

## Product datasheet for RC205269L3V

## OriGene Technologies, Inc.

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## Semaphorin 3c (SEMA3C) (NM\_006379) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** Semaphorin 3c (SEMA3C) (NM\_006379) Human Tagged ORF Clone Lentiviral Particle

Symbol: Semaphorin 3c Synonyms: SEMAE; SemE

**Mammalian Cell** 

Selection:

Puromycin

**Vector:** pLenti-C-Myc-DDK-P2A-Puro (PS100092)

 Tag:
 Myc-DDK

 ACCN:
 NM\_006379

 ORF Size:
 2253 bp

**ORF Nucleotide** 

OTI Disclaimer:

The ORF insert of this clone is exactly the same as(RC205269).

Sequence:

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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

The molecular sequence of this clone aligns with the gene accession number as a point of

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 006379.2

 RefSeq Size:
 5189 bp

 RefSeq ORF:
 2256 bp

 Locus ID:
 10512

 UniProt ID:
 Q99985

 Cytogenetics:
 7q21.11

**Domains:** Sema, IG, PSI

**Protein Families:** Secreted Protein





## Semaphorin 3c (SEMA3C) (NM\_006379) Human Tagged ORF Clone Lentiviral Particle – RC205269L3V

**Protein Pathways:** Axon guidance

**MW:** 85 kDa

**Gene Summary:** This gene encodes a secreted glycoprotein that belongs to the semaphorin class 3 family of

neuronal guidance cues. The encoded protein contains an N-terminal sema domain, integrin and immunoglobulin-like domains, and a C-terminal basic domain. Homodimerization and proteolytic cleavage of the C-terminal propeptide are necessary for the function of the encoded protein. It binds a neuropilin co-receptor before forming a heterotrimeric complex with an associated plexin. An increase in the expression of this gene correlates with an increase in cancer cell invasion and adhesion. Naturally occurring mutations in this gene are

associated with Hirschsprung disease. [provided by RefSeq, May 2017]