

## **Product datasheet for SC204472**

## MYBPC3 (NM 000256) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

Product Name: MYBPC3 (NM 000256) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: MYBPC3

Synonyms: CMD1MM; CMH4; cMyBP-C; FHC; LVNC10; MYBP-C

**ACCN:** NM\_000256

**Insert Size:** 367 bp

Insert Sequence: >SC204472 3'UTR clone of NM\_000256

The sequence shown below is from the reference sequence of NM\_000256. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

AAGTGCAGTCACAGAGAACTCA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

**Restriction Sites:** Sgfl-Mlul

OTI Disclaimer: Our molecular clone sequence data has been matched to the sequence identifier above as a

point of reference. Note that the complete sequence of this clone is largely the same as the

reference sequence but may contain minor differences, e.g., single nucleotide

polymorphisms (SNPs).

**Components:** The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

**RefSeg:** NM 000256.3



**OriGene Technologies, Inc.** 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



## MYBPC3 (NM\_000256) Human 3' UTR Clone - SC204472

Summary: MYBPC3 encodes the cardiac isoform of myosin-binding protein C. Myosin-binding protein C

is a myosin-associated protein found in the cross-bridge-bearing zone (C region) of A bands in striated muscle. MYBPC3, the cardiac isoform, is expressed exclussively in heart muscle. Regulatory phosphorylation of the cardiac isoform in vivo by cAMP-dependent protein kinase (PKA) upon adrenergic stimulation may be linked to modulation of cardiac contraction.

Mutations in MYBPC3 are one cause of familial hypertrophic cardiomyopathy. [provided by

RefSeq, Jul 2008]

**Locus ID:** 4607

**MW:** 13