abcam

Product datasheet

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

重组 RabMAb

★★★★ 4 Abreviews 12 References 6 图像

概述

产品名称 Alexa Fluor® 488荧光Anti-Cytokeratin 8抗体[EP1628Y]

Alexa Fluor® 488荧光兔单克隆抗体[EP1628Y] to Cytokeratin 8 描述

宿主 Rabbit

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

经测试应用 适用于: ICC, Flow Cyt (Intra)

种属反应性 与反应: Human

预测可用于: Mouse

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

proprietary.

Database link: P05787

ICC/IF: HeLa cells Flow Cyt (intra): HeLa cells 阳性对照

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit 常规说明

monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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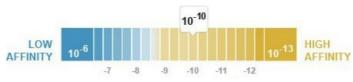
outlicensing@thermofisher.com.

形式 Liquid

存放说明 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.

解离常数(K_D) $K_D = 4.60 \times 10^{-10} M$



Learn more about K_D

存储溶液 pH: 7.40

Preservative: 0.02% Sodium azide

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

纯度 Protein A purified

 克隆
 单克隆

 克隆编号
 EP1628Y

同种型 lgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab192467于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
ICC		1/100.
Flow Cyt (Intra)		1/500.

靶标

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

striated muscle.

组织特异性 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.

疾病相关 Cirrhosis

序列相似性 Belongs to the intermediate filament family.

翻译后修饰 Phosphorylation on serine residues is enhanced during EGF stimulation and mitosis. Ser-74

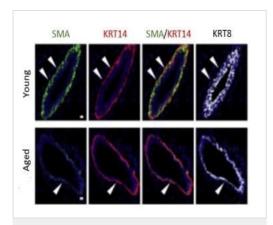
phosphorylation plays an important role in keratin filament reorganization.

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.

细胞定位 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 μm .

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

100pm

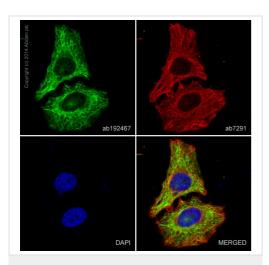
labelling cytokeratin 8 (green) with ab192467 at 1/100 dilution. SYP was also stained using **ab206870** (red). Cells were fixed with formalin and embedded in paraffin. Sections were blocked with PBS-Tween (0.1%) containing 5% of BSA. Primary conjugated antibodies were simultaneously incubated overnight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

Immunofluorescence analysis of mouse postate tumor samples

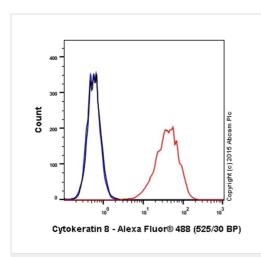
Immunocytochemistry - Alexa Fluor® 488 Anti-

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.

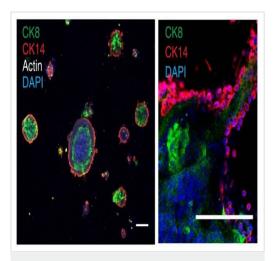
Image was taken with a Confocal microscope (Leica-microsystems, TCS SP8)

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.

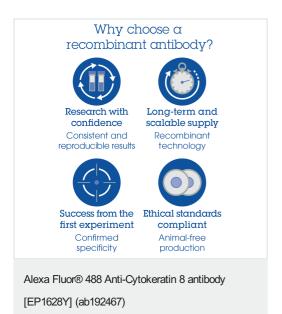
This antibody gave a positive signal in HeLa cells fixed with 4% formaldehyde (10 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.



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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