abcam

Product datasheet

Alexa Fluor® 488 Anti-Cytokeratin 8 antibody [EP1628Y] ab192467

יובעדער RabMAb

*** * * 4 Abreviews 12 References 画像数6

製品の概要

免疫原

製品名 Alexa Fluor® 488 Anti-Cytokeratin 8 antibody [EP1628Y]

製品の詳細 Alexa Fluor® 488 Rabbit monoclonal [EP1628Y] to Cytokeratin 8

由来種 Rabbit

Alexa Fluor® 488. Ex: 495nm, Em: 519nm 標識

アプリケーション 適用あり: ICC, Flow Cyt (Intra)

種交差性 交差種: Human

交差が予測される動物種: Mouse 🔷

Synthetic peptide within Human Cytokeratin 8 aa 300-400 (C terminal). The exact sequence is

proprietary.

Database link: P05787

ポジティブ・コントロール ICC/IF: HeLa cells Flow Cyt (intra): HeLa cells

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit 特記事項 monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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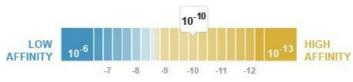
outlicensing@thermofisher.com.

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.

Avoid freeze / thaw cycle. Stable for 12 months at -20°C. Store In the Dark.

解離定数(K_D 値) $K_D = 4.60 \times 10^{-10} M$



Learn more about K_D

バッファー pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 30% Glycerol (glycerin, glycerine), 1% BSA, PBS

精製度 Protein A purified

ポリ/モノ モノクローナル **クローン名** EP1628Y

アイソタイプ lgG

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ab192467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		1/100.
Flow Cyt (Intra)		1/500.

ターゲット情報

機能 Together with KRT19, helps to link the contractile apparatus to dystrophin at the costameres of

striated muscle.

組織特異性 Observed in muscle fibers accumulating in the costameres of myoplasm at the sarcolemma

membrane in structures that contain dystrophin and spectrin. Expressed in gingival mucosa and

hard palate of the oral cavity.

関連疾患 Cirrhosis

配列類似性 Belongs to the intermediate filament family.

翻訳後修飾 Phosphorylation on serine residues is enhanced during EGF stimulation and mitosis. Ser-74

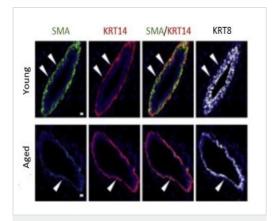
phosphorylation plays an important role in keratin filament reorganization.

O-glycosylated. O-GlcNAcylation at multiple sites increases solubility, and decreases stability by

inducing proteasomal degradation.

O-glycosylated (O-GlcNAcylated), in a cell cycle-dependent manner.

細胞内局在 Cytoplasm. Nucleus, nucleoplasm. Nucleus matrix.



Immunofluorescence analysis of mouse myoepithelial cells labelling KRT8 (right) with ab192467 at 1/100 dilution. The tissue was fixed in 10% neutral buffered formalin overnight. Paraffin embedding and sectioning were performed by the Rodent Histopathology Core at Harvard Medical School. Scale bar, 10 μ m.

Immunocytochemistry - Alexa Fluor® 488 Anti-

Cytokeratin 8 antibody [EP1628Y] (ab192467)

Li CMet al; Cell Rep. 2020 Dec 29;33(13):108566. doi: 10.1016/j.celrep.2020.108566. Reproduced under the Creative Commons license: https://creativecommons.org/licenses/by/4.0/

REG HGPIN/ADENO NEPC

DAPI

100m

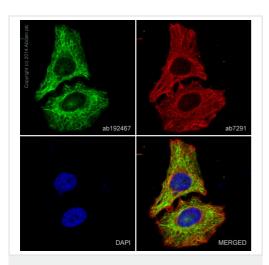
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Immunofluorescence analysis of mouse postate tumor samples labelling cytokeratin 8 (green) with ab192467 at 1/100 dilution. SYP was also stained using <u>ab206870</u> (red). Cells were fixed with formalin and embedded in paraffin. Sections were blocked with PBS-Tween (0.1%) containing 5% of BSA. Primary conjugated antibodies were simultaneously incubated overnight at 4°C. Nuclear DNA was labelled with DAPI (blue). Scale bar = 100 μm.

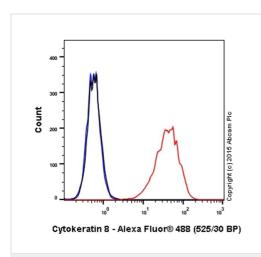
Immunocytochemistry - Alexa Fluor® 488 Anti-

Cytokeratin 8 antibody [EP1628Y] (ab192467)

Sulsenti R et al; Front Immunol. 2021 Mar 2;12:622001. doi: 10.3389/fimmu.2021.622001. Reproduced under the Creative Commons license: https://creativecommons.org/licenses/by/4.0/



Immunocytochemistry - Alexa Fluor® 488 Anti-Cytokeratin 8 antibody [EP1628Y] (ab192467)



Flow Cytometry (Intracellular) - Alexa Fluor® 488 Anti-Cytokeratin 8 antibody [EP1628Y] (ab192467)

Immunofluorescence staining of cytokeratin 8 in HeLa cells using ab192467. The cells were fixed with 4% formaldehyde (10 min) and then blocked in 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Triton X-100 for 1hr. The cells were then incubated with **ab192647** at a working dilution of 1 in 100 (shown in green) and **ab7291** (Mouse monoclonal [DM1A] to alpha Tubulin) at 1 μ g/ml overnight at +4°C, followed by a further incubation at room temperature for 1hr with AlexaFluor[®] 594 Goat anti-mouse lgG (H&L - preadsorbed) (**ab150120**) at 2 μ g/ml (shown in pseudocolor red).

Nuclear DNA was labeled in blue with DAPI.

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

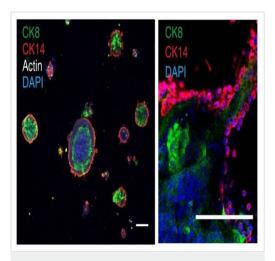
Image was taken with a Confocal microscope (Leica-microsystems, TCS SP8)

Flow cytometry analysis of HeLa cells stained with ab192467. The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab192467, 1/500 dilution) for 30 min at 22°C.

Isotype control antibody (black line) was rabbit monoclonal IgG [EPR25A] Alexa Fluor[®] 488 (<u>ab199091</u>) used at the same concentration and conditions as the primary antibody. Unlabeled sample (blue line) was also used as a control.

Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.

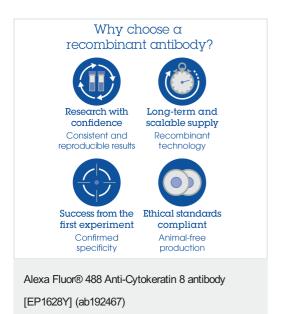
This antibody gave a positive signal in HeLa cells fixed with 4% formaldehyde (10 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.



Immunocytochemistry - Alexa Fluor® 488 Anti-Cytokeratin 8 antibody [EP1628Y] (ab192467)

Rosenbluth J et al., Nat Commun, 11(1), 1711. Fig 1c.; doi: 10.1038/s41467-020-15548-7. Reproduced under the Creative Commons license http://creativecommons.org/licenses/by/4.0/.

Immunofluorescence staining in human mammary organoids of cytokeratin 8 using ab192467 (green), cytokeratin 14 using **ab206100** (red), and actin (white). Cells were fixed with 4% paraformaldehyde for 20 minutes at room temperature and permeabilized with 0.5% Triton X-100 for 10 minutes at 4 °C. Primary antibodies incubated overnight at 4 °C. Nuclear DNA was labelled with DAPI (blue). Scale bar = 100 μ m. Organoids were imaged by confocal microscopy.



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