

## **Product datasheet for TL302811**

## **OGT Human shRNA Plasmid Kit (Locus ID 8473)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** OGT Human shRNA Plasmid Kit (Locus ID 8473)

Locus ID: 8473

Synonyms: HRNT1, FLJ23071, MGC22921, O-GLCNAC

**Vector:** pGFP-C-shLenti (TR30023)

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

**Mammalian Cell** 

Selection:

Puromycin

Format: Lentiviral plasmids

Components: OGT - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 8473). 5µg

purified plasmid DNA per construct

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

RefSeq: NM 003605, NM 181672, NM 181673, NM 181672.1, NM 181672.2, NM 181673.1,

NM 181673.2, NM 003605.2, BC038180, BC038180.1, BC014434, BC015144, NM 181672.3,

NM 181673.3

UniProt ID: 015294

**Summary:** This gene encodes a glycosyltransferase that catalyzes the addition of a single N-

acetylglucosamine in O-glycosidic linkage to serine or threonine residues. Since both

phosphorylation and glycosylation compete for similar serine or threonine residues, the two processes may compete for sites, or they may alter the substrate specificity of nearby sites by steric or electrostatic effects. The protein contains multiple tetratricopeptide repeats that are required for optimal recognition of substrates. Alternatively spliced transcript variants

encoding distinct isoforms have been found for this gene. [provided by RefSeq, Oct 2009]

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>.



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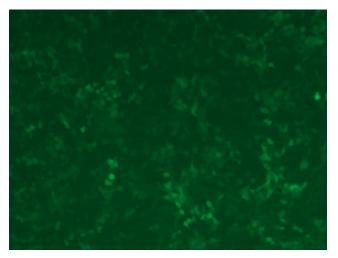


## Performance Guaranteed:

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.

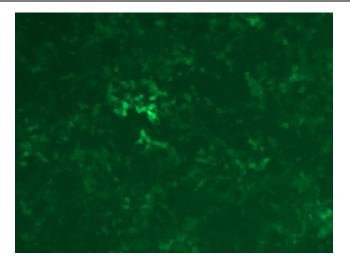
For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).

## **Product images:**

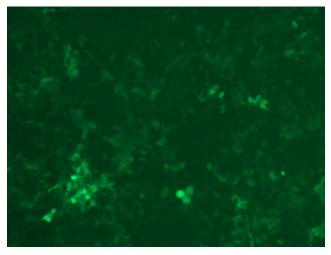


GFP signal was observed under microscope at 48 hours after transduction of TL302811A virus into HEK293 cells. TL302811A virus was prepared using lenti-shRNA TL302811A and [TR30037] packaging kit.

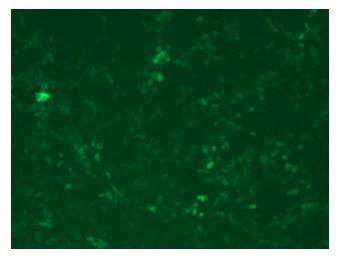




GFP signal was observed under microscope at 48 hours after transduction of TL302811B virus into HEK293 cells. TL302811B virus was prepared using lenti-shRNA TL302811B and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL302811C] virus into HEK293 cells. [TL302811C] virus was prepared using lenti-shRNA [TL302811C] and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL302811D] virus into HEK293 cells. [TL302811D] virus was prepared using lenti-shRNA [TL302811D] and [TR30037] packaging kit.