

## **Product datasheet for TL502286**

## OriGene Technologies, Inc.

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## **Timeless Mouse shRNA Plasmid (Locus ID 21853)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Timeless Mouse shRNA Plasmid (Locus ID 21853)

**Locus ID:** 21853

**Synonyms:** C77407; Debt69; tim

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

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

**Mammalian Cell** 

Selection:

Puromycin

Format: Lentiviral plasmids

Components: Timeless - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID =

21853). 5µg purified plasmid DNA per construct

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

RefSeq: BC058641, BC082770, NM 001136082, NM 001164080, NM 001164081, NM 011589,

NM 001164080.1, NM 011589.1, NM 011589.2, NM 001164081.1, NM 001136082.1,

NM 001136082.2, BC026526, BC052884, BC064788

UniProt ID: Q9R1X4

**Summary:** The protein encoded by this gene is highly conserved and is involved in cell survival after

damage or stress, increase in DNA polymerase epsilon activity, maintenance of telomere length, and epithelial cell morphogenesis. The encoded protein also plays a role in the circadian rhythm autoregulatory loop, interacting with the PERIOD genes (PER1, PER2, and PER3) and others to downregulate activation of PER1 by CLOCK/ARNTL. Changes in this gene or its expression may promote prostate cancer, lung cancer, breast cancer, and mental disorders. Several transcript variants encoding different isoforms have been found for this

gene. [provided by RefSeq, Feb 2014]

**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 custom shRNA service.







## 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).