

Product datasheet for TL300019V

OriGene Technologies, Inc.

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ZPBP2 Human shRNA Lentiviral Particle (Locus ID 124626)

Product data:

Product Type: shRNA Lentiviral Particles

Product Name: ZPBP2 Human shRNA Lentiviral Particle (Locus ID 124626)

Locus ID: 124626 Synonyms: ZPBPL

Vector: pGFP-C-shLenti (TR30023)

Format: Lentiviral particles

Components: ZPBP2 - Human shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble

control), 0.5 ml each, >10^7 TU/ml.

RefSeq: NM 198844, NM 199321, NM 198844.1, NM 198844.2, NM 199321.1, NM 199321.2,

BC043152, NM 198844.3, NM 199321.3

UniProt ID: Q6X784

Summary: Is implicated in sperm-oocyte interaction during fertilization.[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).

