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Product datasheet for TL319855

Cytochrome C Oxidase subunit VIb (COX6B1) Human shRNA Plasmid Kit (Locus ID 1340)

Product data:

Product Type:	shRNA Plasmids
Product Name:	Cytochrome C Oxidase subunit VIb (COX6B1) Human shRNA Plasmid Kit (Locus ID 1340)
Locus ID:	1340
Synonyms:	COX6B; COXG; COXVIb1; MC4DN7
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol
Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	COX6B1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 1340). 5µg purified plasmid DNA per construct Non-effective 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	NM_001863, NM_001863.2, NM_001863.3, NM_001863.4, BC001015, BC001015.2, BC002478, NM_001863.5
UniProt ID:	P14854
Summary:	Cytochrome c oxidase (COX), the terminal enzyme of the mitochondrial respiratory chain, catalyzes the electron transfer from reduced cytochrome c to oxygen. It is a heteromeric complex consisting of 3 catalytic subunits encoded by mitochondrial genes and multiple structural subunits encoded by nuclear genes. The mitochondrially-encoded subunits function in electron transfer, and the nuclear-encoded subunits may be involved in the regulation and assembly of the complex. This nuclear gene encodes subunit VIb. Mutations in this gene are associated with severe infantile encephalomyopathy. Three pseudogenes COX6BP-1, COX6BP-2 and COX6BP-3 have been found on chromosomes 7, 17 and 22q13.1-13.2, respectively. [provided by RefSeq, Jan 2010]
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).