

## **Product datasheet for TR300472**

## OriGene Technologies, Inc.

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## WDR79 (WRAP53) Human shRNA Plasmid Kit (Locus ID 55135)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** WDR79 (WRAP53) Human shRNA Plasmid Kit (Locus ID 55135)

**Locus ID:** 55135

**Synonyms:** DKCB3; TCAB1; WDR79

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Fulbiliycili

Format: Retroviral plasmids

Components: WRAP53 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

55135). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

RefSeq: NM 001143990, NM 001143991, NM 001143992, NM 018081, NM 018081.1, NM 018081.2,

NM 001143990.1, NM 001143991.1, NM 001143992.1, BC002336, BC002336.2,

NM 001143992.2, NM 001143991.2

UniProt ID: Q9BUR4

**Summary:** This gene encodes an essential component of the telomerase holoenzyme complex, a

ribonucleoprotein complex required for telomere synthesis. This protein is enriched in Cajal bodies, nuclear sites of RNP processing that are important for telomerase function. It

interacts with dyskerin, TERT and TERC, other components of active telomerase, and with small Cajal body RNAs (scaRNAs), which are involved in modifying splicing RNAs. This mRNA also functions as a p53 antisense transcript, that regulates endogenous p53 mRNA levels and

further induction of p53 protein by targeting the 5' untranslated region of p53 mRNA. Alternatively spliced transcript variants which differ only in the 5' UTR have been found for

this gene. [provided by RefSeq, Mar 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).