

## Product datasheet for MR211082L4V

## OriGene Technologies, Inc.

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## Dag1 (NM\_010017) Mouse Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** Dag1 (NM\_010017) Mouse Tagged ORF Clone Lentiviral Particle

Symbol: Dag1

Synonyms: D9Wsu13; D9Wsu13e; DG; Dp71; Dp427

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_010017 **ORF Size:** 2679 bp

**ORF Nucleotide** 

OTI Disclaimer:

Sequence:

The ORF insert of this clone is exactly the same as(MR211082).

Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements.

Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA.

Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at

custsupport@origene.com or by calling 301.340.3188 option 3 for pricing and delivery.

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.

RefSeq: <u>NM 010017.3, NP 034147.1</u>

**RefSeq Size:** 5591 bp **RefSeq ORF:** 2682 bp





## Dag1 (NM\_010017) Mouse Tagged ORF Clone Lentiviral Particle - MR211082L4V

**Locus ID:** 13138

 UniProt ID:
 Q62165

 Cytogenetics:
 9 59.08 cM

**Gene Summary:** This gene encodes dystroglycan, a central component of dystrophin-glycoprotein complex

that links the extracellular matrix and the cytoskeleton in the skeletal muscle. The encoded preproprotein undergoes O- and N-glycosylation, and proteolytic processing to generate alpha and beta subunits. A complete lack of the encoded protein in mice results in embryonic lethality due to the disorganization of Reichert's membrane. Chimeric mice deficient in the encoded protein overcome embryonic lethality but develop a progressive muscular dystrophy. Alternative splicing results in multiple transcript variants, all encoding the same

protein. [provided by RefSeq, Nov 2015]