

## Product datasheet for **TL308946V**

### TDP43 (TARDBP) Human shRNA Lentiviral Particle (Locus ID 23435)

#### Product data:

Product Type:	shRNA Lentiviral Particles
Product Name:	TDP43 (TARDBP) Human shRNA Lentiviral Particle (Locus ID 23435)
Locus ID:	23435
Synonyms:	ALS10; TDP-43
Vector:	pGFP-C-shLenti (TR30023)
Format:	Lentiviral particles
Components:	TARDBP - Human shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble control), 0.5 ml each, >10 <sup>7</sup> TU/ml.
RefSeq:	<a href="#">NM_007375</a> , <a href="#">NM_007375.1</a> , <a href="#">NM_007375.2</a> , <a href="#">NM_007375.3</a> , <a href="#">BC095435</a> , <a href="#">BC095435.1</a> , <a href="#">BC001487</a> , <a href="#">BC059955</a> , <a href="#">BC071657</a> , <a href="#">BM716882</a> , <a href="#">NM_007375.4</a>
UniProt ID:	<a href="#">Q13148</a>
Summary:	HIV-1, the causative agent of acquired immunodeficiency syndrome (AIDS), contains an RNA genome that produces a chromosomally integrated DNA during the replicative cycle. Activation of HIV-1 gene expression by the transactivator Tat is dependent on an RNA regulatory element (TAR) located downstream of the transcription initiation site. The protein encoded by this gene is a transcriptional repressor that binds to chromosomally integrated TAR DNA and represses HIV-1 transcription. In addition, this protein regulates alternate splicing of the CFTR gene. A similar pseudogene is present on chromosome 20. [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 <a href="mailto:techsupport@origene.com">techsupport@origene.com</a> . If you need a special design or shRNA sequence, please utilize our <a href="#">custom shRNA service</a> .

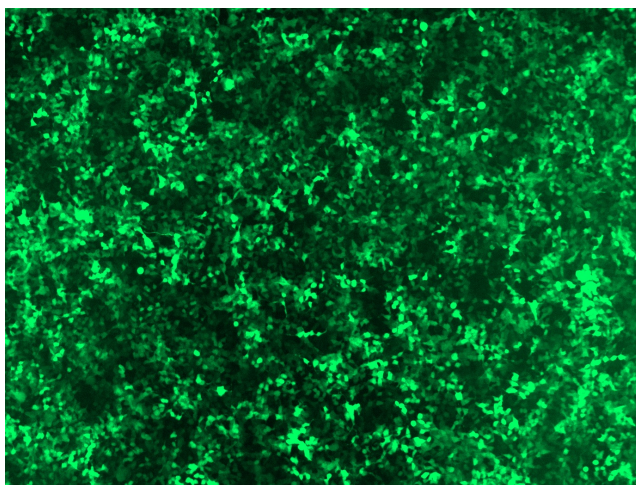


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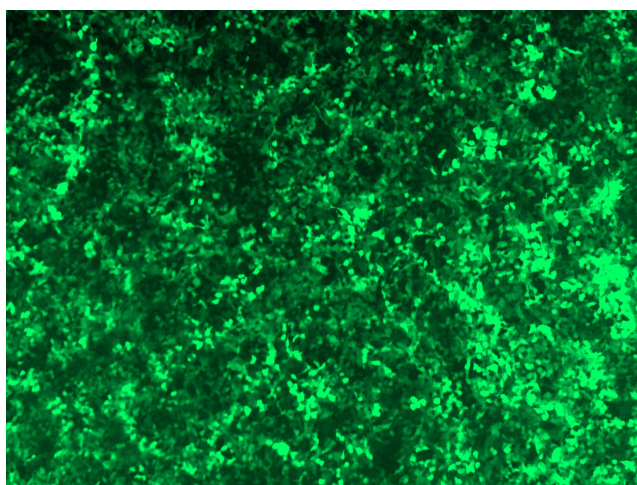
**Performance  
Guaranteed:**

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.

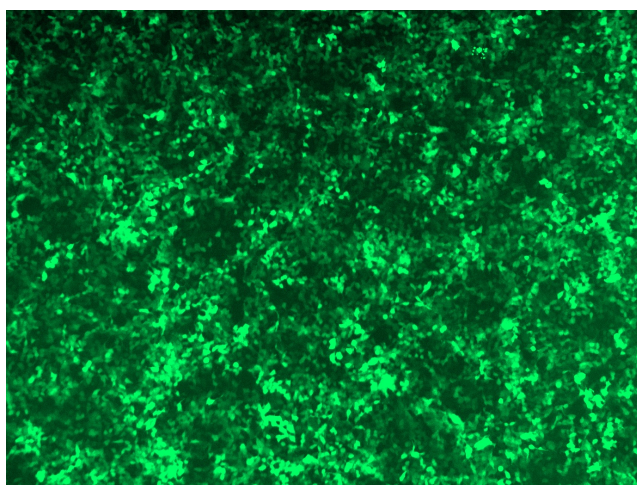
For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at [techsupport@origene.com](mailto:techsupport@origene.com). 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:**

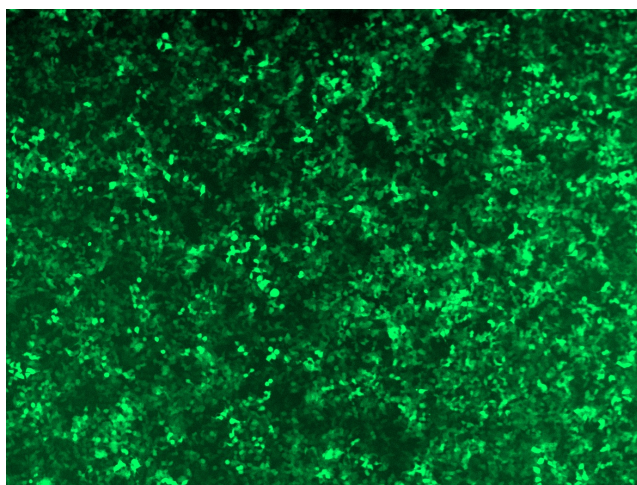
GFP signal was observed under microscope at 48 hours after transduction of TL308946A virus into HEK293 cells. TL308946A virus was prepared using lenti-shRNA TL308946A and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of TL308946B virus into HEK293 cells. TL308946B virus was prepared using lenti-shRNA TL308946B and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL308946C] virus into HEK293 cells. [TL308946C] virus was prepared using lenti-shRNA [TL308946C] and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL308946D] virus into HEK293 cells. [TL308946D] virus was prepared using lenti-shRNA [TL308946D] and [TR30037] packaging kit.