abcam

Product datasheet

Alexa Fluor® 647 Anti-Cytokeratin 13 antibody [EPR3671] ab198585

Recombinant RabMAb

3 Images

Overview

Product name Alexa Fluor® 647 Anti-Cytokeratin 13 antibody [EPR3671]

Description Alexa Fluor® 647 Rabbit monoclonal [EPR3671] to Cytokeratin 13

Host species Rabbit

Conjugation Alexa Fluor® 647. Ex: 652nm. Em: 668nm

Tested applications Suitable for: Flow Cyt (Intra), ICC/IF

Species reactivity Reacts with: Human

Predicted to work with: Mouse

Does not react with: Rat

Synthetic peptide. This information is proprietary to Abcam and/or its suppliers. **Immunogen**

Positive control Flow Cyt (intra): A431 cells. ICC/IF: A431 cells.

This product is a recombinant monoclonal antibody, which offers several advantages including: **General notes**

- 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**.

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1

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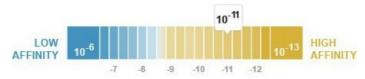
Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle. Store In the Dark.

Dissociation constant (K_D) $K_D = 1.20 \times 10^{-11} M$



Learn more about K_D

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

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

Purity Protein A purified

ClonalityMonoclonalClone numberEPR3671

Isotype IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab198585 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
Flow Cyt (Intra)		1/50. ab199093 - Rabbit monoclonal lgG (Alexa Fluor® 647), is suitable for use as an isotype control with this antibody.
ICC/IF		1/100. This product gave a positive signal in A431 cells fixed with 4% formaldehyde (10 min) and 100% methanol (5 min).

Target

Tissue specificity Expressed in some epidermal sweat gland ducts (at protein level) and in exocervix, esophagus

and placenta.

Involvement in disease

Defects in KRT13 are a cause of white sponge nevus of cannon (WSN) [MIM:193900]. WSN is a rare autosomal dominant disorder which predominantly affects non-cornified stratified squamous epithelia. Clinically, it is characterized by the presence of soft, white, and spongy plaques in the oral mucosa. The characteristic histopathologic features are epithelial thickening, parakeratosis, and vacuolization of the suprabasal layer of oral epithelial keratinocytes. Less frequently the

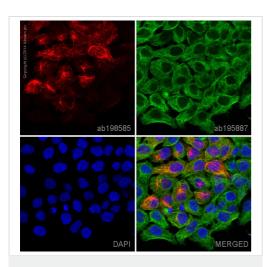
modifications

mucous membranes of the nose, esophagus, genitalia and rectum are involved.

Belongs to the intermediate filament family.

O-glycosylated; glycans consist of single N-acetylglucosamine residues.

Images



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 647 Anti-Cytokeratin 13 antibody [EPR3671] (ab198585)

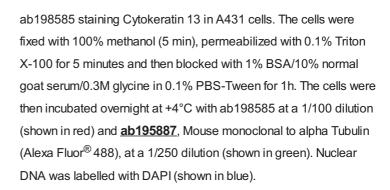
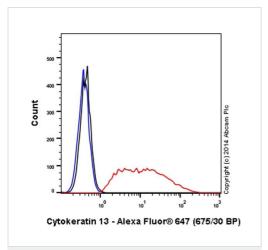


Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

This product also gave a positive signal under the same testing conditions in A431 cells fixed with 4% formaldehyde (10 min).

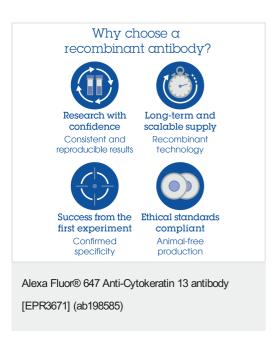


Flow Cytometry (Intracellular) - Alexa Fluor® 647 Anti-Cytokeratin 13 antibody [EPR3671] (ab198585)

Overlay histogram showing A431 cells stained with ab198585 (red line). The cells were fixed with 4% formaldehyde (10 min) and then permeabilized with 0.1% PBS-Triton X-100 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 (ab198585, 1/50 dilution) for 30 min at 22°C. Isotype control antibody (black line) was rabbit IgG (monoclonal) Alexa Fluor[®] 647 used at the same concentration and conditions as the primary antibody. Unlabelled sample (blue line) was also used as a control.

Acquisition of >5,000 events were collected using a solid-state 25mW red diode laser (635 nm) and 675/30 bandpass filter.

This antibody gave a positive signal in A431 fixed with 80% methanol (5 min)/permeabilized with 0.1% PBS-Triton X-100 for 20 min used under the same conditions.



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