abcam

Product datasheet

Alexa Fluor® 488 Anti-muscle Actin antibody [EPR8484] ab190196



重组 RabMAb

2 图像

概述

产品名称 Alexa Fluor® 488荧光Anti-muscle Actin抗体[EPR8484]

描述 Alexa Fluor® 488荧光兔单克隆抗体[EPR8484] to muscle Actin

宿主 Rabbit

偶联物 Alexa Fluor® 488. Ex: 495nm, Em: 519nm

经测试应用 适用干: ICC/IF 种属反应性 与反应: Human

预测可用于: Mouse, Rat 📤

Synthetic peptide within Human muscle Actin aa 350 to the C-terminus. The exact immunogen 免疫原

> sequence used to generate this antibody is proprietary information. If additional detail on the immunogen is needed to determine the suitability of the antibody for your needs, please contact

our Scientific Support team to discuss your requirements.

Database link: P68032

Run BLAST with Run BLAST with

阳性对照 ICC/IF: HeLa cells.

常规说明 This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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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. Stable for 12 months at -20°C. Store In the Dark.

存储溶液 pH: 7.40

Preservative: 0.02% Sodium azide

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

纯**度** Protein A purified

同种型 IgG

应用

The Abpromise guarantee

Abpromise™承诺保证使用ab190196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。

应用	Ab评论	说明
ICC/IF		1/50.

靶标

功能

Actins are highly conserved proteins that are involved in various types of cell motility and are ubiquitously expressed in all eukaryotic cells.

疾病相关

Defects in ACTA1 are the cause of nemaline myopathy type 3 (NEM3) [MIM:161800]. A form of nemaline myopathy. Nemaline myopathies are muscular disorders characterized by muscle weakness of varying severity and onset, and abnormal thread-or rod-like structures in muscle fibers on histologic examination. The phenotype at histological level is variable. Some patients present areas devoid of oxidative activity containg (cores) within myofibers. Core lesions are unstructured and poorly circumscribed.

Defects in ACTA1 are a cause of myopathy, actin, congenital, with excess of thin myofilaments (MPCETM) [MIM:161800]. A congenital muscular disorder characterized at histological level by areas of sarcoplasm devoid of normal myofibrils and mitochondria, and replaced with dense masses of thin filaments. Central cores, rods, ragged red fibers, and necrosis are absent. Defects in ACTA1 are a cause of congenital myopathy with fiber-type disproportion (CFTD) [MIM:255310]; also known as congenital fiber-type disproportion myopathy (CFTDM). CFTD is a genetically heterogeneous disorder in which there is relative hypotrophy of type 1 muscle fibers

compared to type 2 fibers on skeletal muscle biopsy. However, these findings are not specific and can be found in many different myopathic and neuropathic conditions.

序列相似性 Belongs to the actin family.

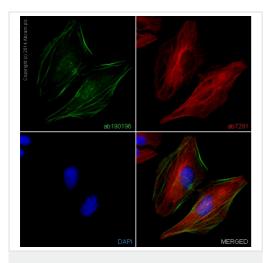
翻译后修饰 Oxidation of Met-46 by MICALs (MICAL1, MICAL2 or MICAL3) to form methionine sulfoxide

 $promotes\ actin\ filament\ depolymerization.\ Methionine\ sulfoxide\ is\ produced\ stereospecifically,\ but$

it is not known whether the (S)-S-oxide or the (R)-S-oxide is produced.

细胞定位 Cytoplasm > cytoskeleton.

图片



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 488 Anti-muscle Actin antibody [EPR8484] (ab190196) ab190196 staining Actin in HeLa cells. The cells were fixed with 100% methanol (5min) and then blocked in 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated with ab190196 at 1/50 dilution and ab7291 at 1 μ g/ml overnight at +4°C, followed by a further incubation at room temperature for 1h with a goat anti-mouse AlexaFluor 594 (ab150120) at 2 μ g/ml (shown in red). Nuclear DNA was labelled in blue with DAPI.



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