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Product datasheet for TL300852

RNF86 (TRIM2) Human shRNA Plasmid Kit (Locus ID 23321)

Product data:

Product Type:	shRNA Plasmids
Product Name:	RNF86 (TRIM2) Human shRNA Plasmid Kit (Locus ID 23321)
Locus ID:	23321
Synonyms:	CMT2R; RNF86
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	TRIM2 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 23321). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	NM_001130067, NM_001302692, NM_001302693, NM_001302694, NM_015271, NM_001351054, NM_001351055, NM_001351056, NM_001351057, NM_015271.1, NM_015271.2, NM_015271.3, NM_015271.4, NM_001130067.1, NM_001302694.1, NM_001302693.1, NM_001302692.1, BC011052, BC011052.1, BC005016, BC019242, BC025417, NM_001130067.2
UniProt ID:	Q9C040
Summary:	The protein encoded by this gene is a member of the tripartite motif (TRIM) family. The TRIM motif includes three zinc-binding domains, a RING, a B-box type 1 and a B-box type 2, and a coiled-coil region. The protein localizes to cytoplasmic filaments. It plays a neuroprotective role and functions as an E3-ubiquitin ligase in proteasome-mediated degradation of target proteins. Mutations in this gene can cause early-onset axonal neuropathy. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Nov 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).