

Product datasheet for **TL302292**

PRDM16 Human shRNA Plasmid Kit (Locus ID 63976)

Product data:

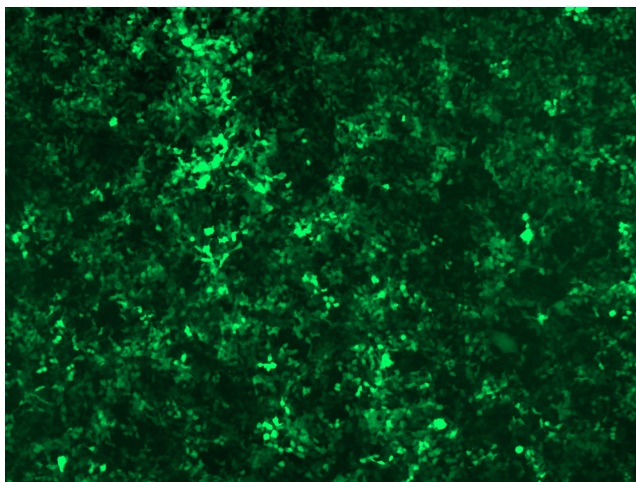
Product Type:	shRNA Plasmids
Product Name:	PRDM16 Human shRNA Plasmid Kit (Locus ID 63976)
Locus ID:	63976
Synonyms:	CMD1LL; KMT8F; LVNC8; MEL1; PFM13
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	PRDM16 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 63976). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	NM_022114 , NM_199454 , NM_199454.1 , NM_199454.2 , NM_022114.1 , NM_022114.2 , NM_022114.3 , BC110593 , BC161614 , BC168363 , NM_022114.4 , NM_199454.3
UniProt ID:	Q9HAZ2
Summary:	The reciprocal translocation t(1;3)(p36;q21) occurs in a subset of myelodysplastic syndrome (MDS) and acute myeloid leukemia (AML). This gene is located near the 1p36.3 breakpoint and has been shown to be specifically expressed in the t(1;3)(p36,q21)-positive MDS/AML. The protein encoded by this gene is a zinc finger transcription factor and contains an N-terminal PR domain. The translocation results in the overexpression of a truncated version of this protein that lacks the PR domain, which may play an important role in the pathogenesis of MDS and AML. Alternatively spliced transcript variants encoding distinct isoforms have been reported. [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 .


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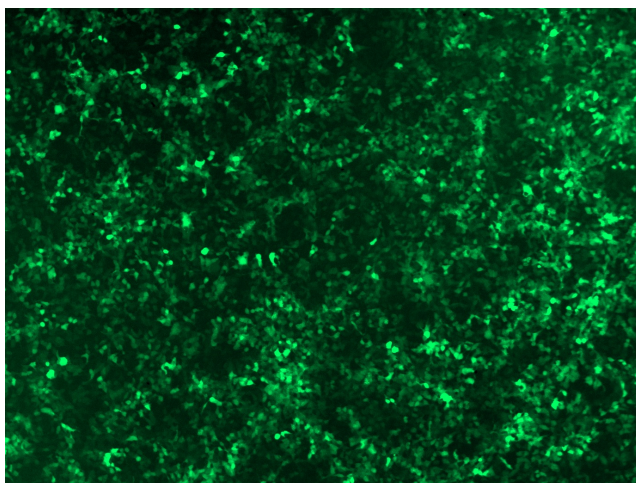
**Performance
Guaranteed:**

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.

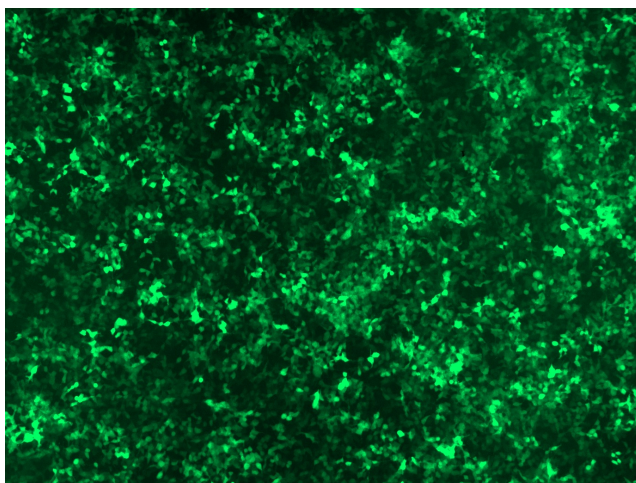
For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating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

Product images:


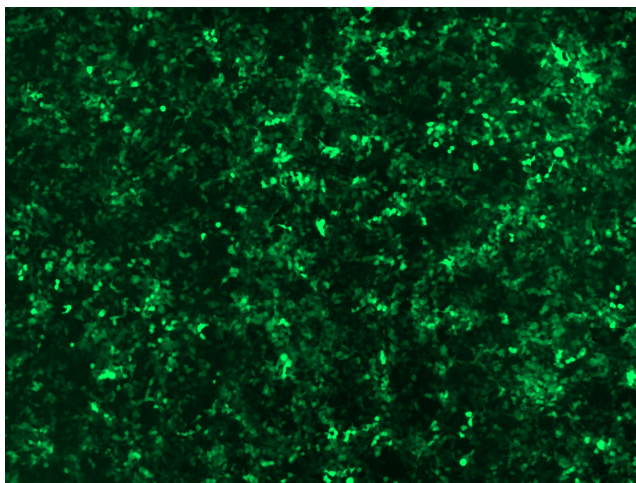
GFP signal was observed under microscope at 48 hours after transduction of TL302292A virus into HEK293 cells. TL302292A virus was prepared using lenti-shRNA TL302292A and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of TL302292B virus into HEK293 cells. TL302292B virus was prepared using lenti-shRNA TL302292B and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL302292C] virus into HEK293 cells. [TL302292C] virus was prepared using lenti-shRNA [TL302292C] and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL302292D] virus into HEK293 cells. [TL302292D] virus was prepared using lenti-shRNA [TL302292D] and [TR30037] packaging kit.