

Product datasheet for TL309074

OriGene Technologies, Inc.

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CDw75 (ST6GAL1) Human shRNA Plasmid Kit (Locus ID 6480)

Product data:

Product Type: shRNA Plasmids

Product Name: CDw75 (ST6GAL1) Human shRNA Plasmid Kit (Locus ID 6480)

Locus ID: 6480

Synonyms: SIAT1; ST6Gall; ST6N

Vector: pGFP-C-shLenti (TR30023)

E. coli Selection: Chloramphenicol (34 ug/ml)

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

Components: ST6GAL1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 6480).

5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

RefSeq: NM 003032, NM 173216, NM 173217, NM 001353916, NM 173216.1, NM 173216.2,

NM 003032.1, NM 003032.2, NM 173217.1, NM 173217.2, BC031476, BC031476.1, BC040009,

BC040009.1, NM 003032.3

UniProt ID: P15907

Summary: This gene encodes a member of glycosyltransferase family 29. The encoded protein is a type

Il membrane protein that catalyzes the transfer of sialic acid from CMP-sialic acid to

galactose-containing substrates. The protein, which is normally found in the Golgi but can be proteolytically processed to a soluble form, is involved in the generation of the cell-surface carbohydrate determinants and differentiation antigens HB-6, CD75, and CD76. This gene has

been incorrectly referred to as CD75. Alternative splicing results in multiple transcript

variants. [provided by RefSeq, Jul 2017]

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).