abcam

Product datasheet

Anti-LTBP2 antibody ab121193

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概述

产**品名称** Anti-LTBP2抗体

描述 兔多克隆抗体to LTBP2

宿主 Rabbit

经测试应用 **适用于:** ICC, IHC-P

种属反应性 与反应: Human

免疫原 antigen sequence, corresponding to internal sequence amino acids 241-376 of Human LTBP2.

阳性对照 IHC-P: Human lung tissue; ICC: U-2 OS cells.

常规说明

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

性能

形式 Liquid

存放说明 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

存储溶液 pH: 7.20

Preservative: 0.02% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine)

纯**度** Immunogen affinity purified

克隆 多克隆

同种型 lgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab121193于以下的经测试应用

"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。

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应用	Ab评论	说明
ICC		Use a concentration of 0.25 - 2 µg/ml.
IHC-P		1/500 - 1/1000. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

靶标

功能 May play an integral structural role in elastic-fiber architectural organization and/or assembly.

组织特异性 Expressed in lung, weakly expressed in heart, placenta, liver and skeletal muscle.

疾病相关 Defects in LTBP2 are the cause of primary congenital glaucoma type 3D (GLC3D) [MIM:613086].

An autosomal recessive form of primary congenital glaucoma (PCG). PCG is characterized by marked increase of intraocular pressure at birth or early childhood, large ocular globes (buphthalmos) and corneal edema. It results from developmental defects of the trabecular meshwork and anterior chamber angle of the eye that prevent adequate drainage of aqueous

humor.

Defects in LTBP2 are the cause of microspherophakia (MCSPH) [MIM:251750]. It is a rare disease characterized by smaller and more spherical lenses than normal bilaterally, an increased anteroposterior thickness of the lens and highly myopic eyes. Lens dislocation or subluxation may

occur, leading to defective accommodation.

序列相似性 Belongs to the LTBP family.

Contains 20 EGF-like domains.

Contains 4 TB (TGF-beta binding) domains.

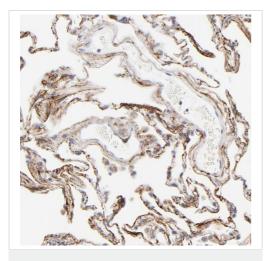
结**构域** Associates covalently with small latent TGF-beta complex via Repeat B and Repeat C.

翻译后修饰 Contains hydroxylated asparagine residues.

细胞定位 Secreted. Localized in nuchal ligament and aorta to the fibrillin-containing, microfibrillar

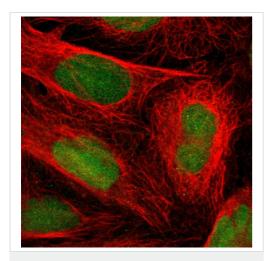
component of elastic fibers.

图片



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-LTBP2 antibody (ab121193)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human lung tissue labelling MLTBP2 with ab121193 at 1/500 dilution. Heat mediated antigen retrieval performed with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunocytochemistry - Anti-LTBP2 antibody (ab121193)

Immunocytochemistry analysis of U-2 OS (human bone osteosarcoma epithelial cell line) cells labelling MLTBP2 with ab121193 at 2 μ g/mL.

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

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