

## **Product datasheet for TL308472**

## OriGene Technologies, Inc.

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## **USH1C Human shRNA Plasmid Kit (Locus ID 10083)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** USH1C Human shRNA Plasmid Kit (Locus ID 10083)

**Locus ID:** 10083

**Synonyms:** AIE-75; DFNB18; DFNB18A; NY-CO-37; NY-CO-38; PDZ-45; PDZ-73; PDZ-73/NY-CO-38; PDZ73;

PDZD7C; ush1cpst

**Vector:** pGFP-C-shLenti (TR30023)

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

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

**Components:** USH1C - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 10083).

5µg purified plasmid DNA per construct

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

RefSeq: NM 001297764, NM 005709, NM 153676, NR 123738, NM 005709.1, NM 005709.2,

NM 005709.3, NM 153676.1, NM 153676.2, NM 153676.3, NM 001297764.1, BC016057,

NM 001297764.2, NM 153676.4, NM 005709.4

UniProt ID: Q9Y6N9

**Summary:** This gene encodes a scaffold protein that functions in the assembly of Usher protein

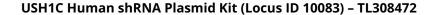
complexes. The protein contains PDZ domains, a coiled-coil region with a bipartite nuclear localization signal and a PEST degradation sequence. Defects in this gene are the cause of Usher syndrome type 1C and non-syndromic sensorineural deafness autosomal recessive type 18. Multiple transcript variants encoding different isoforms have been found for this

gene. [provided by RefSeq, Mar 2009]

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).