

## **Product datasheet for TL305018**

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## **DGCR8 Human shRNA Plasmid Kit (Locus ID 54487)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** DGCR8 Human shRNA Plasmid Kit (Locus ID 54487)

**Locus ID:** 54487

Synonyms: C22orf12; DGCRK6; Gy1; pasha

**Vector:** pGFP-C-shLenti (TR30023)

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

**Mammalian Cell** 

Selection:

Puromycin

Format: Lentiviral plasmids

**Components:** DGCR8 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 54487).

5µg purified plasmid DNA per construct

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

RefSeq: NM 001190326, NM 022720, NM 022720.1, NM 022720.2, NM 022720.3, NM 022720.4,

NM 022720.5, NM 022720.6, NM 001190326.1, BC078147, BC009323, BC009984, BC037564,

BC071568, NM 022720.7

UniProt ID: Q8WYQ5

**Summary:** This gene encodes a subunit of the microprocessor complex which mediates the biogenesis

of microRNAs from the primary microRNA transcript. The encoded protein is a double-

stranded RNA binding protein that functions as the non-catalytic subunit of the

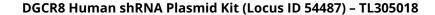
microprocessor complex. This protein is required for binding the double-stranded RNA substrate and facilitates cleavage of the RNA by the ribonuclease III protein, Drosha. Alternate

splicing results in multiple transcript variants. [provided by RefSeq, Jun 2010]

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).