

Product datasheet for **TL316692**

RAD52 Human shRNA Plasmid Kit (Locus ID 5893)

Product data:

Product Type:	shRNA Plasmids
Product Name:	RAD52 Human shRNA Plasmid Kit (Locus ID 5893)
Locus ID:	5893
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	RAD52 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 5893). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	NM_001297419 , NM_001297420 , NM_001297421 , NM_001297422 , NM_002879 , NM_134422 , NM_134423 , NM_134424 , NR_123713 , NM_134424.1 , NM_134424.2 , NM_134424.3 , NM_001297420.1 , NM_001297422.1 , NM_001297421.1 , NM_001297419.1 , NM_134422.1 , BC042136 , BC104015 , BC104016 , BC104017 , BC114507 , BC114954 , BC144458 , BC160123 , NM_001297422.2 , NM_134424.4
UniProt ID:	P43351
Summary:	The protein encoded by this gene shares similarity with <i>Saccharomyces cerevisiae</i> Rad52, a protein important for DNA double-strand break repair and homologous recombination. This gene product was shown to bind single-stranded DNA ends, and mediate the DNA-DNA interaction necessary for the annealing of complementary DNA strands. It was also found to interact with DNA recombination protein RAD51, which suggested its role in RAD51 related DNA recombination and repair. A pseudogene of this gene is present on chromosome 2. Alternative splicing results in multiple transcript variants. Additional alternatively spliced transcript variants of this gene have been described, but their full-length nature is not known. [provided by RefSeq, Jul 2014]
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).