

Product datasheet for TR313593

OriGene Technologies, Inc.

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Cytochrome P450 2C9 (CYP2C9) Human shRNA Plasmid Kit (Locus ID 1559)

Product data:

Product Type: shRNA Plasmids

Product Name: Cytochrome P450 2C9 (CYP2C9) Human shRNA Plasmid Kit (Locus ID 1559)

Locus ID: 1559

Synonyms: CPC9; CYP2C; CYP2C10; CYPIIC9; P450IIC9

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection: Format:

Retroviral plasmids

Components: CYP2C9 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

1559). 5µg purified plasmid DNA per construct

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

RefSeq: <u>BC020754</u>, <u>NM 000771</u>, <u>NM 000771.1</u>, <u>NM 000771.2</u>, <u>NM 000771.3</u>, <u>BC020754.1</u>, <u>BC070317</u>,

BC125054, NM 000771.4

UniProt ID: P11712

Summary: This gene encodes a member of the cytochrome P450 superfamily of enzymes. The

cytochrome P450 proteins are monooxygenases which catalyze many reactions involved in drug metabolism and synthesis of cholesterol, steroids and other lipids. This protein localizes to the endoplasmic reticulum and its expression is induced by rifampin. The enzyme is known to metabolize many xenobiotics, including phenytoin, tolbutamide, ibuprofen and S-warfarin. Studies identifying individuals who are poor metabolizers of phenytoin and tolbutamide suggest that this gene is polymorphic. The gene is located within a cluster of cytochrome

P450 genes on chromosome 10q24. [provided by RefSeq, Jul 2008]

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>.



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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).