconds



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-NAT10 antibody
[EPR18663] (ab194297)

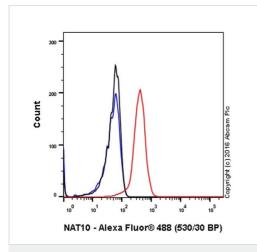
Immunohistochemical analysis of paraffin-embedded Rat colon tissue labeling NAT10 with ab194297 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Nucleus staining on rat colon tissue is observed.

Counter stained with Hematoxylin.

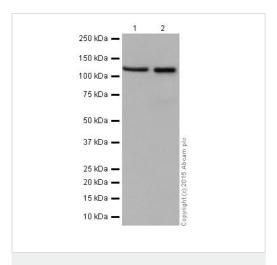
Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Flow Cytometry (Intracellular) - Anti-NAT10 antibody [EPR18663] (ab194297)

Intracellular Flow Cytometry analysis of NIH/3T3 (mouse embryo) cells labelling NAT10 (red) with purified ab194297 at dilution of 1/600. The secondary antibody used was Alexa Fluorr® 488 goatanti-rabbit IgG (1/2000). Cells were fixed with 4% paraformaldehyde and permeabilised with 90% methanol. Isotype control antibody used was Rabbit Monoclonal IgG (black). The blue line shows cells without incubation with primary antibody and secondary antibody.



Western blot - Anti-NAT10 antibody [EPR18663] (ab194297)

**All lanes :** Anti-NAT10 antibody [EPR18663] (ab194297) at 1/2000 dilution

Lane 1: Human fetal brain lysate

Lane 2: Human fetal heart lysate

Lysates/proteins at 10 µg per lane.

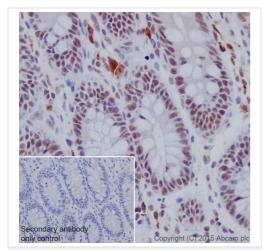
## **Secondary**

**All lanes :** Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/10000 dilution

**Predicted band size:** 116 kDa **Observed band size:** 116 kDa

Exposure time: 30 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-NAT10 antibody
[EPR18663] (ab194297)

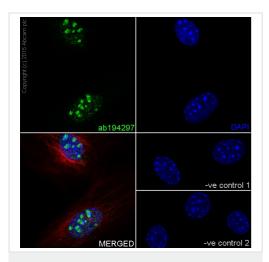
Immunohistochemical analysis of paraffin-embedded Human colon tissue labeling NAT10 with ab194297 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Nucleus staining on Human colon tissue 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-NAT10 antibody [EPR18663] (ab194297)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized NIH/3T3 (Mouse embyro fibroblast cells) cells labeling NAT10 with ab194297 at 1/2000 dilution, followed by Goat anti-rabbit lgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green).

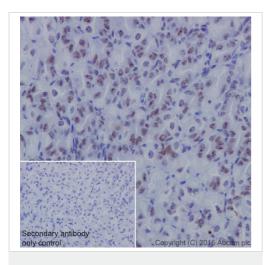
Confocal image showing nuclear staining on NIH/3T3 cell line.

The nuclear counterstain is DAPI (blue).

Tubulin is detected with <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution and <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution (red).

The negative controls are as follows:-

-ve control 1: ab194297 at 1/2000 dilution followed by <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution. -ve control 2: <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> (Alexa Fluor®488 Goat Anti-Rabbit lgG H&L) at 1/1000 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-NAT10 antibody
[EPR18663] (ab194297)

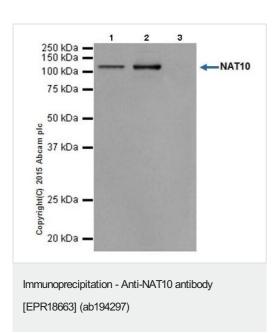
Immunohistochemical analysis of paraffin-embedded Mouse stomach tissue labeling NAT10 with ab194297 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Nucleus staining on mouse stomach tissue 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.



NAT10 was immunoprecipitated from 1mg of HeLa (Human epithelial cells from cervix adenocarcinoma) cell lysate with ab194297 at 1/80 dilution.

Lane 1: HeLa cell lysate 10ug (Input).

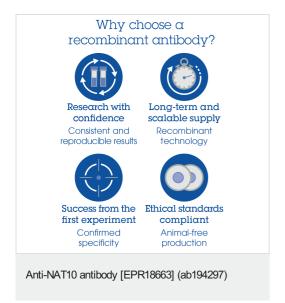
Lane 2: ab194297 IP in HeLa cell lysate.

Lane 3: Rabbit monoclonal IgG (<u>ab172730</u>) instead of ab194297 in HeLa cell lysate.

Western blot was performed from the immunoprecipitate using ab194297 at 1/1000 dilution. Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG, was used as secondary antibody at 1/1500.

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

Exposure time: 3 seconds.



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