

Product datasheet for SC201051

Myosin 8 (MYH8) (NM_002472) Human 3' UTR Clone

Product data:

Product Type:	3' UTR Clones
Product Name:	Myosin 8 (MYH8) (NM_002472) Human 3' UTR Clone
Symbol:	Myosin 8
Synonyms:	DA7; gtMHC-F; MyHC-peri; MyHC-pn
Mammalian Cell Selection:	Neomycin
Vector:	pMirTarget (PS100062)
ACCN:	NM_002472
Insert Size:	162 bp
Insert Sequence:	<p>>SC201051 3'UTR clone of NM_002472</p> <p>The sequence shown below is from the reference sequence of NM_002472. The complete sequence of this clone may contain minor differences, such as SNPs.</p> <p>Blue=Stop Codon Red=Cloning site</p> <pre> GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG TAACAATTGGCAGAGCTCAGAATTCAAACGATCGCC GAGGTTACACAAAAATCAGTGCAGAGTAAACACACCTGCCTGATGCTATCAAGAGGCTGAAGAAAGGC ACAAAATGTGCTATTTTGGTCACTTGCTTTATGACGTTTATTTCTGTAAAGCTGAATAAATAAAA ACTACAGTAAATGTATACATTA ACGCGTAAGCGGCCGCGCATCTAGATTGAAGAAAATGACCGACCAAGCGACGCCAACCTGCCATCA CGAGATTCGATTCCACCGCCGCTTCTATGAAAGG </pre>
Restriction Sites:	SgfI-MluI
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.
RefSeq:	NM_002472.3


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Summary:

Myosins are actin-based motor proteins that function in the generation of mechanical force in eukaryotic cells. Muscle myosins are heterohexamers composed of 2 myosin heavy chains and 2 pairs of nonidentical myosin light chains. This gene encodes a member of the class II or conventional myosin heavy chains, and functions in skeletal muscle contraction. This gene is predominantly expressed in fetal skeletal muscle. This gene is found in a cluster of myosin heavy chain genes on chromosome 17. A mutation in this gene results in trismus-pseudocamptodactyly syndrome. [provided by RefSeq, Sep 2009]

Locus ID:

4626

MW:

6.2