

## **Product datasheet for TL309182**

## OriGene Technologies, Inc.

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## Sorbitol Dehydrogenase (SORD) Human shRNA Plasmid Kit (Locus ID 6652)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Sorbitol Dehydrogenase (SORD) Human shRNA Plasmid Kit (Locus ID 6652)

**Locus ID:** 6652

**Synonyms:** HEL-S-95n; RDH; SORD1; SORDD; XDH

Vector:pGFP-C-shLenti (TR30023)E. coli Selection:Chloramphenicol (34 ug/ml)

**Mammalian Cell** 

**Cell** Puromycin

Selection:

Format: Lentiviral plasmids

**Components:** SORD - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 6652).

5µg purified plasmid DNA per construct

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

RefSeq: NM 003104, NR 034039, NM 003104.1, NM 003104.2, NM 003104.3, NM 003104.4,

NM 003104.5, BC021085, BC021085.2, BC025295

UniProt ID: 000796

Summary: Sorbitol dehydrogenase (SORD; EC 1.1.1.14) catalyzes the interconversion of polyols and their

corresponding ketoses, and together with aldose reductase (ALDR1; MIM 103880), makes up the sorbitol pathway that is believed to play an important role in the development of diabetic

complications (summarized by Carr and Markham, 1995 [PubMed 8535074]). The first

reaction of the pathway (also called the polyol pathway) is the reduction of glucose to sorbitol by ALDR1 with NADPH as the cofactor. SORD then oxidizes the sorbitol to fructose using

NAD(+) cofactor.[supplied by OMIM, Jul 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).