

## **Product datasheet for TL316953**

### OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## **SNRNP27 Human shRNA Plasmid Kit (Locus ID 11017)**

#### **Product data:**

**Product Type:** shRNA Plasmids

**Product Name:** SNRNP27 Human shRNA Plasmid Kit (Locus ID 11017)

**Locus ID:** 11017 **Synonyms:** 27K; RY1

Vector:pGFP-C-shLenti (TR30023)E. coli Selection:Chloramphenicol (34 ug/ml)

Mammalian Cell Pur

Selection:

Puromycin

Format: Lentiviral plasmids

Components: SNRNP27 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID =

11017). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

RefSeq: <u>BC017890</u>, <u>NM 006857</u>, <u>NR 037862</u>, <u>NM 006857.1</u>, <u>NM 006857.2</u>, <u>BC017890.1</u>, <u>NM 006857.3</u>

UniProt ID: Q8WVK2

Summary: This gene encodes a serine/arginine-rich (SR) protein. SR proteins play important roles in pre-

mRNA splicing by facilitating the recognition and selection of splice sites. The encoded protein associates with the 25S U4/U6.U5 tri-snRNP, a major component of the U2-type spiceosome. The expression of this gene may be altered in cells infected with the human T-cell lymphotropic virus type 1 (HTLV-1) retrovirus. A pseudogene of this gene is located on the long arm of chromosome 5. Alternatively spliced transcript variants have been observed for

this gene. [provided by RefSeq, Feb 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 <u>techsupport@origene.com</u>. If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u>.





# 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).