

Product datasheet for RC226697

MYH14 (NM_001145809) Human Tagged ORF Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	MYH14 (NM_001145809) Human Tagged ORF Clone
Tag:	Myc-DDK
Symbol:	MYH14
Synonyms:	DFNA4; DFNA4A; FP17425; MHC16; MYH17; myosin; NMHC-II-C; NMHC II-C; PNMHH
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
ORF Nucleotide Sequence:	>RC226697 representing NM_001145809 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC
GCC**CGATCGCC**

ATGGCAGCCGTGACCATGTCGGTGCCCGGGCGGAAGGCGCCCCCAGGCCGGGCCAGTGCCCGAGGCGG
CCCAGCCGTTCTGTTCACGCCCGCGGGCCAGCGGGGTGGCGGGCTGGCTCGGGCACCTCCCCGCA
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GAGATCTTCCAGCTGAACTCCTTCGAGCAGCTCTGCATCAACTACACCAACGAGAAGCTGCAGCAGCTCT
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ACGCGTACGCGGCCGCTCGAGCAGAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGATT
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Protein Sequence: >RC226697 representing NM_001145809
 Red=Cloning site Green=Tags(s)

MAAVTMSVPRKAPRPGVPVEAAQPFLFTPRGPSAGGGPGSGTSPQVEWTARRLVWVPELHGFEAAAL
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 FCVVINPYKQLPIYTEAIVEMYRGGKRHEVPPHYAVTEGAYRSMQDREDQSILCTGESGAGKTENTKK
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 VLQARAQELQKVQELQQQSAREVGELQGRVAQLEEERARLAEQLRAEAELCAEAEEETRGRLLAARKQEL
 VVSELEARVGEEECSRQMQTEKKRLQQHIQLEAHLEAEEGARQKLQLEKVTTEAKMKKFEEDLLLLLED
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 ARTKAEKQRRDLGEELEALRGELEDTL DSTNAQQELRSKREQEVTTELKKTLEEE TRIHEAAVQELRQRHG
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 EGEELELKAQMASAGQGKEEAVKQLRKMQAQMKELWREVEETRTSREEIFSQNRESEKRLKGLEAEVRL
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 LLQVESLTTLSAERSFSAKAESGRQQLERQIQELRGRGLEDAGARARHKMTIAALESKLAQAEQLEQ
 ETRERILSGKLVRRRAEKRLKEVVLQVEEERRVADQLRDQLEKGNLRVKQLKRQLEEAEEEA SRAQAGRRR
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 PPAHPQ

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Chromatograms: https://cdn.origene.com/chromatograms/mk8031_b08.zip

Restriction Sites: SgfI-MluI

Cloning Scheme:


ACCN: NM_001145809

ORF Size: 6108 bp

OTI Disclaimer: Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at custsupport@origene.com or by calling 301.340.3188 option 3 for pricing and delivery.

The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. [More info](#)

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.

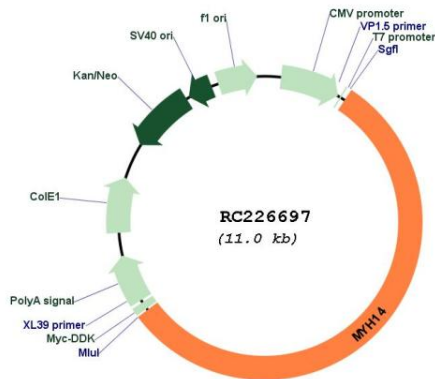
Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

- Reconstitution Method:**
1. Centrifuge at 5,000xg for 5min.
 2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.
 3. Close the tube and incubate for 10 minutes at room temperature.
 4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.
 5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.

RefSeq: [NM_001145809.2](#)
RefSeq ORF: 6111 bp
Locus ID: 79784
UniProt ID: [Q7Z406](#)
Cytogenetics: 19q13.33
Protein Pathways: Regulation of actin cytoskeleton, Tight junction, Viral myocarditis
MW: 231.8 kDa

Gene Summary: This gene encodes a member of the myosin superfamily. The protein represents a conventional non-muscle myosin; it should not be confused with the unconventional myosin-14 (MYO14). Myosins are actin-dependent motor proteins with diverse functions including regulation of cytokinesis, cell motility, and cell polarity. Mutations in this gene result in one form of autosomal dominant hearing impairment. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Dec 2011]

Product images:



Circular map for RC226697