

Anti-alpha smooth muscle Actin (acetyl E3) + ACTG2 (acetyl E3) antibody [E184] - BSA and Azide free ab215368

リコンビナント **RabMAb**

26 References **画像数 12**

製品の概要

製品名	Anti-alpha smooth muscle Actin (acetyl E3) + ACTG2 (acetyl E3) antibody [E184] - BSA and Azide free
製品の詳細	Rabbit monoclonal [E184] to alpha smooth muscle Actin (acetyl E3) + ACTG2 (acetyl E3) - BSA and Azide free
由来種	Rabbit
特異性	<p>For immunohistochemistry, this antibody only detects actin in smooth muscle and not cardiac muscle.</p> <p>This antibody has been shown to detect P62736-ACTA (gene <i>ACTA2</i>): (Ac)-EEEDSTALVC and P63267-ACTH (gene <i>ACTG2</i>): (Ac)-EEETTALVC in indirect ELISA.</p>
アプリケーション	適用あり: IHC-Fr, Flow Cyt (Intra), ICC/IF, WB, IHC-P, ELISA
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: A431, HeLa, C6, RAW264.7, PC-12, NIH/3T3 and MCF-7 cell lysates. IHC-P: Human uterus, human smooth muscle and mouse smooth muscle tissues. Flow Cyt (intra): HeLa cells. ICC/IF: A431
特記事項	<p>ab215368 is the carrier-free version of ab32575.</p> <p>Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.</p> <p>This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.</p> <p>Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.</p> <p>This product is compatible with the Maxpar[®] Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] is a trademark of Fluidigm Canada Inc.</p> <p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p>

- 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. Do Not Freeze.
バッファー	pH: 7.20 Constituent: PBS
キャリア・フリー	はい
精製度	Protein A purified
ポリ/モノ	モノクローナル
クローン名	E184
アイソタイプ	IgG

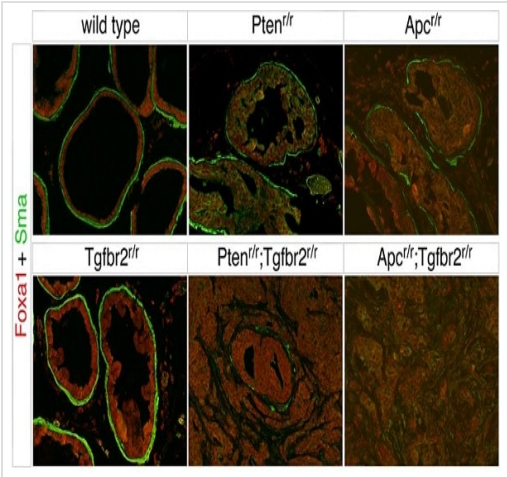
アプリケーション

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ab215368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
IHC-Fr		Use at an assay dependent concentration.
Flow Cyt (Intra)		Use at an assay dependent concentration. ab199376 - Rabbit monoclonal IgG, is suitable for use as an isotype control with this antibody.
ICC/IF		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Detects a band of approximately 42 kDa (predicted molecular weight: 42 kDa).
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval before commencing with IHC staining protocol. See IHC antigen retrieval protocols .
ELISA		Use a concentration of 2e-005 - 1 µg/ml.

ターゲット情報

画像



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-alpha smooth muscle Actin (acetyl E3) + ACTG2 (acetyl E3) antibody [E184] - BSA and Azide free (ab215368)

Image from Bjerke GA. et al PLoS One. 2014 Mar 20;9(3):e92800. doi: 10.1371/journal.pone.0092800. eCollection 2014.

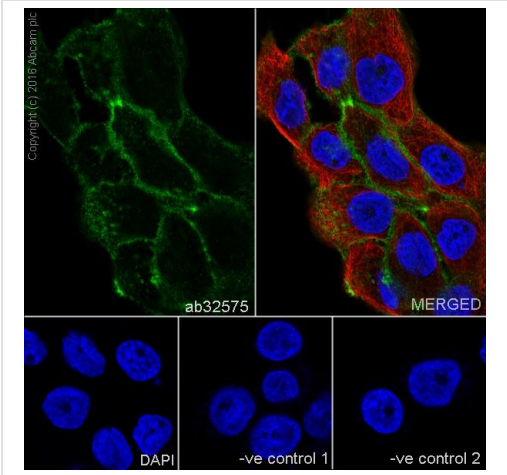
FoxA1 (red) and alpha smooth muscle actin (green) staining are shown by indirect immunofluorescence on sections of prostate from mice of the indicated genotypes.

Wild type: 21 weeks, *Tgfb^{2fl/r}*: 44 weeks, *Pten^{fl/r}*: 21 weeks, *Pten^{fl/r};Tgfb^{2fl/r}*: 11 weeks, *Apc^{fl/r}*: 36 weeks, and *Apc^{fl/r};Tgfb^{2fl/r}*: 24 weeks old.

IF images were captured on an Olympus BX51 microscope and DP70 digital camera, or on a Nikon Eclipse NI-U and captured with a DS-Q11 camera with NIS Elements software.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide ([ab32575](#)).

Perform heat mediated antigen retrieval before commencing with IHC staining protocol.

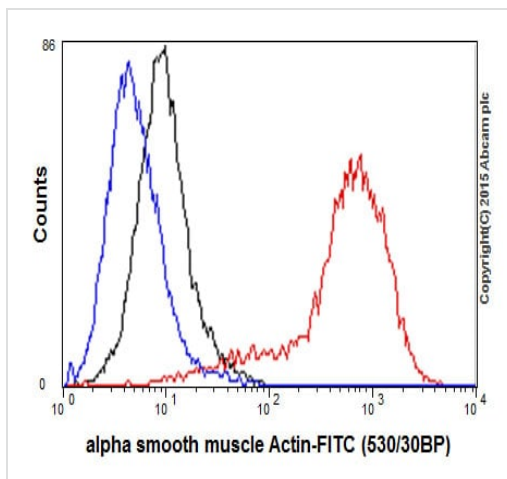


Immunocytochemistry/ Immunofluorescence - Anti-alpha smooth muscle Actin (acetyl E3) + ACTG2 (acetyl E3) antibody [E184] - BSA and Azide free (ab215368)

Immunocytochemistry/Immunofluorescence analysis of A431 (human epidermoid carcinoma) cells labeling alpha smooth muscle Actin (green) with purified [ab32575](#) at 1/500. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. [ab150077](#), Alexa Fluor[®] 488-conjugated goat anti-rabbit IgG (1/1000) was used as the secondary antibody. Cells were counterstained with [ab7291](#), anti-Tubulin (mouse mAb) at 1/1000 followed by [ab150120](#) Alexa Fluor[®]594 goat anti-mouse secondary (1/1000). Nuclei were counterstained with DAPI (blue).

For negative control 1, rabbit primary antibody and anti-mouse secondary antibody ([ab150120](#)) were used. For negative control 2, [ab7291](#) (mouse primary antibody) was used followed by anti-rabbit secondary antibody ([ab150077](#)).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide ([ab32575](#)).

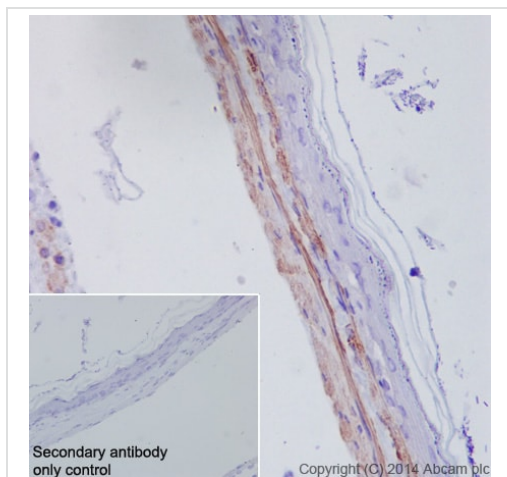


Flow Cytometry (Intracellular) - Anti-alpha smooth muscle Actin (acetyl E3) + ACTG2 (acetyl E3) antibody [E184] - BSA and Azide free (ab215368)

Intracellular Flow Cytometry analysis of HeLa (Human epithelial cell line from cervix adenocarcinoma) cells labeling alpha smooth muscle Actin with purified **ab32575** at a dilution of 1/20 (red).

Cells were fixed with 2% paraformaldehyde. A FITC-conjugated goat anti-rabbit IgG (1/150) was used as the secondary antibody. Black - Isotype control, rabbit monoclonal IgG. Blue - Unlabeled control, cells without incubation with primary and secondary antibodies.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab32575**).



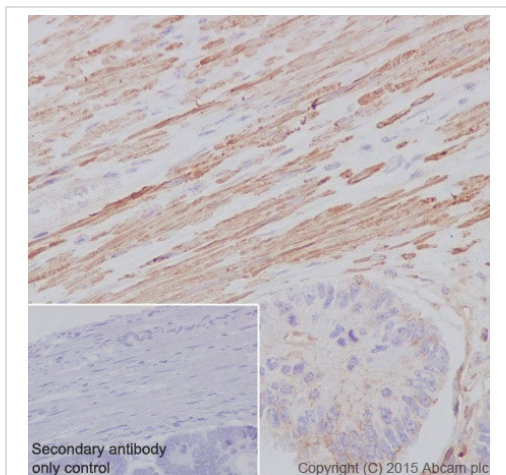
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-alpha smooth muscle Actin (acetyl E3) + ACTG2 (acetyl E3) antibody [E184] - BSA and Azide free (ab215368)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of mouse smooth muscle tissue labeling alpha smooth muscle Actin with purified **ab32575** at a dilution of 1/200. Heat mediated antigen retrieval was performed using EDTA buffer pH 9. **ab97051**, an HRP-conjugated goat anti-rabbit IgG (H+L) was used as the secondary antibody (1/500).

Negative control using PBS instead of primary antibody.

Counterstained with hematoxylin.

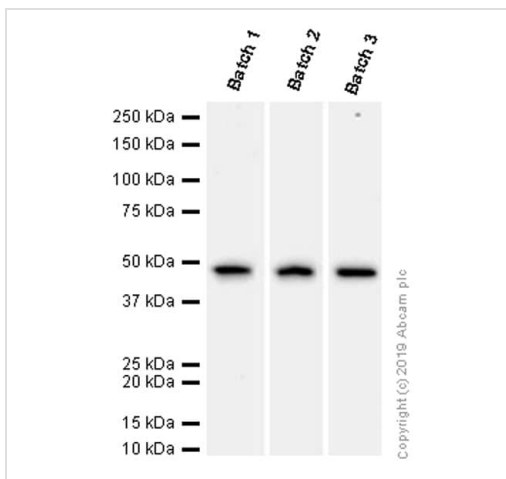
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab32575**).



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-alpha smooth muscle Actin (acetyl E3) + ACTG2 (acetyl E3) antibody [E184] - BSA and Azide free (ab215368)

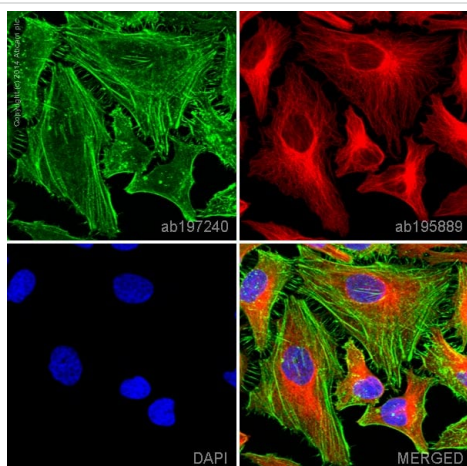
This IHC data was generated using the same anti-alpha smooth muscle Actin antibody clone, E184, in a different buffer formulation (cat# **ab32575**).

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human smooth muscle tissue labelling alpha smooth muscle Actin with purified **ab32575** at a dilution of 1/200. Heat mediated antigen retrieval was performed using EDTA buffer pH 9. **ab97051**, a HRP-conjugated goat anti-rabbit IgG (H+L) was used as the secondary antibody (1/500). Negative control using PBS instead of primary antibody. Counterstained with hematoxylin.



Western blot - Anti-alpha smooth muscle Actin (acetyl E3) + ACTG2 (acetyl E3) antibody [E184] - BSA and Azide free (ab215368)

This data was developed using **ab32575**, the same antibody clone in a different buffer formulation. Different batches of **ab32575** were tested on A431 (Human epidermoid carcinoma epithelial cell) lysate at 0.005 µg/ml. 15 µg of lysate was loaded in each lane. Bands observed at 42 kDa.



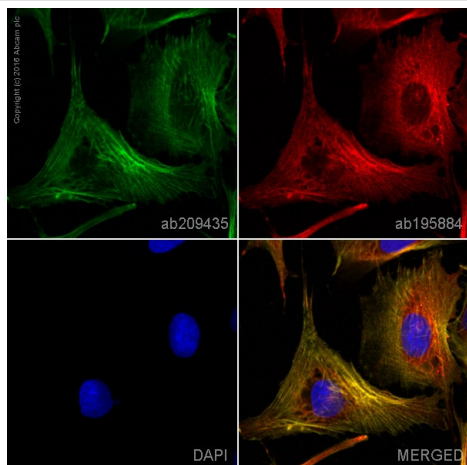
Immunocytochemistry/ Immunofluorescence - Anti-alpha smooth muscle Actin (acetyl E3) + ACTG2 (acetyl E3) antibody [E184] - BSA and Azide free (ab215368)

Clone E184 (ab215368) has been successfully conjugated by Abcam. This image was generated using Anti-alpha smooth muscle Actin antibody [E184] (Alexa Fluor® 488). Please refer to [ab197240](#) for protocol details.

[ab197240](#) staining alpha smooth muscle actin in HeLa cells. The cells were fixed with 4% formaldehyde (10 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated overnight at +4°C with [ab197240](#) at a 1/100 dilution (shown in green) and [ab195889](#), Mouse monoclonal to alpha Tubulin (Alexa Fluor® 594), at a 1/250 dilution (shown in red). Nuclear DNA was labelled with DAPI (shown in blue).

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

This product also gave a positive signal under the same testing conditions in HeLa cells fixed with 100% methanol (5 min).

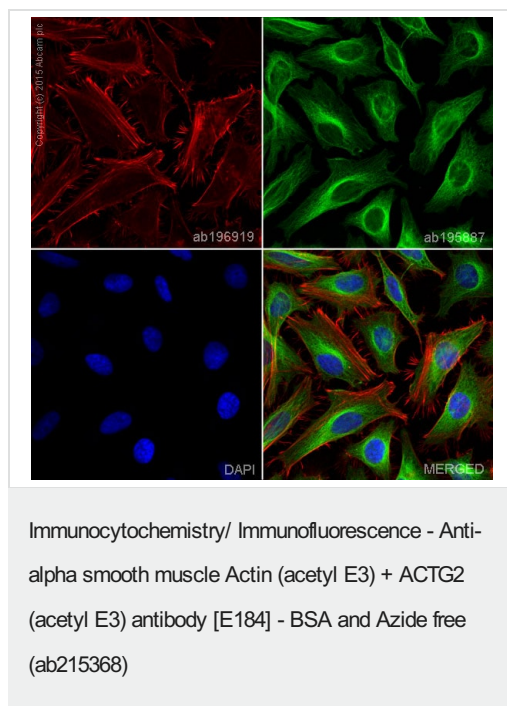


Immunocytochemistry/ Immunofluorescence - Anti-alpha smooth muscle Actin (acetyl E3) + ACTG2 (acetyl E3) antibody [E184] - BSA and Azide free (ab215368)

Clone E184 (ab215368) has been successfully conjugated by Abcam. This image was generated using Anti-alpha smooth muscle Actin antibody [E184] (PE). Please refer to [ab209435](#) for protocol details.

[ab209435](#) staining alpha smooth muscle Actin in SV40LT-SMC cells. The cells were fixed with 4% formaldehyde (10 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with [ab209435](#) at 1/500 dilution (Pseudocolored in green) and [ab195884](#), Rat monoclonal to Tubulin (Alexa Fluor® 647), at 1/250 dilution (shown in red). Nuclear DNA was labelled with DAPI (shown in blue).

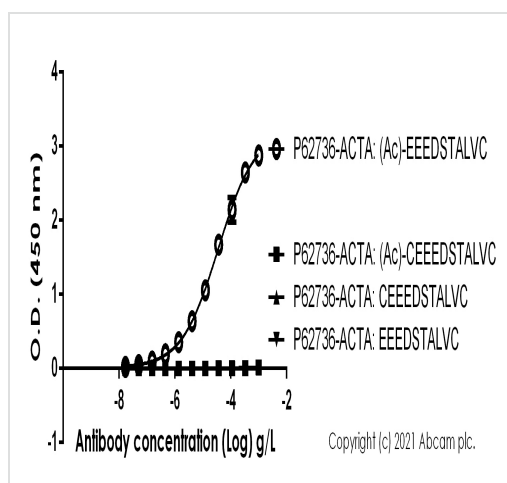
Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).



Clone E184 (ab215368) has been successfully conjugated by Abcam. This image was generated using Anti-alpha smooth muscle Actin antibody [E184] (Alexa Fluor® 647). Please refer to [ab196919](#) for protocol details.

[ab196919](#) staining alpha Smooth Muscle Actin in HeLa cells. The cells were fixed with 4% PFA (10min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with [ab196919](#) at 1/100 dilution (shown in red) and [ab195887](#), Mouse monoclonal to alpha Tubulin (Alexa Fluor® 488), at 2µg/ml (shown in green). Nuclear DNA was labelled with DAPI (shown in blue).

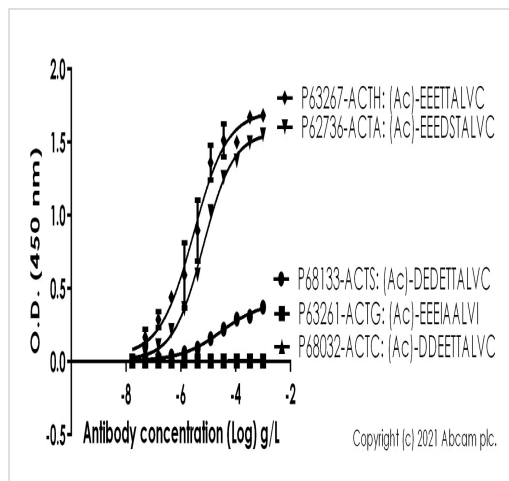
Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).



Following reconstitution in PBS with 10% DMSO, peptides (1 ug per mL) were immobilised in PBS on an ELISA plate overnight. After blocking in 5% BSA, primary antibody ([ab271180](#)) was added in a concentration range of 0.017-1000 ng per mL.

Pre-adsorbed secondary antibody goat anti-rabbit IgG H&L (HRP, [ab97080](#)) was used at 1/20000 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide ([ab32575](#)).



ELISA - Anti-alpha smooth muscle Actin (acetyl E3)
+ ACTG2 (acetyl E3) antibody [E184] - BSA and
Azide free (ab215368)

Following reconstitution in PBS with 10% DMSO, peptides (1 ug per mL) were immobilised in PBS on an ELISA plate overnight. After blocking in 5% BSA, primary antibody ([ab271180](#)) was added in a concentration range of 0.017-1000 ng per mL.

Pre-adsorbed secondary antibody goat anti-rabbit IgG H&L (HRP, [ab97080](#)) was used at 1/20000 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide ([ab32575](#)).

Why choose a recombinant antibody?



Research with confidence
Consistent and reproducible results



Long-term and scalable supply
Recombinant technology



Success from the first experiment
Confirmed specificity



Ethical standards compliant
Animal-free production

Anti-alpha smooth muscle Actin (acetyl E3) +
ACTG2 (acetyl E3) antibody [E184] - BSA and
Azide free (ab215368)

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