

## **Product datasheet for SC202838**

## OriGene Technologies, Inc.

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## C19orf46 (SYNE4) (NM\_001039876) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

Product Name: C19orf46 (SYNE4) (NM\_001039876) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: SYNE4

Synonyms: C19orf46; DFNB76; KASH4; Nesp4

**ACCN:** NM\_001039876

**Insert Size:** 60 bp

Insert Sequence: >SC202838 3'UTR clone of NM\_001039876

The sequence shown below is from the reference sequence of NM\_001039876. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

AGCTATGTCAATGGTCTTCCCCCAGTCTGATGTGTGTAATAAATGGTCACTGTCAAAGGA

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.

**RefSeq:** <u>NM 001039876.3</u>





## C19orf46 (SYNE4) (NM\_001039876) Human 3' UTR Clone - SC202838

**Summary:** 

This gene is a member of the nesprin family of genes, that encode KASH (Klarsicht, Anc-1, Syne Homology) domain-containing proteins. In addition to the KASH domain, this protein also contains a coiled-coil and leucine zipper region, a spectrin repeat, and a kinesin-1 binding region. This protein localizes to the outer nuclear membrane, and is part of the linker of nucleoskeleton and cytoskeleton (LINC) complex in the nuclear envelope. LINC complexes are formed by SUN (Sad1, UNC-84)-KASH pairs, and are thought to mechanically couple nuclear components to the cytoskeleton. Mutations in this gene have been associated with progressive high-frequency hearing loss. The absence of this protein in mice also caused hearing loss, and changes in hair cell morphology in the ears. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Aug 2015]

**Locus ID:** 163183

MW: 2.1