

Anti-MYL9 Antibody

Catalog # AP53909

Product Information

ApplicationWB, IFPrimary AccessionP24844Other AccessionP19105

Reactivity Human, Mouse, Rat

Host Rabbit
Clonality Polyclonal
Calculated MW 19827

Additional Information

Gene ID 10398

Other Names MYL9; MLC2; MRLC1; MYRL2; Myosin regulatory light polypeptide 9; 20 kDa

myosin light chain; LC20; MLC-2C; Myosin RLC; Myosin regulatory light chain 2, smooth muscle isoform; Myosin regulatory light chain 9; Myosin regulatory light chain MRLC1; MYL12A; MLCB; MRLC3; RLC; Myosin regulatory light chain 12A; MLC-2B; Myosin RLC; Myosin regulatory light chain 2, nonsarcomeric;

Myosin regulatory light chain MRLC3

Target/Specificity Recognizes endogenous levels of MYL9 protein.

Dilution WB~~1/500 - 1/1000 IF~~1/50 - 1/200

Format Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

Protein Information

Name MYL9

Synonyms MLC2, MRLC1, MYRL2

Function 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 (PubMed: 11942626, PubMed: 2526655). In myoblasts, may regulate PIEZO1-dependent cortical actomyosin assembly involved in myotube

formation (By similarity).

Cellular Location Cytoplasm, cytoskeleton {ECO:0000250 | UniProtKB:Q9CQ19}. Cytoplasm, cell

cortex {ECO:0000250 | UniProtKB:Q9CQ19}. Note=Colocalizes with F-actin,

MYH9 and PIEZO1 at the actomyosin cortex in myoblasts {ECO:0000250|UniProtKB:Q9CQ19}

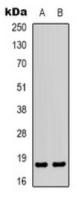
Tissue Location

Smooth muscle tissues and in some, but not all, nonmuscle cells.

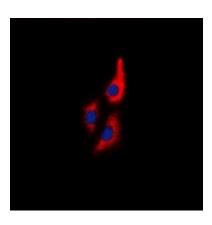
Background

Rabbit polyclonal antibody to MYL9

Images



Western blot analysis of MYL9 expression in Jurkat (A), COLO205 (B) whole cell lysates.



Immunofluorescent analysis of MYL9 staining in Jurkat cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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