abcam

Product datasheet

Anti-Cytokeratin 10 antibody [rKRT10/1275] - BSA and Azide free ab236306



2 Images

Overview

Product name Anti-Cytokeratin 10 antibody [rKRT10/1275] - BSA and Azide free

Description Mouse monoclonal [rKRT10/1275] to Cytokeratin 10 - BSA and Azide free

Host species Mouse

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

Immunogen Recombinant full length protein corresponding to Human Cytokeratin 10 aa 1 to the C-terminus.

Database link: P13645

Run BLAST with
Run BLAST with

Positive control IHC-P: Human skin tissue.

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 long

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A/G purified

Purification notes Purified from Bioreactor Concentrate by Protein A/G.

Clonality Monoclonal
Clone number rKRT10/1275

Light chain type lgG1 kappa

Applications

1

The Abpromise guarantee

Our Abpromise guarantee covers the use of ab236306 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
IHC-P		Use a concentration of 0.1 - 0.2 µg/ml. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. (Primary incubation for 30 min at room temperature).

Target

Tissue specificity Involvement in disease

Seen in all suprabasal cell layers including stratum corneum.

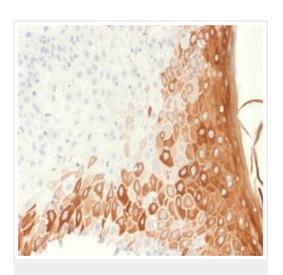
Defects in KRT10 are a cause of bullous congenital ichthyosiform erythroderma (BCIE) [MIM:113800]; also known as epidermolytic hyperkeratosis (EHK) or bullous erythroderma ichthyosiformis congenita of Brocq. BCIE is an autosomal dominant skin disorder characterized by widespread blistering and an ichthyotic 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.

Sequence similarities

Belongs to the intermediate filament family.

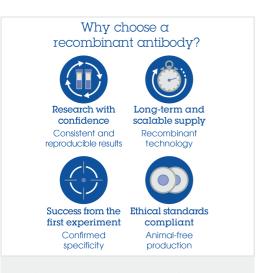
Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Cytokeratin 10 antibody [rKRT10/1275] - BSA and Azide free (ab236306)

Formalin-fixed, paraffin-embedded human skin tissue stained for Cytokeratin 10 using <u>ab234330</u> at 0.2 µg/ml in immunohistochemical analysis.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab234330).



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