

## Product datasheet for RC212796L2V

## OriGene Technologies, Inc.

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## WBSCR17 (GALNT17) (NM\_022479) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** WBSCR17 (GALNT17) (NM\_022479) Human Tagged ORF Clone Lentiviral Particle

Symbol: GALNT17

Synonyms: GalNAc-T5L; GalNAc-T17; GalNAc-T19; GALNACT17; GALNT16; GALNT20; GALNTL3; WBSCR17

**Mammalian Cell** 

Selection:

None

**Vector:** pLenti-C-mGFP (PS100071)

Tag: mGFP

**ACCN:** NM\_022479 **ORF Size:** 1794 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC212796).

OTI Disclaimer:

Sequence:

The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

**RefSeg:** NM 022479.1

 RefSeq Size:
 3298 bp

 RefSeq ORF:
 1797 bp

 Locus ID:
 64409

 UniProt ID:
 Q6IS24

 Cytogenetics:
 7q11.22

**Domains:** RICIN, Glycos\_transf\_2

**Protein Families:** Transmembrane





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**Protein Pathways:** Metabolic pathways, O-Glycan biosynthesis

**MW:** 67.6 kDa

**Gene Summary:** This gene encodes an N-acetylgalactosaminyltransferase. This gene is located centromeric to

the common deleted region in Williams-Beuren syndrome (WBS), a multisystem

developmental disorder caused by the deletion of contiguous genes at 7q11.23. This protein

may play a role in membrane trafficking. [provided by RefSeq, Jan 2013]