abcam

Product datasheet

Anti-MYH2 antibody [EPR5280] ab124937

Recombinant RabMAb

7 References 4 Images

Overview

Product name Anti-MYH2 antibody [EPR5280]

Description Rabbit monoclonal [EPR5280] to MYH2

Host species Rabbit

Suitable for: WB **Tested applications**

Unsuitable for: IHC-P or IP

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Synthetic peptide within Human MYH2 aa 50-150. The exact sequence is proprietary.

Positive control Mouse muscle, Rat muscle and Human trachea 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.

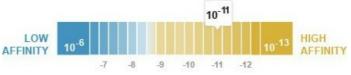
Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

Properties

Form Liquid

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

 $K_D = 6.90 \times 10^{-11} M$ Dissociation constant (K_D)



Learn more about K_D

Storage buffer pH: 7.2

Preservative: 0.05% Sodium azide

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

supernatant

Purity Protein A purified

Clonality Monoclonal
Clone number EPR5280

Isotype IgG

Applications

The Abpromise quarantee Our Abpromise quarantee covers the use of ab124937 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		1/10000 - 1/50000. Detects a band of approximately 220 kDa (predicted molecular weight: 223 kDa).

Application notes Is unsuitable for IHC-P or IP.

Target

Function Muscle contraction. Required for cytoskeleton organization.

Involvement in disease Defects in MYH2 are the cause of inclusion body myopathy type 3 (IBM3) [MIM:605637].

Hereditary inclusion body myopathies constitute a group of neuromuscular disorders

characterized by slowly progressive distal and proximal weakness and a typical muscle pathology including rimmed vacuoles and filamentous inclusions. IBM3 is a variant of hereditary inclusion body myopathies and is characterized by autosomal dominant myopathy with joint contracture, ophthalmoplegia and rimmed vacuoles. Morphological analysis of muscle biopsies from patients indicate that the type 2A fibers frequently were abnormal, whereas other fiber types appeared

normal.

Sequence similarities Contains 1 IQ domain.

Contains 1 myosin head-like domain.

Domain The rodlike tail sequence is highly repetitive, showing cycles of a 28-residue repeat pattern

composed of 4 heptapeptides, characteristic for alpha-helical coiled coils.

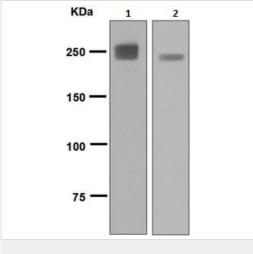
Each myosin heavy chain can be split into 1 light meromyosin (LMM) and 1 heavy meromyosin

(HMM). It can later be split further into 2 globular subfragments (S1) and 1 rod-shaped

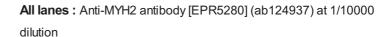
subfragment (S2).

Cellular localization Cytoplasm > myofibril. Thick filaments of the myofibrils.

Images



Western blot - Anti-MYH2 antibody [EPR5280] (ab124937)



Lane 1 : Mouse muscle tissue lysate

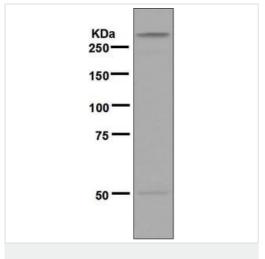
Lane 2 : Rat muscle tissue lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes: HRP labelled Goat anti Rabbit lgG at 1/2000 dilution

Predicted band size: 223 kDa **Observed band size:** 220 kDa



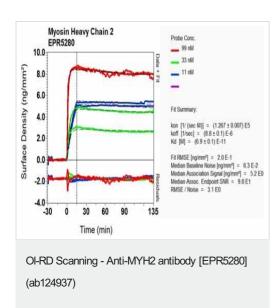
Western blot - Anti-MYH2 antibody [EPR5280] (ab124937)

Anti-MYH2 antibody [EPR5280] (ab124937) at 1/10000 dilution + Human trachea lysate at 10 μg

Secondary

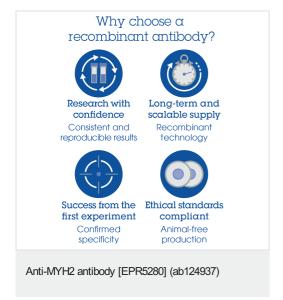
HRP labelled Goat anti Rabbit lgG at 1/2000 dilution

Predicted band size: 223 kDa **Observed band size:** 220 kDa



Equilibrium disassociation constant (K_D) Learn more about K_D

Click here to learn more about K_D



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