

Product datasheet for TL301441

JLP (SPAG9) Human shRNA Plasmid Kit (Locus ID 9043)

Product data:

Product Type: shRNA Plasmids **Product Name:** JLP (SPAG9) Human shRNA Plasmid Kit (Locus ID 9043) Locus ID: 9043 CT89; HLC-6; HLC4; HLC6; JIP-4; JIP4; JLP; PHET; PIG6 Synonyms: Vector: pGFP-C-shLenti (TR30023) E. coli Selection: Chloramphenicol (34 ug/ml) Mammalian Cell Puromycin Selection: Format: Lentiviral plasmids **Components:** SPAG9 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 9043). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free. BC007524, NM 001130527, NM 001130528, NM 001251971, NM 003971, NM 172345, RefSeq: NM 003971.1, NM 003971.3, NM 003971.4, NM 003971.5, NM 001130528.1, <u>NM 001130528.2</u>, <u>NM 001251971.1</u>, <u>NM 001130527.1</u>, <u>NM 001130527.2</u>, <u>BC007524.1</u>, NM 172345.1, BC059946, BC146755, BC153878, NM 001130527.3, NM 003971.6, NM 001251971.2, NM 001130528.3 **UniProt ID:** 060271 Summary: This gene encodes a member of the cancer testis antigen gene family. The encoded protein functions as a scaffold protein that structurally organizes mitogen-activated protein kinases and mediates c-Jun-terminal kinase signaling. This protein also binds to kinesin-1 and may be involved in microtubule-based membrane transport. This protein may play a role in tumor growth and development. Alternate splicing results in multiple transcript variants. [provided by RefSeq, Oct 2011] 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 techsupport@origene.com. If you need a special design or shRNA sequence, please utilize our custom shRNA service.



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SPAG9) Human shRNA Plasmid Kit (Locus ID 9043) – TL301441 JLP (SPAG9) Human shRNA Plasmid Kit (Locus ID 9043) – TL301441

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).

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