abcam

Product datasheet

Anti-Collagen VII antibody - BSA and Azide free ab93350

★★★★★ 2 Abreviews 14 References 画像数 1

製品の概要

製品名 Anti-Collagen VII antibody - BSA and Azide free

製品の詳細 Rabbit polyclonal to Collagen VII - BSA and Azide free

由来種 Rabbit

アプリケーション 適用あり: ICC/IF

種交差性 交差種: Human, Common marmoset

免疫原 Full length native protein (purified) corresponding to Human Collagen VII.

特記事項

The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.

パッファー Constituent: PBS

キャリア・フリー はい

精製度 Affinity purified ポリクローナル

アイソタイプ IgG

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab93350の使用に適用されますアプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
ICC/IF		

追加情報 ELISA: Use at an assay dependent dilution.

EIA: Use at an assay dependent dilution. Flow Cyt: Use at an assay dependent dilution. ICC/IF: Use at a concentration of 2.5 - 10 µg/ml.

IP: Use at a concentration of 2.5 - 10 μg/ml.

WB: Use at a concentration of 0.2 - 0.8 µg/ml. Predicted molecular weight: 295 kDa.

Not yet tested in other applications.

Optimal dilutions/concentrations should be determined by the end user.

ターゲット情報

機能 Stratified squamous epithelial basement membrane protein that forms anchoring fibrils which may

contribute to epithelial basement membrane organization and adherence by interacting with

extracellular matrix (ECM) proteins such as type IV collagen.

関連疾患 Epidermolysis bullosa acquisita (EBA) is an autoimmune acquired blistering skin disease

resulting from autoantibodies to type VII collagen.

Epidermolysis bullosa dystrophica, autosomal dominant Epidermolysis bullosa dystrophica, autosomal recessive

Epidermolysis bullosa dystrophica, Pasini type

Epidermolysis bullosa dystrophica, Hallopeau-Siemens type

Transient bullous dermolysis of the newborn Epidermolysis bullosa dystrophica, pretibial type Epidermolysis bullosa dystrophica, Bart type

Epidermolysis bullosa pruriginosa

Nail disorder, non-syndromic congenital, 8

Epidermolysis bullosa dystrophica, with subcorneal cleavage

配列類似性 Contains 1 BPTI/Kunitz inhibitor domain.

Contains 9 fibronectin type-III domains.

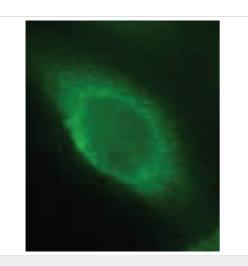
Contains 2 VWFA domains.

翻訳後修飾 Prolines at the third position of the tripeptide repeating unit (G-X-Y) are hydroxylated in some or all

of the chains.

細胞内局在 Secreted > extracellular space > extracellular matrix > basement membrane.

画像



Immunocytochemistry/ Immunofluorescence - Anti-Collagen VII antibody - BSA and Azide free (ab93350)

Immunofluorescence staining of Collagen VII in normal human keratinocytes. Second antibody for staining of cells: goat-anti-rabbit IgG Alexa 488

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

Our Abpromise to you: Quality guaranteed and expert technical support

- Replacement or refund for products not performing as stated on the datasheet
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- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

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