

Rabbit Anti-Myozenin 2 antibody

SL18975R

Muozenin 2
III内结己性蛋白MVO72结体
加风行开任度口MIOZZ加体
CAOTTS; Calcineurin binding protein calsarcin 1; Calsarci
CMH16; CS 1; CS1; FA1Z related protein 2; FA1Z-related protein 2; Muscle specific
protein; MYOZ 2; MYOZ2; MYOZ2 HUMAN; Myozenin-2; Myozenin2.
Rabbit
Polyclonal
Human,Mouse,Rat,Dog,Pig,Cow,Horse,Rabbit,Sheep,
WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=1:100-
500IF=1:100-500 (Paraffin sections need antigen repair)
not yet tested in other applications.
optimal dilutions/concentrations should be determined by the end user.
30kDa
cytoplasmic
Lyophilized or Liquid
1mg/ml
KLH conjugated synthetic peptide derived from human Myozenin 2:1-100/264
IgG
affinity purified by Protein A
0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized
antibody is stable at room temperature for at least one month and for greater than a year
when kept at -20°C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of
antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed
The protein encoded by this gene belongs to a family of sarcomeric proteins that bind to
calcineurin, a phosphatase involved in calcium-dependent signal transduction in diverse
cell types. These family members tether calcineurin to alpha-actinin at the z-line of the
sarcomere of cardiac and skeletal muscle cells, and thus they are important for
calcineurin signaling. Mutations in this gene cause cardiomyopathy familial

hypertrophic type 16, a hereditary heart disorder. [provided by RefSeq, Aug 2011]

Function:

Myozenins may serve as intracellular binding proteins involved in linking Z line proteins such as alpha-actinin, gamma-filamin, TCAP/telethonin, LDB3/ZASP and localizing calcineurin signaling to the sarcomere. Plays an important role in the modulation of calcineurin signaling. May play a role in myofibrillogenesis.

Subcellular Location:

Cytoplasm > myofibril > sarcomere > Z line. Colocalizes with ACTN1 and PPP3CA at the Z-line of heart and skeletal muscle.

Tissue Specificity: Expressed specifically in heart and skeletal muscle.

DISEASE:

Defects in MYOZ2 are the cause of familial hypertrophic cardiomyopathy type 16 (CMH16) [MIM:613838]. CMH16 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.

Similarity: Belongs to the myozenin family.

SWISS: Q9NPC6

Gene ID: 51778

Database links:

Entrez Gene: 51778 Human

Entrez Gene: 540487 Cow

Entrez Gene: 59006 Mouse

Entrez Gene: 295426 Rat

Entrez Gene: 100526817 Sheep

<u>Omim: 605602</u> Human

	SwissProt: Q5E9V3 Cow
	SwissProt: Q9NPC6 Human
	SwissProt: Q9JJW5 Mouse
	Unigene: 732122 Human
	Unigene: 141157 Mouse
	Unigene: 12931 Rat
Picture:	Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications. 100 treat 100 5 63 63 63 63 63 63 63 63 63 63 63 63 63
	Sample: Heart (Mouse) Lysate at 40 ug
	Primary: Anti- Myozenin 2 (SL18975R) at 1/300 dilution
	Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution

	Predicted band size: 30kD
	Observed band size: 30kD

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