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Product datasheet for TR311735

DNA Ligase III (LIG3) Human shRNA Plasmid Kit (Locus ID 3980)

Product data:

Product Type:	shRNA Plasmids
Product Name:	DNA Ligase III (LIG3) Human shRNA Plasmid Kit (Locus ID 3980)
Locus ID:	3980
Synonyms:	LIG2; LIG3alpha
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	LIG3 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 3980). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	<u>NM 002311, NM 013975, NM 013975.1, NM 013975.2, NM 013975.3, NM 002311.1, NM 002311.2, NM 002311.3, NM 002311.4, BC068005, BC068005.1, BC009026, BM467239, BM726768, BM759573, NM 013975.4</u>
UniProt ID:	<u>P49916</u>
Summary:	This gene is a member of the DNA ligase family. Each member of this family encodes a protein that catalyzes the joining of DNA ends but they each have a distinct role in DNA metabolism. The protein encoded by this gene is involved in excision repair and is located in both the mitochondria and nucleus, with translation initiation from the upstream start codon allowing for transport to the mitochondria and translation initiation from a downstream start codon allowing for transport to the nucleus. Additionally, alternate transcriptional splice variants, encoding different isoforms, have been characterized. [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 <u>techsupport@origene.com</u> . If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u> .



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GRIGENE DNA Ligase III (LIG3) Human shRNA Plasmid Kit (Locus ID 3980) – TR311735

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).

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