abcam

Product datasheet

HRP Anti-Cytokeratin 10 antibody [EP1607IHCY] ab194232



重组 RabMAb

3 图像

概述

产品名称 HRP Anti-Cytokeratin 10抗体[EP1607IHCY]

描述 HRP兔单克隆抗体[EP1607IHCY] to Cytokeratin 10

宿主 Rabbit 偶联物 HRP

经测试应用 适用于: IHC-P, WB

种属反应性 与反应: Human

预测可用于: Mouse, Rat 🔷

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

阳性对照 WB: HaCat whole cell lysate. IHC-P: Normal human skin tissue.

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**.

性能

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. Store In the Dark.

存储溶液 pH: 7.40

Preservative: 0.1% Proclin 300 Solution

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

纯度 Protein A purified

克隆 单克隆

克隆编号 EP1607IHCY

同种型 ΙgG

应用

Abpromise™承诺保证使用ab194232于以下的经测试应用 The Abpromise guarantee

"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
IHC-P		1/100. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.
WB		1/5000. Detects a band of approximately 60 kDa (predicted molecular weight: 60 kDa).

靶标

组织**特异性**

疾病相关

Seen in all suprabasal cell layers including stratum corneum.

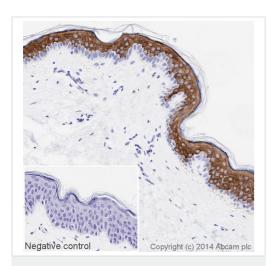
Defects in KRT10 are a cause of bullous congenital ichthyosiform erythroderma (BCIE) [MIM:113800]; also known as epidermolytic hyperkeratosis (EHK) or bullous erythroderma ichthyosiformis congenita of Brocq. BCIE is an autosomal dominant skin disorder characterized by widespread blistering and an ichthyotic erythroderma at birth that persist into adulthood. Histologically there is a diffuse epidermolytic degeneration in the lower spinous layer of the epidermis. Within a few weeks from birth, erythroderma and blister formation diminish and hyperkeratoses develop.

Defects in KRT10 are a cause of ichthyosis annular epidermolytic (AEI) [MIM:607602]; also known as cyclic ichthyosis with epidermolytic hyperkeratosis. AEI is a skin disorder resembling bullous congenital ichthyosiform erythroderma. Affected individuals present with bullous ichthyosis in early childhood and hyperkeratotic lichenified plaques in the flexural areas and extensor surfaces at later ages. The feature that distinguishes AEI from BCIE is dramatic episodes of flares of annular polycyclic plaques with scale, which coalesce to involve most of the body surface and can persist for several weeks or even months.

序列相似性

Belongs to the intermediate filament family.

图片



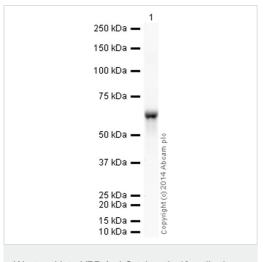
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - HRP Anti-Cytokeratin 10 antibody [EP1607IHCY] (ab194232)

IHC image of Cytokeratin 10 staining in a section of formalin-fixed paraffin-embedded normal human skin*, performed on a Leica BOND. 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 ab194232 at 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



Western blot - HRP Anti-Cytokeratin 10 antibody [EP1607IHCY] (ab194232) HRP Anti-Cytokeratin 10 antibody [EP1607lHCY] (ab194232) at 1/5000 dilution + HaCat whole cell lysate at 10 μg

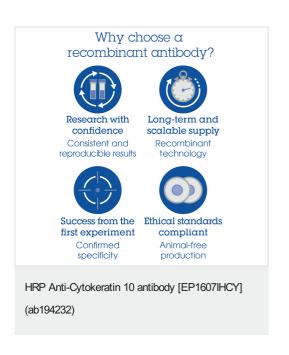
Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 60 kDa **Observed band size:** 60 kDa

Exposure time: 2 minutes

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 3% milk before being incubated with ab194232 overnight at 4°C. Antibody binding was visualised using ECL development solution **ab133406**.



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