

## Product datasheet for **TL311222**

### NDUFB9 Human shRNA Plasmid Kit (Locus ID 4715)

#### Product data:

Product Type:	shRNA Plasmids
Product Name:	NDUFB9 Human shRNA Plasmid Kit (Locus ID 4715)
Locus ID:	4715
Synonyms:	B22; CI-B22; LYRM3; MC1DN24; UQOR22
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	NDUFB9 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 4715). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	<a href="#">NM_001278645</a> , <a href="#">NM_001278646</a> , <a href="#">NM_001311168</a> , <a href="#">NM_005005</a> , <a href="#">NM_005005.1</a> , <a href="#">NM_005005.2</a> , <a href="#">NM_001278645.1</a> , <a href="#">NM_001278646.1</a> , <a href="#">BC007672</a> , <a href="#">BC007672.2</a> , <a href="#">NM_005005.3</a>
UniProt ID:	<a href="#">Q9Y6M9</a>
Summary:	The protein encoded by this gene is a subunit of the mitochondrial oxidative phosphorylation complex I (nicotinamide adenine dinucleotide: ubiquinone oxidoreductase). Complex I is localized to the inner mitochondrial membrane and functions to dehydrogenate nicotinamide adenine dinucleotide and to shuttle electrons to coenzyme Q. Complex I deficiency is the most common defect found in oxidative phosphorylation disorders and results in a range of conditions, including lethal neonatal disease, hypertrophic cardiomyopathy, liver disease, and adult-onset neurodegenerative disorders. Pseudogenes of this gene are found on chromosomes five, seven and eight. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jul 2015]
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).