



MYL2 Monoclonal Antibody

Catalog No	YP-mAb-03242
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	MYL2
Immunogen	Synthesized peptide derived from human MYL2. at AA range: 91-140
Specificity	MYL2 Monoclonal Antibody detects endogenous levels of MYL2
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	Cardiac myosin light chain-2;Cardiac ventricular myosin light chain 2;CMH1;MLC 2v;MLC-2;MLC-2v;MLC2;MLRV_HUMAN;MYL 2;MYL2;Myosin light chain 2 regulatory cardiac slow;Myosin light polypeptide 2 regulatory cardiac slow;Myosin regulatory light chain 2;Myosin regulatory light chain 2 ventricular/cardiac muscle isoform;Regulatory light chain of myosin;RLC of myosin;Slow cardiac myosin regulatory light chain 2;ventricular/cardiac muscle isoform;ventricular/cardiac muscle isoform (MLC-2) (MLC-2v); ventricular/cardiac muscle isoform (MLC-2) (MLC-2v)
Observed Band	18kD
Calculated Molecular Weight	18kD
Cell Pathway	Cytoplasm, myofibril, sarcomere, A band .
Tissue Specificity	Highly expressed in type I muscle fibers.
Function	disease:Defects in MYL2 are the cause of cardiomyopathy familial hypertrophic type 10 (CMH10) [MIM:608758]. 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.,disease:Defects in MYL2 are the cause of cardiomyopathy hypertrophic with mid-left ventricular chamber type 2 (MVC2) [MIM:608758]. MVC2 is a very rare variant of familial hypertrophic cardiomyopathy, characterized by mid-left ventricular chamber thickening., miscellaneous:This chain binds calcium.,similarity:Contains 3 EF-hand doma

Background

Thus gene encodes the regulatory light chain associated with cardiac myosin beta (or slow) heavy chain. Ca⁺ triggers the phosphorylation of regulatory light chain that in turn triggers contraction. Mutations in this gene are associated with mid-left ventricular chamber type hypertrophic cardiomyopathy. [provided by RefSeq, Jul 2008],

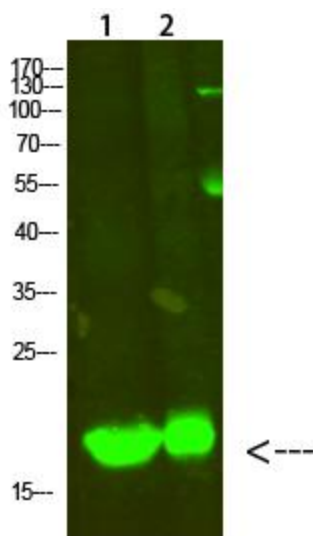
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western Blot analysis of various cells using MYL2 Monoclonal Antibody