abcam

Product datasheet

HRP Anti-Cytokeratin 13 antibody [EPR3671] ab199092



RabMAb

3 Images

Overview

Product name HRP Anti-Cytokeratin 13 antibody [EPR3671]

Description HRP Rabbit monoclonal [EPR3671] to Cytokeratin 13

Host species Rabbit
Conjugation HRP

Tested applications
Suitable for: WB, IHC-P
Species reactivity
Reacts with: Human

Predicted to work with: Mouse

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

Positive control WB: A431 whole cell lysate. IHC-P: FFPE human tonsil (normal) tissue sections.

General notesOur RabMAb[®] technology is a patented hybridoma-based technology for making rabbit

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

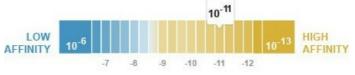
Properties

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 KD

Storage buffer pH: 7.40

Preservative: 0.1% Proclin 300 Solution

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

Purity Protein A purified

ClonalityMonoclonalClone numberEPR3671

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Isotype IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab199092 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
WB		1/5000. Detects a band of approximately 50 kDa (predicted molecular weight: 50 kDa).
IHC-P		1/100. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

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

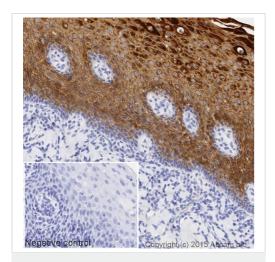
Sequence similarities

Belongs to the intermediate filament family.

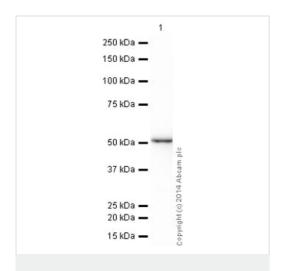
Post-translational modifications

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

Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - HRP Anti-Cytokeratin 13 antibody [EPR3671] (ab199092)



Western blot - HRP Anti-Cytokeratin 13 antibody [EPR3671] (ab199092)

IHC image of Cytokeratin 13 staining in a section of formalin-fixed paraffin-embedded normal human tonsil tissue*, performed on a Leica BONDTM. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20mins. The section was then incubated with ab199092, 1/100 dilution, for 15 mins at room temperature. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX. The inset negative control image is taken from an identical assay without primary antibody.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

*Tissue obtained from the Human Research Tissue Bank, supported by the NIHR Cambridge Biomedical Research Centre

HRP Anti-Cytokeratin 13 antibody [EPR3671] (ab199092) at 1/5000 dilution + A431 (Human epithelial carcinoma cell line) Whole Cell Lysate at 10 µg

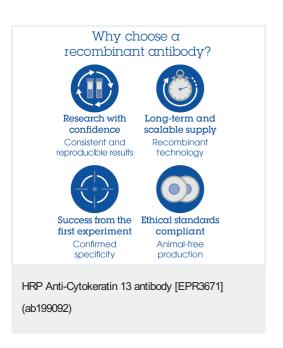
Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 50 kDa **Observed band size:** 50 kDa

Exposure time: 10 seconds

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 2% Bovine Serum Albumin before being incubated with ab199092 overnight at 4°C. Antibody binding was visualised using ECL development solution **ab133406**.



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