

## Product datasheet for TL309425

#### OriGene Technologies, Inc.

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### SnoN (SKIL) Human shRNA Plasmid Kit (Locus ID 6498)

#### **Product data:**

**Product Type:** shRNA Plasmids

**Product Name:** SnoN (SKIL) Human shRNA Plasmid Kit (Locus ID 6498)

Locus ID:

Synonyms: SNO; SnoA; SnoI; SnoN

Vector: pGFP-C-shLenti (TR30023)

E. coli Selection: Chloramphenicol (34 ug/ml)

Mammalian Cell

Puromycin

Selection:

Format: Lentiviral plasmids

SKIL - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 6498). 5µg Components:

purified plasmid DNA per construct

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

NM 001145097, NM 001145098, NM 001248008, NM 005414, NM 005414.1, NM 005414.2, RefSeq:

> NM 005414.3, NM 005414.4, NM 001145097.1, NM 001145097.2, NM 001145098.1, NM 001145098.2, NM 001248008.1, BC059386, BC059386.1, BC041913, NM 001145098.3

**UniProt ID:** P12757

Summary: The protein encoded by this gene is a component of the SMAD pathway, which regulates cell

> growth and differentiation through transforming growth factor-beta (TGFB). In the absence of ligand, the encoded protein binds to the promoter region of TGFB-responsive genes and recruits a nuclear repressor complex. TGFB signaling causes SMAD3 to enter the nucleus and degrade this protein, allowing these genes to be activated. Four transcript variants encoding

three different isoforms have been found for this gene. [provided by RefSeq, Oct 2011]

These shRNA constructs were designed against multiple splice variants at this gene locus. To shRNA Design:

> be certain that your variant of interest is targeted, please contact <a href="techsupport@origene.com">techsupport@origene.com</a>. If you need a special design or shRNA sequence, please utilize our custom shRNA service.



# 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).