

## **Product datasheet for TR301038**

#### OriGene Technologies, Inc.

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### TMEM108 Human shRNA Plasmid Kit (Locus ID 66000)

#### **Product data:**

**Product Type:** shRNA Plasmids

**Product Name:** TMEM108 Human shRNA Plasmid Kit (Locus ID 66000)

**Locus ID:** 66000

Synonyms: CT124; RTLN

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Components: TMEM108 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID

= 66000). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

RefSeq: NM 001136469, NM 001282865, NM 023943, NM 023943.1, NM 023943.2, NM 023943.3,

NM 001136469.1, NM 001136469.2, NM 001282865.1, BC000568, BC000568.2, BM667108,

NM 023943.4

UniProt ID: Q6UXF1

**Summary:** Transmembrane protein required for proper cognitive functions. Involved in the development

of dentate gyrus (DG) neuron circuitry, is neccessary for AMPA receptors surface expression

and proper excitatory postsynaptic currents of DG granule neurons. Regulates the

organization and stability of the microtubule network of sensory neurons to allow axonal transport. Through the interaction with DST, mediates the docking of the dynein/dynactin motor complex to vesicle cargos for retrograde axonal transport. In hippocampal neurons, required for BDNF-dependent dendrite outgrowth. Cooperates with SH3GL2 and recruits the WAVE1 complex to facilitate actin-dependent BDNF:NTRK2 early endocytic trafficking and

mediate signaling from early endosomes.[UniProtKB/Swiss-Prot Function]

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).