

## Product datasheet for **TL505621V**

### Hsd3b7 Mouse shRNA Lentiviral Particle (Locus ID 101502)

#### Product data:

Product Type:	shRNA Lentiviral Particles
Product Name:	Hsd3b7 Mouse shRNA Lentiviral Particle (Locus ID 101502)
Locus ID:	101502
Synonyms:	A1195443; BB098564
Vector:	pGFP-C-shLenti (TR30023)
Format:	Lentiviral particles
Components:	Hsd3b7 - Mouse shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble control), 0.5 ml each, >10 <sup>7</sup> TU/ml.
RefSeq:	<a href="#">BC132605</a> , <a href="#">NM_001040684</a> , <a href="#">NM_133943</a> , <a href="#">NM_001040684.1</a> , <a href="#">NM_133943.1</a> , <a href="#">NM_133943.2</a> , <a href="#">BC006822</a> , <a href="#">BC012954</a> , <a href="#">BC018525</a> , <a href="#">BC138589</a>
UniProt ID:	<a href="#">Q9EQC1</a>
Summary:	The 3-beta-HSD enzymatic system plays a crucial role in the biosynthesis of all classes of hormonal steroids. HSD VII is active against four 7-alpha-hydroxylated sterols. Does not metabolize several different C(19/21) steroids as substrates. Involved in bile acid synthesis (PubMed:11067870). Plays a key role in cell positioning and movement in lymphoid tissues by mediating degradation of 7-alpha,25-dihydroxycholesterol (7-alpha,25-OHC): 7-alpha,25-OHC acts as a ligand for the G protein-coupled receptor GPR183/EBI2, a chemotactic receptor for a number of lymphoid cells (PubMed:22999953).[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 <a href="mailto:techsupport@origene.com">techsupport@origene.com</a> . If you need a special design or shRNA sequence, please utilize our <a href="#">custom shRNA service</a> .



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**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](mailto: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).