

## Product datasheet for TR302297

## OriGene Technologies, Inc.

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## PRDM10 Human shRNA Plasmid Kit (Locus ID 56980)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** PRDM10 Human shRNA Plasmid Kit (Locus ID 56980)

Locus ID:

PFM7; TRIS Synonyms:

pRS (TR20003) Vector:

E. coli Selection: Ampicillin Mammalian Cell

Selection:

Puromycin

Format: Retroviral plasmids

PRDM10 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = Components:

56980). 5µg purified plasmid DNA per construct

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

NM 020228, NM 199437, NM 199438, NM 199439, NM 199437.1, NM 199439.1, RefSeq:

> NM 020228.1, NM 020228.2, NM 199438.1, BC112934, BC112934.1, BC117415, BC143612, BC143613, NM 001367890, NM 001367894, NM 001367897, NM 001367891, NM 001367892,

NM 001367893, NM 001367895, NM 001367896, NM 001367898, NM 001367899,

NM 199439.2, NM 020228.3, NM 199438.2

**UniProt ID:** Q9NQV6

**Summary:** The protein encoded by this gene is a transcription factor that contains C2H2-type zinc-

> fingers. It also contains a positive regulatory domain, which has been found in several other zinc-finger transcription factors including those involved in B cell differentiation and tumor suppression. Studies of the mouse counterpart suggest that this protein may be involved in the development of the central nerve system (CNS), as well as in the pathogenesis of

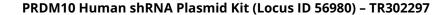
> neuronal storage disease. Multiple alternatively spliced transcript variants encoding distinct

isoforms have been observed. [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 techsupport@origene.com. 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).