# abcam

## Product datasheet

## Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401] ab201015

יועדער RabMAb

★★★★★ 4 Abreviews 157 References 画像数 9

#### 製品の概要

製品名 Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401]

製品の詳細 Rabbit monoclonal [EPR19401] to ERK1 (phospho T202) + ERK2 (phospho T185)

由来種 Rabbit

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

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.

ポジティブ・コントロール WB: Jurkat, treated with 200 ng/ml PMA for 30 minutes whole cell lysate; NIH/3T3, untreated and

> treated with 50 ng/ml PDGF for 40 minutes whole cell lysates; PC-12, untreated and treated with 200 ng/ml NGF for 4 days whole cell lysates. IHC-P: Human breast and glioma tissues. IP: NIH/3T3 treated with 50ng/ml PDGF for 40min whole cell lysate; PC-12 treated with 100ng/ml

NGF for 10min whole cell lysate. ICC/IF:Jurkat

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

精製度 Protein A purified

**ポリ**(モノ モノクローナル **クローン名** EPR19401

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ab201015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。

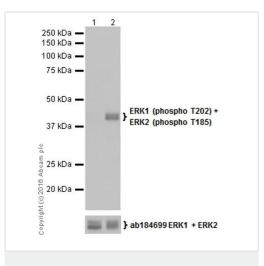
アプリケーション	Abreviews	特記事項
ICC/IF	<b>★★★★ (3)</b>	Use at an assay dependent concentration.
IP		1/30.
IHC-P		1/100. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. IHC is recommended for human only.
WB	**** <u>(1)</u>	1/1000. Detects a band of approximately 44, 42 kDa (predicted molecular weight: 41 kDa).
Dot blot		1/1000.

#### ターゲット情報

細胞内局在

ERK2: Nucleus.

#### 画像



Western blot - Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401] (ab201015)

**All lanes**: Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401] (ab201015) at 1/1000 dilution

**Lane 1 :** Untreated Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate

Lane 2: Jurkat (Human T cell leukemia cell line from peripheral blood) treated with 200 ng/ml PMA for 30 minutes whole cell lysate

Lysates/proteins at 20 µg per lane.

## **Secondary**

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

Predicted band size: 41 kDa

Observed band size: 42,44 kDa

Exposure time: 15 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

### AD201015 DAPI MERGED

### AD201015 DAPI MERGED

### AD201015 DAPI MERGED

\*\*\*CONTROL 1\*\*\*

\*\*\*CONTROL 2\*\*\*

\*\*\*CONTROL 2\*\*

\*

Immunocytochemistry/ Immunofluorescence - Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401] (ab201015)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized Jurkat (Human T cell leukemia cell line from peripheral blood) cells labeling ERK1 (phospho T202) and ERK2 (phospho T185)

ERK1 (phospho T202) + ERK2 (phospho T185) with ab201015 at 1/500 dilution, followed by Goat anti-rabbit lgG (Alexa Fluor<sup>®</sup> 488) (ab150077) secondary antibody at 1/1000 dilution (green).

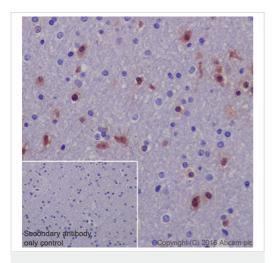
Confocal image showing staining on M phase cells (PMID:26529125). After PMA treatment (200 ng/ml, 30min), the staining was increased, and LP treatment decreased the PMA induced staining.

The nuclear counter stain is DAPI (blue).

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

The negative controls are as follows:

-ve control 1: ab201015 at 1/500 dilution followed by <u>ab150120</u> (Alexa Fluor<sup>®</sup> 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.

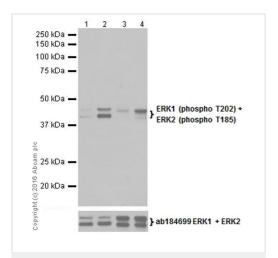


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401] (ab201015)

Immunohistochemical analysis of paraffin-embedded Human glioma tissue labeling ERK2 (phospho T185) with ab201015 at 1/100 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear with weak cytoplasm staining on Human glioma is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is <u>ab97051</u> at 1/500 dilution.

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



Western blot - Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401] (ab201015)

All lanes: Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401] (ab201015) at 1/1000 dilution

**Lane 1 :** Untreated NIH/3T3 (Mouse embryonic fibroblast cell line) whole cell lysate

**Lane 2:** NIH/3T3 (Mouse embryonic fibroblast cell line) treated with 50 ng/ml PDGF for 40 minutes whole cell lysate

Lane 3: Untreated PC-12 (Rat adrenal gland pheochromocytoma cell line) whole cell lysate

**Lane 4**: PC-12 (Rat adrenal gland pheochromocytoma cell line) treated with 200 ng/ml NGF for 4 days whole cell lysate

Lysates/proteins at 20 µg per lane.

## Secondary

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

Predicted band size: 41 kDa

Observed band size: 42.44 kDa

Exposure time: 10 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

The induction conditions refer to PMID:12454035; PMID:17026715;

PMID:22206868.

1 2 3
5ng
1ng
0.1ng
0.01ng

Dot Blot - Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401] (ab201015)

Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401] (ab201015)

Dot blot analysis of ERK2 (phospho T185) labeled with ab201015 at 1/1000 dilution.

**Lane 1:** ERK2 (pT185) phospho peptide: DHTGFLT(p)EYVATR aa179-191 peptide.

**Lane 2:** ERK2 Non-phospho peptide: DHTGFLTEYVATR aa179-191 peptide.

**Lane 3:** ERK2 (pY187) phospho peptide: DHTGFLTEY(p)VATR aa179-191 peptide.

Goat Anti-Rabbit IgG, (H+L),Peroxidase conjugated (<u>ab97051</u>) at 1/100000 dilution was used as secondary antibody.

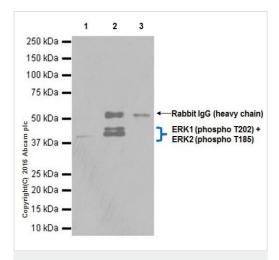
Blocking/Dilution buffer: 5% NFDM/TBST.

Exposure time: 3 minutes.

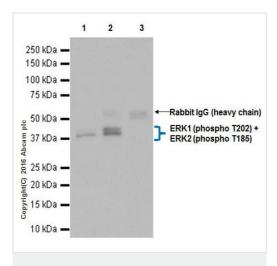
Immunohistochemical analysis of paraffin-embedded Human breast tissue labeling ERK2 (phospho T185) with ab201015 at 1/100 dilution, followed by Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear staining on Human breast is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is <u>ab97051</u> at 1/500 dilution.

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



Immunoprecipitation - Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401] (ab201015)



Immunoprecipitation - Anti-ERK1 (phospho T202) + ERK2 (phospho T185) antibody [EPR19401] (ab201015)

ERK2 (phospho T185) was immunoprecipitated from 0.35 mg of NIH/3T3 (Mouse embryonic fibroblast cell line) treated with 50ng/ml PDGF for 40min whole cell lysate with ab201015 at 1/30 dilution. Western blot was performed from the immunoprecipitate using ab201015 at 1/1000 dilution. VeriBlot for IP Detection Reaction (HRP) (ab131366), was used for detection at 1/1000 dilution.

Lane 1: NIH/3T3 treated with 50ng/ml PDGF for 40min whole cell lysate,  $10\mu g$  (lnput).

Lane 2: ab201015 IP in NIH/3T3 treated with 50ng/ml PDGF for 40min whole cell lysate.

Lane 3: Rabbit lgG,monoclonal [EPR25A]-lsotype
Control (<u>ab172730</u>) instead of ab201015 in NIH/3T3 treated with
50ng/ml PDGF for 40min whole cell lysate.

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

ERK2 (phospho T185) was immunoprecipitated from 0.35 mg of PC-12 (Rat adrenal gland pheochromocytoma cell line) treated with 100ng/ml NGF for 10min whole cell lysate with ab201015 at 1/30 dilution. Western blot was performed from the immunoprecipitate

using ab201015 at 1/1000 dilution. VeriBlot for IP Detection Reaction (HRP) (ab131366), was used for detection at 1/1000 dilution.

Lane 1: PC-12 treated with 100ng/ml NGF for 10min whole cell lysate,  $10\mu g$  (lnput).

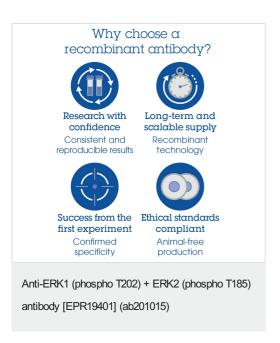
Lane 2: ab201015 IP in PC-12 treated with 100ng/ml NGF for 10min whole cell lysate.

Lane 3: Rabbit lgG,monoclonal [EPR25A]-lsotype
Control (ab172730) instead of ab201015 in PC-12 treated with
100ng/ml NGF for 10min whole cell lysate

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

Exposure time: 3 minutes.

Exposure time: 3 minutes.



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