

## Product datasheet for RC207088L2V

## OriGene Technologies, Inc.

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## LMAN1 (NM\_005570) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: LMAN1 (NM\_005570) Human Tagged ORF Clone Lentiviral Particle

Symbol: LMAN1

Synonyms: ERGIC-53; ERGIC53; F5F8D; FMFD1; gp58; MCFD1; MR60

Mammalian Cell

Selection:

None

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

Tag: mGFP

**ACCN:** NM\_005570 **ORF Size:** 1530 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC207088).

Sequence:
OTI Disclaimer:

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 005570.2

 RefSeq Size:
 4848 bp

 RefSeq ORF:
 1533 bp

 Locus ID:
 3998

 UniProt ID:
 P49257

 Cytogenetics:
 18q21.32

Domains: Lectin\_leg-like

**Protein Families:** Druggable Genome, Transmembrane





ORIGENE

**MW:** 57.5 kDa

**Gene Summary:** 

The protein encoded by this gene is a membrane mannose-specific lectin that cycles between the endoplasmic reticulum, endoplasmic reticulum-Golgi intermediate compartment, and cis-Golgi, functioning as a cargo receptor for glycoprotein transport. The protein has an N-terminal signal sequence, a calcium-dependent and pH-sensitive carbohydrate recognition domain, a stalk region that functions in oligomerization, a transmembrane domain, and a short cytoplasmic domain required for organelle targeting. Allelic variants of this gene are associated with the autosomal recessive disorder combined factor V-factor VIII deficiency. [provided by RefSeq, Jul 2015]