

## Product datasheet for RC203767L4V

## 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

## SERPING1 (NM\_000062) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** SERPING1 (NM\_000062) Human Tagged ORF Clone Lentiviral Particle

Symbol: SERPING1

Synonyms: C1IN; C1INH; C1NH; HAE1; HAE2

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_000062 **ORF Size:** 1500 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC203767).

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 000062.2

 RefSeq Size:
 1984 bp

 RefSeq ORF:
 1503 bp

 Locus ID:
 710

 UniProt ID:
 P05155

Cytogenetics: 11q12.1

Domains: SERPIN

**Protein Families:** Druggable Genome, Secreted Protein





## SERPING1 (NM\_000062) Human Tagged ORF Clone Lentiviral Particle - RC203767L4V

**Protein Pathways:** Complement and coagulation cascades

**MW:** 55.2 kDa

**Gene Summary:** This gene encodes a highly glycosylated plasma protein involved in the regulation of the

complement cascade. Its encoded protein, C1 inhibitor, inhibits activated C1r and C1s of the first complement component and thus regulates complement activation. It is synthesized in the liver, and its deficiency is associated with hereditary angioneurotic oedema (HANE). Alternative splicing results in multiple transcript variants encoding the same isoform.

[provided by RefSeq, May 2020]