

## **Product datasheet for TL518894**

## **OriGene Technologies, Inc.** 9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US

Rockville, MD 20850, US
Phone: +1-888-267-4436
https://www.origene.com
techsupport@origene.com
EU: info-de@origene.com
CN: techsupport@origene.cn

## Spocd1 Mouse shRNA Plasmid (Locus ID 622480)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Spocd1 Mouse shRNA Plasmid (Locus ID 622480)

**Locus ID:** 622480

Synonyms: OTTMUSG00000009522

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

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

**Mammalian Cell** 

Selection:

Puromycin

Format: Lentiviral plasmids

**Components:** Spocd1 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 622480).

5µg purified plasmid DNA per construct

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

**RefSeq:** <u>NM 001144949</u>

**Summary:** Essential excecutor of PIWIL4-piRNA pathway directed transposon DNA methylation and

silencing in the male embryonic germ cells (PubMed:32674113). Associates with the de novo

DNA methylation machinery and repressive chromatin remodeling complexes

(PubMed:32674113). Tethering of PIWIL4 to a nascent transposable element transcript recruits repressive chromatin remodeling activities and the de novo methylation apparatus

through SPOCD1 (PubMed:32674113). Not required for piRNA biosynthesis

(PubMed:32674113).[UniProtKB/Swiss-Prot Function]

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).