

# Produktinformation



Forschungsprodukte & Biochemikalien
Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

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Lieferung & Zahlungsart siehe unsere Liefer- und Versandbedingungen

## Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

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#### SANTA CRUZ BIOTECHNOLOGY, INC.

## MYLK2 (m): 293 Lysate: sc-178974



#### BACKGROUND

Myosin, the major component of thick muscle filaments, is a long asymmetric molecule containing a globular head and a long tail. Activation of smooth and cardiac muscle primarily involves pathways which increase calcium and Myosin phosphorylation resulting in contraction. Myosin in vertebrate striated muscle is composed of two heavy chains and four light chains. There are two distinct types of light chains: the phosphorylatable, regulatory or MLC2 type; and the nonphosphorylatable, alkali or MLC1 and MLC3 types. Myosin light chain phosphatase acts to regulate muscle contraction by dephosphorylating activated Myosin light chain. The role of Myosin alkali light chains in vertebrate skeletal muscle is poorly understood, although alkali light chains in smooth muscle may be involved with the active site of Myosin. Several isoforms of Myosin alkali light chains have been identified; each is associated with different muscle types and is encoded by a family of Myosin light chain genes. Human Myosin light chain has application as a cardiac marker. The human MLC1 gene maps to the same region in which the IDH1 (isocitrate dehydrogenase) gene is located. The MLC1 locus is closely linked to IDH1 on chromosome 1 in mouse, thus indicating that this is a conserved linkage.

#### REFERENCES

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- 2. Seidel, U., et al. 1987. The complete nucleotide sequences of cDNA clones coding for human Myosin light chains 1 and 3. Nucleic Acids Res. 15: 4989.
- 3. Cohen-Haguenauer, O., et al. 1988. Assignment of the human fast skeletal muscle Myosin alkali light chains gene (MLC1F/MLC3F) to 2g 32.1-2gter. Hum. Genet. 78: 65-70.
- 4. Castaneda, F., et al. 1990. Perineal abscess after prostatic urethroplastv with balloon catheter: report of a case. Radiology 174: 49-50.
- 5. Katoh, H., et al. 1992. Development of an immunoradiometric assay kit for ventricular Myosin light chain I with monoclonal antibodies. Clin. Chem. 38: 170-171.
- 6. Sanbe, A., et al. 1999. Abnormal cardiac structure and function in mice expressing nonphosphorylatable cardiac regulatory Myosin light chain 2. J. Biol. Chem. 274: 21085-21094.
- 7. Davis, J.S., et al. 2001. The overall pattern of cardiac contraction depends on a spatial gradient of Myosin regulatory light chain phosphorylation. Cell 107: 631-641.
- 8. Yamashita, H., et al. 2003. Myosin light chain isoforms modify force-generating ability of cardiac Myosin by changing the kinetics of actin-myosin interaction. Cardiovasc. Res. 60: 580-588.
- 9. Bicer, S. and Reiser, P.J. 2003. Myosin light chain 1 isoforms in slow fibers from global and orbital layers of canine rectus muscles. Invest. Ophthalmol. Vis. Sci 45: 138-143.

#### **STORAGE**

Store at -20° C. Repeated freezing and thawing should be minimized. Sample vial should be boiled once prior to use. Non-hazardous. No MSDS required.

#### CHROMOSOMAL LOCATION

Genetic locus: Mylk2 (mouse) mapping to 2 H1.

#### PRODUCT

MYLK2 (m): 293 Lysate represents a lysate of mouse MYLK2 transfected 293 cells and is provided as 100 µg protein in 200 µl SDS-PAGE buffer.

#### **APPLICATIONS**

MYLK2 (m): 293 Lysate is suitable as a Western Blotting positive control for mouse reactive MYLK2 antibodies. Recommended use: 10-20 µl per lane.

Control 293 Lysate: sc-110760 is available as a Western Blotting negative control lysate derived from non-transfected 293 cells.

#### **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

#### **PROTOCOLS**

See our web site at www.scbt.com for detailed protocols and support products.