abcam

Product datasheet

Anti-Myosin light chain 3 antibody [EPR4161] ab108516

Recombinant RabMAb

2 Images

Overview

Product name Anti-Myosin light chain 3 antibody [EPR4161]

Description Rabbit monoclonal [EPR4161] to Myosin light chain 3

Host species Rabbit

Suitable for: WB **Tested applications**

Unsuitable for: Flow Cyt,ICC/IF,IHC-P or IP

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control Human skeletal muscle, Human heart, Mouse heart, Mouse liver, and Rat heart lysates

General notes 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**.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

Storage buffer pH: 7.2

Preservative: 0.05% Sodium azide

Constituents: 0.1% BSA, 40% Glycerol (glycerin, glycerine), 9.85% Tris glycine, 50% Tissue

culture supernatant

Purity Protein A purified

Clonality Monoclonal Clone number **EPR4161**

Isotype ΙgG

Applications

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab108516 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		1/1000 - 1/10000. Predicted molecular weight: 22 kDa.

Application notes

Is unsuitable for Flow Cyt,ICC/IF,IHC-P or IP.

Target

Function

Regulatory light chain of myosin. Does not bind calcium.

Involvement in disease

Defects in MYL3 are the cause of cardiomyopathy familial hypertrophic type 8 (CMH8) [MIM:608751]. Familial hypertrophic cardiomyopathy is a hereditary heart disorder characterized by ventricular hypertrophy, which is usually asymmetric and often involves the interventricular septum. The symptoms include dyspnea, syncope, collapse, palpitations, and chest pain. They can be readily provoked by exercise. The disorder has inter- and intrafamilial variability ranging from benign to malignant forms with high risk of cardiac failure and sudden cardiac death. CMH8 inheritance can be autosomal dominant or recessive.

Defects in MYL3 are the cause of cardiomyopathy familial hypertrophic with mid-left ventricular chamber type 1 (MVC1) [MIM:608751]. MVC1 is a very rare variant of familial hypertrophic cardiomyopathy, characterized by mid-left ventricular chamber thickening.

Sequence similarities

Contains 3 EF-hand domains.

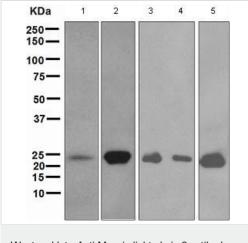
Post-translational

The N-terminus is blocked.

modifications

N-terminus is methylated by METTL11A/NTM1.

Images



Western blot - Anti-Myosin light chain 3 antibody [EPR4161] (ab108516)

All lanes: Anti-Myosin light chain 3 antibody [EPR4161]

(ab108516) at 1/1000 dilution

Lane 1: Human skeletal muscle lysate

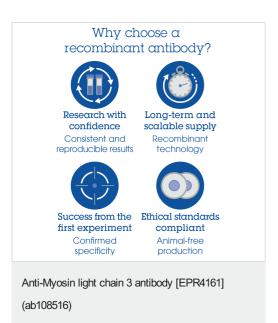
Lane 2 : Human heart lysate
Lane 3 : Mouse heart lysate

Lane 4: Mouse liver lysate

Lane 5: Rat heart lysate

Lysates/proteins at 10 µg per lane.

Predicted band size: 22 kDa



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

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