

## Product datasheet for RC215501L4V

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## ALG12 (NM\_024105) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** ALG12 (NM\_024105) Human Tagged ORF Clone Lentiviral Particle

Symbol: ALG12

Synonyms: CDG1G; ECM39; hALG12; PP14673

**Mammalian Cell** 

Selection:

Puromycin

**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

**ACCN:** NM\_024105 **ORF Size:** 1464 bp

**ORF Nucleotide** 

•

Sequence:
OTI Disclaimer:

The ORF insert of this clone is exactly the same as(RC215501).

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 024105.3

 RefSeq Size:
 2387 bp

 RefSeq ORF:
 1467 bp

 Locus ID:
 79087

 UniProt ID:
 Q9BV10

 Cytogenetics:
 22q13.33

**Protein Families:** Transmembrane

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





## ALG12 (NM\_024105) Human Tagged ORF Clone Lentiviral Particle - RC215501L4V

**MW:** 54.6 kDa

**Gene Summary:** This gene encodes a member of the glycosyltransferase 22 family. The encoded protein

catalyzes the addition of the eighth mannose residue in an alpha-1,6 linkage onto the dolichol-PP-oligosaccharide precursor (dolichol-PP-Man(7)GlcNAc(2)) required for protein glycosylation. Mutations in this gene have been associated with congenital disorder of glycosylation type Ig (CDG-Ig)characterized by abnormal N-glycosylation. [provided by RefSeq,

Jul 2008]