

Product datasheet for **TL511318V**

Smg1 Mouse shRNA Lentiviral Particle (Locus ID 233789)

Product data:

Product Type:	shRNA Lentiviral Particles
Product Name:	Smg1 Mouse shRNA Lentiviral Particle (Locus ID 233789)
Locus ID:	233789
Synonyms:	2610207I05Rik; 5430435M13Rik; C130002K18Rik; mKIAA0421
Vector:	pGFP-C-shLenti (TR30023)
Format:	Lentiviral particles
Components:	Smg1 - Mouse shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble control), 0.5 ml each, >10 ⁷ TU/ml.
RefSeq:	NM_001031814 , BC024431 , BC042594 , NM_177180
Summary:	<p>Serine/threonine protein kinase involved in both mRNA surveillance and genotoxic stress response pathways. Recognizes the substrate consensus sequence [ST]-Q. Plays a central role in nonsense-mediated decay (NMD) of mRNAs containing premature stop codons by phosphorylating UPF1/RENT1. Recruited by release factors to stalled ribosomes together with SMG8 and SMG9 (forming the SMG1C protein kinase complex), and UPF1 to form the transient SURF (SMG1-UPF1-eRF1-eRF3) complex. In EJC-dependent NMD, the SURF complex associates with the exon junction complex (EJC) through UPF2 and allows the formation of an UPF1-UPF2-UPF3 surveillance complex which is believed to activate NMD. Also acts as a genotoxic stress-activated protein kinase that displays some functional overlap with ATM. Can phosphorylate p53/TP53 and is required for optimal p53/TP53 activation after cellular exposure to genotoxic stress. Its depletion leads to spontaneous DNA damage and increased sensitivity to ionizing radiation (IR). May activate PRKCI but not PRKCZ (By similarity). [UniProtKB/Swiss-Prot Function]</p>
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 .

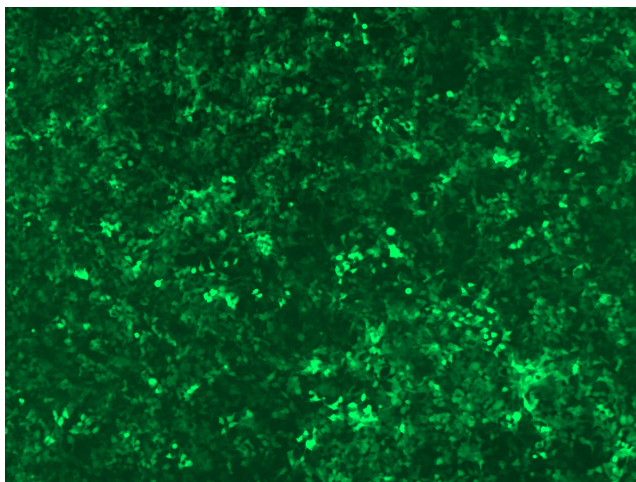


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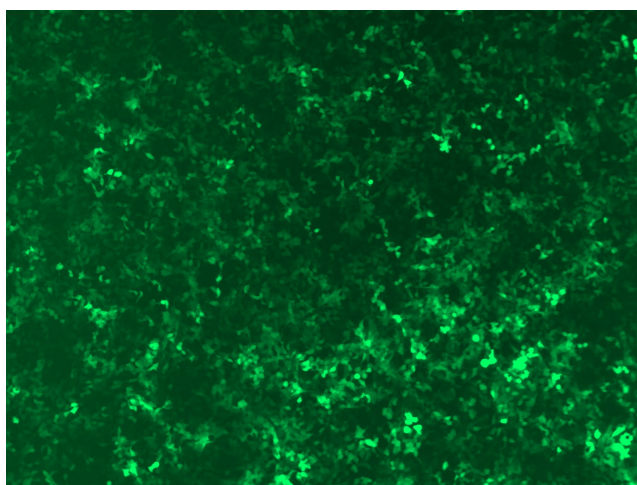
**Performance
Guaranteed:**

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.

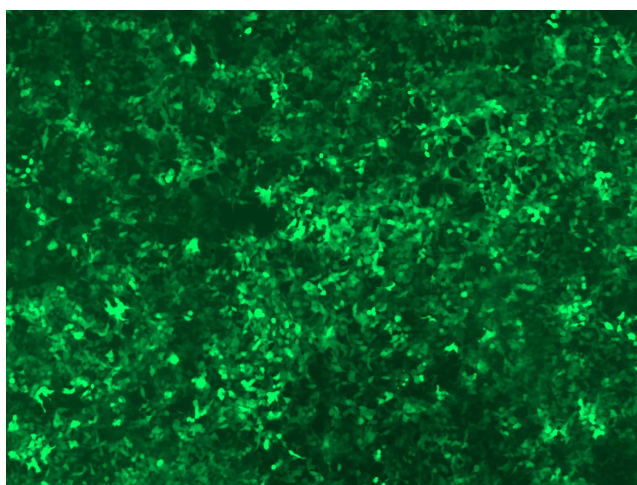
For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating 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 TL511318B virus into HEK293 cells. TL511318B virus was prepared using lenti-shRNA TL511318B and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL511318C] virus into HEK293 cells. [TL511318C] virus was prepared using lenti-shRNA [TL511318C] and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL511318D] virus into HEK293 cells. [TL511318D] virus was prepared using lenti-shRNA [TL511318D] and [TR30037] packaging kit.