

Product datasheet for RC206340L3V

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

SPG7 (NM_003119) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: SPG7 (NM 003119) Human Tagged ORF Clone Lentiviral Particle

Symbol: SPG7

Synonyms: CAR; CMAR; PGN; SPG5C

Mammalian Cell

Selection:

Puromycin

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

 Tag:
 Myc-DDK

 ACCN:
 NM_003119

ORF Size: 2385 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC206340).

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 003119.2

 RefSeq Size:
 3102 bp

 RefSeq ORF:
 2388 bp

 Locus ID:
 6687

 UniProt ID:
 Q9UQ90

 Cytogenetics:
 16q24.3

Domains: Peptidase_M41, AAA, AAA

Protein Families: Protease, Transmembrane





ORIGENE

MW: 88.2 kDa

Gene Summary: This gene encodes a mitochondrial metalloprotease protein that is a member of the AAA

family. Members of this protein family share an ATPase domain and have roles in diverse cellular processes including membrane trafficking, intracellular motility, organelle biogenesis, protein folding, and proteolysis. Mutations in this gene cause autosomal recessive spastic paraplegia 7. Two transcript variants encoding distinct isoforms have been identified.

[provided by RefSeq, Mar 2014]