abcam

Product datasheet

Anti-TGFBI antibody ab122975

★★★★★ 4 Abreviews 2 Images

Overview

Product name Anti-TGFBI antibody

Description Goat polyclonal to TGFBI

Host species Goat

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

Predicted to work with: Mouse, Rat, Dog

A

Immunogen Synthetic peptide from an internal region of Human TGFBI, conjugated to KLH

Positive control Human Kidney, Renal Duct Human Kidney lysate

General notes

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

Properties

Form Liquid

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

Storage buffer pH: 7.30

Preservative: 0.02% Sodium azide

Constituents: 99% Tris buffered saline, 0.5% BSA

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab122975 in the following tested applications.

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

| Application | Abreviews | Notes |
|-------------|---------------------|---|
| WB | *** <u>*</u> ** (2) | Use a concentration of 0.2 - 0.5 μg/ml. Predicted molecular weight: 75 kDa. |
| IHC-P | | Use a concentration of 3.75 μg/ml. |

Target

Function

Binds to type I, II, and IV collagens. This adhesion protein may play an important role in cell-collagen interactions. In cartilage, may be involved in endochondral bone formation.

Tissue specificity

Involvement in disease

Highly expressed in the corneal epithelium.

Defects in TGFBI are the cause of epithelial basement membrane corneal dystrophy (EBMD) [MIM:121820]; also known as Cogan corneal dystrophy or map-dot-fingerprint type corneal dystrophy. EBMD is a bilateral anterior corneal dystrophy characterized by grayish epithelial fingerprint lines, geographic map-like lines, and dots (or microcysts) on slit-lamp examination. Pathologic studies show abnormal, redundant basement membrane and intraepithelial lacunae filled with cellular debris. Although this disorder usually is not considered to be inherited, families with autosomal dominant inheritance have been identified.

Defects in TGFBI are the cause of corneal dystrophy Groenouw type 1 (CDGG1) [MIM:121900]; also known as corneal dystrophy granular type. Inheritance is autosomal dominant. Corneal dystrophies show progressive opacification of the cornea leading to severe visual handicap. Defects in TGFBI are the cause of corneal dystrophy lattice type 1 (CDL1) [MIM:122200]. Inheritance is autosomal dominant.

Defects in TGFBI are a cause of corneal dystrophy Thiel-Behnke type (CDTB) [MIM:602082]; also known as corneal dystrophy of Bowman layer type 2 (CDB2).

Defects in TGFBI are the cause of Reis-Buecklers corneal dystrophy (CDRB) [MIM:608470]; also known as corneal dystrophy of Bowman layer type 1 (CDB1).

Defects in TGFBI are the cause of lattice corneal dystrophy type 3A (CDL3A) [MIM:608471].

CDL3A clinically resembles to lattice corneal dystrophy type 3, but differs in that its age of onset is 70 to 90 years. It has an autosomal dominant inheritance pattern.

Defects in TGFBI are the cause of Avellino corneal dystrophy (ACD) [MIM:607541]. ACD could be considered a variant of granular dystrophy with a significant amyloidogenic tendency. Inheritance is autosomal dominant.

Sequence similarities

Contains 1 EMI domain. Contains 4 FAS1 domains.

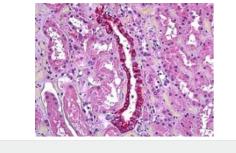
Post-translational modifications

Gamma-carboxyglutamate residues are formed by vitamin K dependent carboxylation. These residues are essential for the binding of calcium.

Cellular localization

Secreted > extracellular space > extracellular matrix. May be associated both with microfibrils and with the cell surface.

Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-TGFBI antibody (ab122975)

Immunohistochemistry of Formalin-Fixed, Paraffin-Embedded Human Kidney, Renal Duct, with ab122975 at 3.75ug/ml.



Western blot - Anti-TGFBI antibody (ab122975)

Anti-TGFBI antibody (ab122975) at 0.3 μ g/ml + Human Kidney lysate at 35 μ g/ml

Developed using the ECL technique.

Predicted band size: 75 kDa

Exposure time: 1 hour

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

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