abcam

Product datasheet

Anti-MYL9 antibody [EPR13013(2)(B)] ab187152





重组 RabMAb

1 References 6 图像

概述

产品名称 Anti-MYL9抗体[EPR13013(2)(B)]

描述 兔单克隆抗体[EPR13013(2)(B)] to MYL9

宿主 Rabbit

适用于: WB, IHC-P 经测试应用

种属反应性 与反应: Mouse, Human

预测可用于: Rat 📤

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

阳性对照 WB: HeLa and human colon cell lysate; Human stomach, colon and uterus lysates; Mouse muscle

lysate. IHC-P: Human colon tissue; Mouse stomach tissue.

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.

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 long

term. Avoid freeze / thaw cycle.

存储溶液 pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

纯度 Protein A purified

克隆 单克隆

克隆编号 EPR13013(2)(B)

应用

The Abpromise guarantee Abpromise™承诺保证使用ab187152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
WB		1/1000 - 1/10000. Predicted molecular weight: 20 kDa.
IHC-P		1/250 - 1/500. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

靶标

功能 Myosin regulatory subunit that plays an important role in regulation of both smooth muscle and

nonmuscle cell contractile activity via its phosphorylation. Implicated in cytokinesis, receptor

capping, and cell locomotion.

组织特异性 Smooth muscle tissues and in some, but not all, nonmuscle cells.

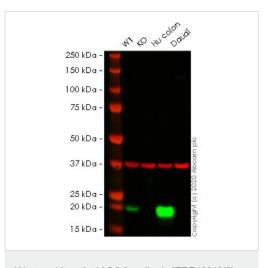
序列相似性 Contains 3 EF-hand domains.

翻译后修饰 Phosphorylation increases the actin-activated myosin ATPase activity and thereby regulates the

contractile activity. It is required to generate the driving force in the migration of the cells but not

necessary for localization of myosin-2 at the leading edge.

图片



Western blot - Anti-MYL9 antibody [EPR13013(2)

(B)] (ab187152)

All lanes : Anti-MYL9 antibody [EPR13013(2)(B)] (ab187152) at 1/1000 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: MYL9 knockout HeLa cell lysate

Lane 3: Human colon cell lysate

Lane 4: Daudi cell lysate

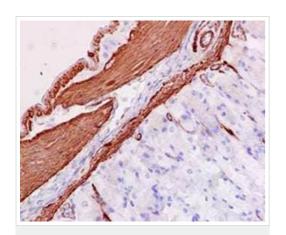
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 20 kDa

Lanes 1-4: Merged signal (red and green). Green - ab187152 observed at 20 kDa. Red - loading control **ab8245** observed at 37 kDa.

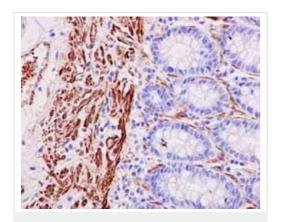
ab187152 Anti-MYL9 antibody [EPR13013(2)(B)] was shown to specifically react with MYL9 in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab266036 (knockout cell lysate ab256999) was used. Wild-type and MYL9 knockout samples were subjected to SDS-PAGE. ab187152 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-MYL9 antibody
[EPR13013(2)(B)] (ab187152)

Immunohistochemical analysis of paraffin-embedded mouse stomach tissue labeling MYL9 with ab187152 at 1/500 dilution. Anti-rabbit IgG HRP was used as a secondary antibody and slides were counterstained with hematoxylin.

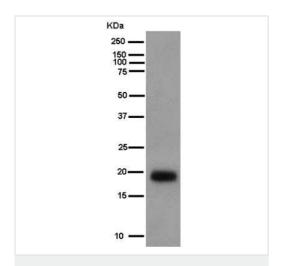
Perform heat mediated antigen retrieval with EDTA buffer pH 9 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-MYL9 antibody
[EPR13013(2)(B)] (ab187152)

Immunohistochemical analysis of paraffin-embedded human colon tissue labeling MYL9 with ab187152 at 1/500 dilution. Anti-rabbit IgG HRP was used as a secondary antibody and slides were counterstained with hematoxylin.

Perform heat mediated antigen retrieval with EDTA buffer pH 9 before commencing with IHC staining protocol.



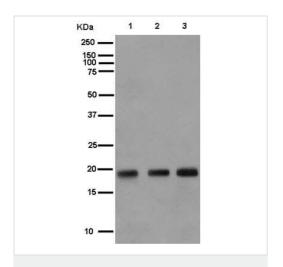
Western blot - Anti-MYL9 antibody [EPR13013(2) (B)] (ab187152)

Anti-MYL9 antibody [EPR13013(2)(B)] (ab187152) at 1/1000 dilution + Mouse muscle lysate at 10 μg

Secondary

Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 20 kDa



Western blot - Anti-MYL9 antibody [EPR13013(2) (B)] (ab187152)

All lanes : Anti-MYL9 antibody [EPR13013(2)(B)] (ab187152) at 1/5000 dilution

Lane 1: Human stomach lysate

Lane 2: Human colon lysate

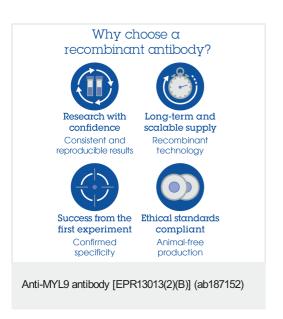
Lane 3: Human uterus lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/1000 dilution

Predicted band size: 20 kDa



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