

## Product datasheet for **SC337974**

### MYO6 (NM\_001300899) Human Untagged Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	MYO6 (NM_001300899) Human Untagged Clone
Tag:	Tag Free
Symbol:	MYO6
Synonyms:	DFNA22; DFNB37
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
Fully Sequenced ORF:	>NCBI ORF sequence for NM_001300899, the custom clone sequence may differ by one or more nucleotides

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ATGGAGGATGGAAAGCCCGTTTGGGCGCCACACCTACAGATGGATTTTCAGATGGCAATATTGTGGATA
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GCCAAGTGATCAACACTTTACATCTGCAGTTCACCAAAAAGCACAAGGATCATTTTCGACTCACTATTCCC  
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 TTAAGTAG

**Restriction Sites:**

Sgfl-Mlul

**ACCN:**

NM\_001300899

**OTI Disclaimer:**

Our molecular clone sequence data has been matched to the reference identifier above as a point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative RNA splicing form or single nucleotide polymorphism (SNP).

**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\\_001300899.1](#), [NP\\_001287828.1](#)

**RefSeq Size:** 8609 bp

**RefSeq ORF:** 3789 bp

**Locus ID:** 4646

**UniProt ID:** [Q9UM54](#)

**Cytogenetics:** 6q14.1

**Gene Summary:** This gene encodes a reverse-direction motor protein that moves toward the minus end of actin filaments and plays a role in intracellular vesicle and organelle transport. The protein consists of a motor domain containing an ATP- and an actin-binding site and a globular tail which interacts with other proteins. This protein maintains the structural integrity of inner ear hair cells and mutations in this gene cause non-syndromic autosomal dominant and recessive hearing loss. Alternative splicing results in multiple transcript variants encoding distinct isoforms. [provided by RefSeq, Jul 2014]

**Transcript Variant:** This variant (2) lacks two in-frame exons in the 3' coding region, compared to variant 1. It encodes isoform 2 which has a shorter C-terminus, compared to isoform 1.

**Sequence Note:** This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments.