abcam

Product datasheet

HRP Anti-muscle Actin antibody [EPR8484] - Loading Control ab185058

Recombinant RabMAb

* ★ ★ ★ ★ ↑ 1 Abreviews 2 References 2 Images

Overview

Product name HRP Anti-muscle Actin antibody [EPR8484] - Loading Control

Description HRP Rabbit monoclonal [EPR8484] to muscle Actin - Loading Control

Host species Rabbit

Conjugation HRP

Tested applications Suitable for: WB

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Synthetic peptide within Human muscle Actin aa 350-450. The exact immunogen 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

Positive control NIH 3T3 cell lysate; Human fetal artery, kidney and heart lisates; Human uterus, stomach and

skeletal muscle lysates; Human skeletal muscle tissue; Human smooth muscle in colon tissue;

Human heart muscle tissue.

General notesThis 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 $\underline{\text{see here}}$.

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

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.

Avoid freeze / thaw cycle. Store In the Dark.

1

Storage buffer pH: 7.40

Preservative: 0.1% Proclin 300 Solution

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

Purity Protein A purified

Clonality Monoclonal
Clone number EPR8484

Isotype IgG

Applications

The Abpromise guarantee

Our Abpromise guarantee covers the use of ab185058 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
WB	★★★★☆(1)	1/10000. Detects a band of approximately 42 kDa (predicted molecular weight: 42 kDa).

Target

Function

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

Involvement in disease

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.

Sequence similarities

Belongs to the actin family.

Post-translational modifications

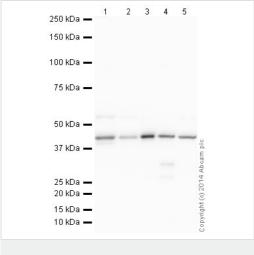
Oxidation of Met-46 by MICALs (MICAL1, MICAL2 or MICAL3) to form methionine sulfoxide promotes actin filament depolymerization. Methionine sulfoxide is produced stereospecifically, but

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

Cellular localization

Cytoplasm > cytoskeleton.

Images



Western blot - HRP Anti-muscle Actin antibody
[EPR8484] - Loading Control (ab185058)

All lanes : HRP Anti-muscle Actin antibody [EPR8484] - Loading Control (ab185058) at 1/10000 dilution

Lane 1 : NIH 3T3 (Mouse embryonic fibroblast cell line) Whole Cell Lysate at 20 μg

Lane 2 : Human kidney tissue lysate - total protein (ab30203) at 20 µq

Lane 3: Human Fetal Heart at 5 µg

Lane 4: Human skeletal muscle tissue lysate - total protein

(ab29330) at 10 µg

Lane 5: Skeletal Muscle (Rat) Tissue Lysate at 10 µg

Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 42 kDa

Additional bands at: 42 kDa. We are unsure as to the identity of

these extra bands.

Exposure time: 5 seconds

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 5% Bovine Serum Albumin before being incubated with ab185058 overnight at 4°C. Antibody binding was visualised using ECL development solution ab133406







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