

Product datasheet for TL302742

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Otoferlin (OTOF) Human shRNA Plasmid Kit (Locus ID 9381)

Product data:

Product Type: shRNA Plasmids

Product Name: Otoferlin (OTOF) Human shRNA Plasmid Kit (Locus ID 9381)

Locus ID: 9381

Synonyms: AUNB1; DFNB6; DFNB9; FER1L2; NSRD9

Vector: pGFP-C-shLenti (TR30023)

E. coli Selection: Chloramphenicol (34 ug/ml)

Mammalian Cell

Puromycin

Selection:

Format: Lentiviral plasmids

Components: OTOF - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 9381).

5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

RefSeq: NM 001287489, NM 004802, NM 194248, NM 194322, NM 194323, NM 194323.1,

NM 194323.2, NM 004802.1, NM 004802.2, NM 004802.3, NM 194322.1, NM 194248.2, NM 194248.1, NM 194248.2, NM 001287489.1, BC156051, BC156938, NM 004802.4,

NM 194322.3, NM 194248.3, NM 194323.3, NM 001287489.2

UniProt ID: Q9HC10

Summary: Mutations in this gene are a cause of neurosensory nonsyndromic recessive deafness,

DFNB9. The short form of the encoded protein has 3 C2 domains, a single carboxy-terminal transmembrane domain found also in the C. elegans spermatogenesis factor FER-1 and human dysferlin, while the long form has 6 C2 domains. The homology suggests that this protein may be involved in vesicle membrane fusion. Several transcript variants encoding

multiple isoforms have been found for this gene. [provided by RefSeq, Jul 2008]

shRNA Design: These shRNA constructs were designed against multiple splice variants at this gene locus. To

be certain that your variant of interest is targeted, please contact <u>techsupport@origene.com</u>. If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u>.







Performance Guaranteed:

OriGene guarantees that the sequences in the shRNA expression cassettes are verified to correspond to the target gene with 100% identity. One of the four constructs at minimum are guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must be used in comparison with the target-specific shRNA transfected samples.

For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).