

# Product datasheet for TR307647

# HOXA9 Human shRNA Plasmid Kit (Locus ID 3205)

## **Product data:**

#### **Product Type:** shRNA Plasmids **Product Name:** HOXA9 Human shRNA Plasmid Kit (Locus ID 3205) Locus ID: 3205 ABD-B; homeo box A9; homeobox A9; homeobox protein HOXA9; homeodomain protein Synonyms: HOXA9; HOX1; HOX1, ABD-B, HOX1G, HOX1.7, MGC1934; HOX1.7; HOX1G; MGC1934 Vector: pRS (TR20003) E. coli Selection: Ampicillin **Mammalian Cell** Puromycin Selection: Format: **Retroviral plasmids** HOXA9 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = **Components:** 3205). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free. RefSeq: NM 002142, NM 152739, NM 152739.1, NM 152739.2, NM 152739.3, BC006537, BC006537.2, BC010023, NM 152739.4 **UniProt ID:** P31269 Summary: In vertebrates, the genes encoding the class of transcription factors called homeobox genes are found in clusters named A, B, C, and D on four separate chromosomes. Expression of these proteins is spatially and temporally regulated during embryonic development. This gene is part of the A cluster on chromosome 7 and encodes a DNA-binding transcription factor which may regulate gene expression, morphogenesis, and differentiation. This gene is highly similar to the abdominal-B (Abd-B) gene of Drosophila. A specific translocation event which causes a fusion between this gene and the NUP98 gene has been associated with myeloid leukemogenesis. Read-through transcription exists between this gene and the upstream homeobox A10 (HOXA10) gene.[provided by RefSeq, Mar 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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### **GRIGENE** HOXA9 Human shRNA Plasmid Kit (Locus ID 3205) – TR307647

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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