

## Product datasheet for **TL315488**

### PARP1 Human shRNA Plasmid Kit (Locus ID 142)

#### Product data:

Product Type:	shRNA Plasmids
Product Name:	PARP1 Human shRNA Plasmid Kit (Locus ID 142)
Locus ID:	142
Synonyms:	ADPRT; ADPRT 1; ADPRT1; ARTD1; pADPRT-1; PARP; PARP-1; PPOL
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	PARP1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 142). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	<a href="#">NM_001618</a> , <a href="#">NM_001618.1</a> , <a href="#">NM_001618.2</a> , <a href="#">NM_001618.3</a> , <a href="#">BC037545</a> , <a href="#">BC037545.1</a> , <a href="#">BC008660</a> , <a href="#">BC014206</a> , <a href="#">BC018620</a> , <a href="#">BC021045</a>
UniProt ID:	<a href="#">P09874</a>
Summary:	This gene encodes a chromatin-associated enzyme, poly(ADP-ribosyl)transferase, which modifies various nuclear proteins by poly(ADP-ribosyl)ation. The modification is dependent on DNA and is involved in the regulation of various important cellular processes such as differentiation, proliferation, and tumor transformation and also in the regulation of the molecular events involved in the recovery of cell from DNA damage. In addition, this enzyme may be the site of mutation in Fanconi anemia, and may participate in the pathophysiology of type I diabetes. [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> .

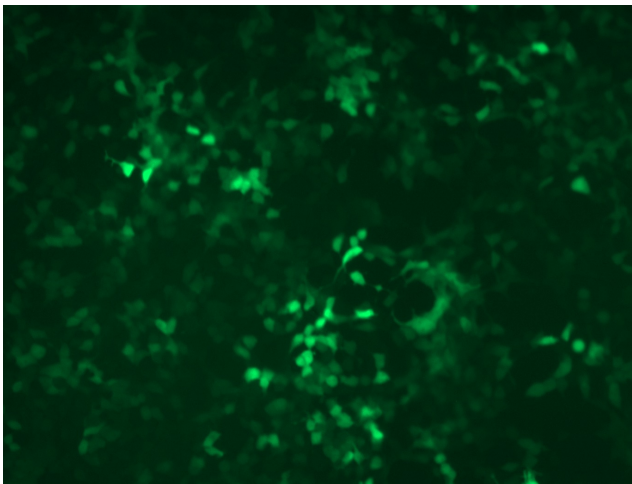


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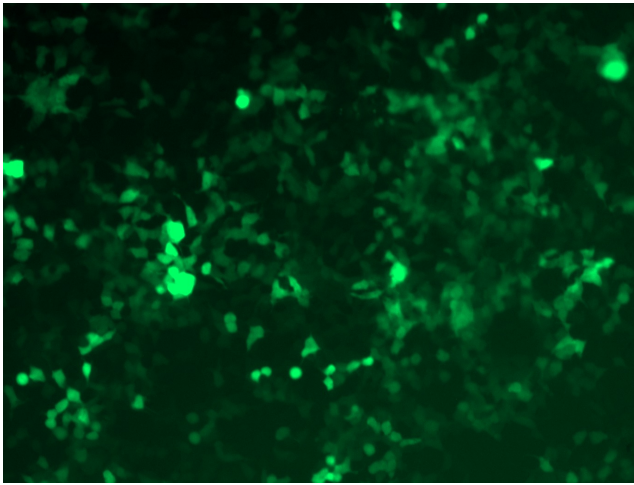
**Performance  
Guaranteed:**

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.

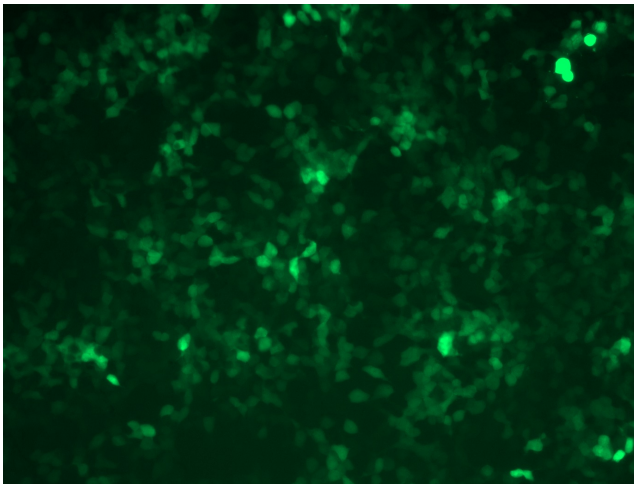
For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at [techsupport@origene.com](mailto:techsupport@origene.com).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

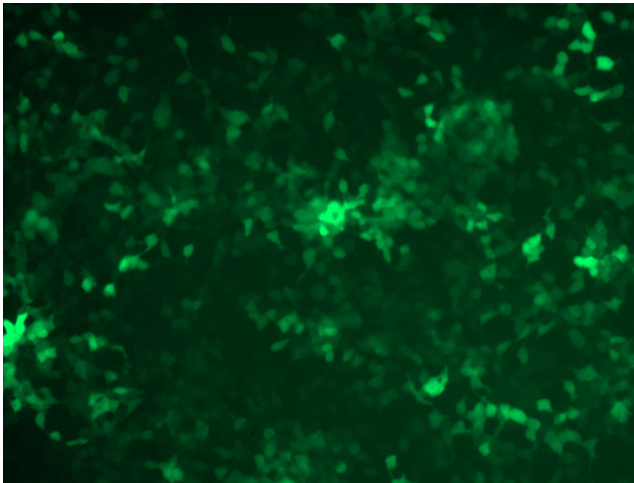
GFP signal was observed under microscope at 48 hours after transduction of TL315488A virus into HEK293 cells. TL315488A virus was prepared using lenti-shRNA TL315488A and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of TL315488B virus into HEK293 cells. TL315488B virus was prepared using lenti-shRNA TL315488B and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL315488C] virus into HEK293 cells. [TL315488C] virus was prepared using lenti-shRNA [TL315488C] and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL315488D] virus into HEK293 cells. [TL315488D] virus was prepared using lenti-shRNA [TL315488D] and [TR30037] packaging kit.