

Product datasheet for TL308035V

OriGene Technologies, Inc.

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B4GALT4 Human shRNA Lentiviral Particle (Locus ID 8702)

Product data:

Product Type: shRNA Lentiviral Particles

Product Name: B4GALT4 Human shRNA Lentiviral Particle (Locus ID 8702)

Locus ID: 8702

Synonyms: B4Gal-T4; beta4Gal-T4

Vector: pGFP-C-shLenti (TR30023)

Format: Lentiviral particles

Components: B4GALT4 - Human shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1

scramble control), 0.5 ml each, >10^7 TU/ml.

RefSeg: NM 003778, NM 212543, NM 003778.1, NM 003778.2, NM 003778.3, NM 212543.1,

BC062618, BC062618.1, BC004523, NM 212543.2, NM 003778.4

UniProt ID: 060513

Summary: This gene is one of seven beta-1,4-galactosyltransferase (beta4GalT) genes. They encode type

Il membrane-bound glycoproteins that appear to have exclusive specificity for the donor substrate UDP-galactose; all transfer galactose in a beta1,4 linkage to similar acceptor sugars: GlcNAc, Glc, and Xyl. Each beta4GalT has a distinct function in the biosynthesis of different glycoconjugates and saccharide structures. As type Il membrane proteins, they have an N-terminal hydrophobic signal sequence that directs the protein to the Golgi apparatus and which then remains uncleaved to function as a transmembrane anchor. By sequence similarity, the beta4GalTs form four groups: beta4GalT1 and beta4GalT2, beta4GalT3 and beta4GalT4, beta4GalT5 and beta4GalT6, and beta4GalT7. The enzyme encoded by this gene appears to mainly play a role in glycolipid biosynthesis. Two alternatively spliced transcript

variants have been found for this gene. [provided by RefSeq, Jul 2008]

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).