

## Product datasheet for RC217883L3V

## OriGene Technologies, Inc.

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## MEGF10 (NM\_032446) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** MEGF10 (NM\_032446) Human Tagged ORF Clone Lentiviral Particle

Symbol: MEGF10

**Synonyms:** EMARDD; SR-F3

Mammalian Cell

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Puromycin

Selection: Vector:

pLenti-C-Myc-DDK-P2A-Puro (PS100092)

 Tag:
 Myc-DDK

 ACCN:
 NM\_032446

ORF Size: 3420 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC217883).

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 032446.1

 RefSeq Size:
 7522 bp

 RefSeq ORF:
 3423 bp

 Locus ID:
 84466

 UniProt ID:
 Q96KG7

 Cytogenetics:
 5q23.2

**Domains:** EGF\_Lam, EGF, EGF

**Protein Families:** Transmembrane





ORIGENE

**MW:** 122 kDa

**Gene Summary:** This gene encodes a member of the multiple epidermal growth factor-like domains protein

family. The encoded protein plays a role in cell adhesion, motility and proliferation, and is a critical mediator of apoptotic cell phagocytosis as well as amyloid-beta peptide uptake in the brain. Expression of this gene may be associated with schizophrenia, and mutations in this gene are a cause of early-onset myopathy, areflexia, respiratory distress, and dysphagia (EMARDD) as well as congenital myopathy with minicores. Alternatively spliced transcript

variants have been observed for this gene. [provided by RefSeq, Apr 2012]