

## Product datasheet for **TL311101V**

### NRAS Human shRNA Lentiviral Particle (Locus ID 4893)

#### Product data:

Product Type:	shRNA Lentiviral Particles
Product Name:	NRAS Human shRNA Lentiviral Particle (Locus ID 4893)
Locus ID:	4893
Synonyms:	ALPS4; CMNS; N-ras; NCMS; NRAS1; NS6
Vector:	pGFP-C-shLenti (TR30023)
Format:	Lentiviral particles
Components:	NRAS - Human shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble control), 0.5 ml each, >10 <sup>7</sup> TU/ml.
RefSeq:	<a href="#">NM_002524</a> , <a href="#">NM_002524.1</a> , <a href="#">NM_002524.2</a> , <a href="#">NM_002524.3</a> , <a href="#">NM_002524.4</a> , <a href="#">BC005219</a> , <a href="#">BC005219.1</a> , <a href="#">BC013214</a> , <a href="#">NM_002524.5</a>
UniProt ID:	<a href="#">P01111</a>
Summary:	This is an N-ras oncogene encoding a membrane protein that shuttles between the Golgi apparatus and the plasma membrane. This shuttling is regulated through palmitoylation and depalmitoylation by the ZDHHC9-GOLGA7 complex. The encoded protein, which has intrinsic GTPase activity, is activated by a guanine nucleotide-exchange factor and inactivated by a GTPase activating protein. Mutations in this gene have been associated with somatic rectal cancer, follicular thyroid cancer, autoimmune lymphoproliferative syndrome, Noonan syndrome, and juvenile myelomonocytic leukemia. [provided by RefSeq, Jun 2011]
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).