

Product datasheet for TL310119

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PSG9 Human shRNA Plasmid Kit (Locus ID 5678)

Product data:

Product Type: shRNA Plasmids

Product Name: PSG9 Human shRNA Plasmid Kit (Locus ID 5678)

Locus ID: 5678

Synonyms: PS34; PSBG-9; PSBG-11; PSG11; PSGII

Vector: pGFP-C-shLenti (TR30023)

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

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

Components: PSG9 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 5678).

5µg purified plasmid DNA per construct

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

RefSeq: NM 001301707, NM 001301708, NM 001301709, NM 002784, NM 002784.1, NM 002784.2,

NM 002784.4, NM 001301709.1, NM 001301707.1, NM 001301708.1, BC020759, BC020759.1,

BC005925, NM 001301708.2, NM 001301707.2, NM 001301709.2

UniProt ID: Q00887

Summary: The protein encoded by this gene is a member of the pregnancy-specific glycoprotein (PSG)

family. This protein family and the closely related carcinoembryonic antigen cell adhesion molecule (CEACAM) gene family are both members of the immunoglobulin superfamily, and are organized as a large gene cluster. This protein is thought to inhibit platelet-fibrinogen interactions. Several studies suggest that reduced serum concentrations of PSGs are associated with fetal growth restrictions, while up-regulation of this gene has been observed

in colorectal cancers. Several pseudogenes of this gene are found on chromosome 19.

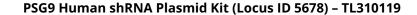
Alternative splicing results in multiple transcript variants that encode multiple protein

isoforms. [provided by RefSeq, Sep 2014]

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).