

# Product datasheet for TL309077

# ST3GAL4 Human shRNA Plasmid Kit (Locus ID 6484)

## **Product data:**

#### **Product Type:** shRNA Plasmids **Product Name:** ST3GAL4 Human shRNA Plasmid Kit (Locus ID 6484) Locus ID: 6484 CGS23; gal-NAc6S; NANTA3; SAT3; SIAT4; SIAT4C; ST-4; ST3GalA.2; ST3GalIV; STZ Synonyms: pGFP-C-shLenti (TR30023) Vector: E. coli Selection: Chloramphenicol (34 ug/ml) Mammalian Cell Puromycin Selection: Format: Lentiviral plasmids ST3GAL4 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 6484). **Components:** 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free. NM 001254757, NM 001254758, NM 001254759, NM 006278, NM 001348396, RefSeq: NM 001348397, NM 001348398, NM 001348399, NM 001348400, NR 145671, NM 006278.1, NM 006278.2, NM 001254759.1, NM 001254757.1, NM 001254758.1, BC010645, NM 001254759.2, NM 006278.3, NM 001254757.2 **UniProt ID:** Q11206 This gene encodes a member of the glycosyltransferase 29 family, a group of enzymes Summary: involved in protein glycosylation. The encoded protein is targeted to Golgi membranes but may be proteolytically processed and secreted. The gene product may also be involved in the increased expression of sialyl Lewis X antigen seen in inflammatory responses. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Dec 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** ST3GAL4 Human shRNA Plasmid Kit (Locus ID 6484) – TL309077

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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