SANTA CRUZ BIOTECHNOLOGY, INC.

MYBPC3 (H-120): sc-67354

SANTA CRUZ BIOTECHNOLOGY

The Power to Question

BACKGROUND

MYBPC3 (myosin-binding protein C, cardiac) encodes the cardiac isoform of the thick-filament Myosin-binding protein C. It is found in the crossbridge-bearing zone (C region) of A bands in vertebrate striated muscle. Regulatory phosphorylation of MYBPC3 by cAMP-dependent protein kinase (PKA) upon adrenergic stimulation may be linked to modulation of cardiac contraction. MYBPC3 binds F-Actin, MHC and native thin filaments, and modifies the activity of Actin-activated Myosin ATPase. Mutations in the MYBPC3 gene lead mainly to truncation of the protein, which results in one cause of familial hypertrophic cardiomyopathy type 4 (CMH4), a heart disorder characterized by ventricular hypertrophy, which often involves the interventricular septum and is usually asymmetric. The MYBPC3 gene maps to chromosome 11p11.2.

REFERENCES

1. Gautel, M., et al. 1995. Phosphorylation switches specific for the cardiac isoform of myosin-binding protein C: a modulator of cardiac contraction? EMBO J. 14: 1952-1960.
2. Bonne, G., et al. 1996. Cardiac myosin-binding protein C gene splice acceptor site mutation is associated with familial hypertrophic cardiomyopathy. Nat. Genet. 11: 438-440.
3. Carrier, L., et al. 1997. Organization and sequence of human cardiac myosin-binding protein C gene (MYBPC3) and identification of mutations predicted to produce truncated proteins in familial hypertrophic cardiomyopathy. Circ. Res. 80: 427-434.
4. Yu, B., et al. 1998. Molecular pathology of familial hypertrophic cardiomyopathy caused in the cardiac myosin-binding protein C gene. J. Med. Genet. 35: 205-210.
5. Moolman-Smook, J.C., et al. 1998. Identification of a new missense mutation in cardiomyopathy. J. Med. Genet. 35: 253-254.
6. Niimura, H., et al. 1998. Mutations in the gene for cardiac myosin-binding protein C and late-onset familial hypertrophic cardiomyopathy. N. Engl. J. Med. 338: 1248-1257.
7. Moolman-Smook, J.C., et al. 1999. The origins of hypertrophic cardiomyopathy-causing mutations in two South African subpopulations: a unique profile of both independent and founder events. Am. J. Hum. Genet. 65: 1308-1320.

CHROMOSOMAL LOCATION

Genetic locus: MYBPC3 (human) mapping to 11p11.2; Mybpc3 (mouse) mapping to 2 E1.

SOURCE

MYBPC3 (H-120) is a rabbit polyclonal antibody raised against amino acids 1-120 mapping at the N-terminus of MYBPC3 of human origin.

PRODUCT

Each vial contains 200 µg IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

MYBPC3 (H-120) is recommended for detection of MYBPC3 of human and, to a lesser extent, mouse and rat origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

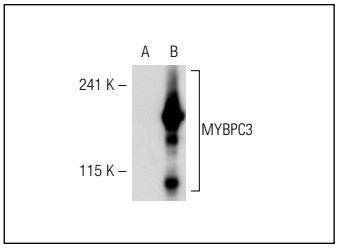
MYBPC3 (H-120) is also recommended for detection of MYBPC3 in additional species, including equine and porcine.

Suitable for use as control antibody for MYBPC3 siRNA (h): sc-61111, MYBPC3 siRNA (m): sc-61112, MYBPC3 shRNA Plasmid (h): sc-61111-SH, MYBPC3 shRNA Plasmid (m): sc-61112-SH, MYBPC3 shRNA (h) Lentiviral Particles: sc-61111-V and MYBPC3 shRNA (m) Lentiviral Particles: sc-61112-V.

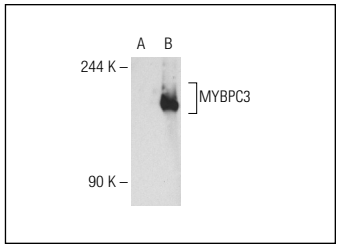
Molecular Weight of MYBPC3: 144 kDa.

Positive Controls: MYBPC3 (h2): 293T Lysate: sc-112857.

DATA



MYBPC3 (H-120): sc-67354. Western blot analysis of MYBPC3 expression in non-transfected: sc-117752 (**A**) and human MYBPC3 transfected: sc-112857 (**B**) 293T whole cell lysates.



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STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

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

PROTOCOLS

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