

Product datasheet for RC200359L4V

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

LONP1 (NM_004793) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: LONP1 (NM_004793) Human Tagged ORF Clone Lentiviral Particle

Symbol: LONP1

Synonyms: CODASS; hLON; LON; LONP; PIM1; PRSS15

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

ACCN: NM_004793 **ORF Size:** 2877 bp

ORF Nucleotide

2077 59

Sequence:

The ORF insert of this clone is exactly the same as(RC200359).

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 004793.2

 RefSeq Size:
 3221 bp

 RefSeq ORF:
 2880 bp

 Locus ID:
 9361

 UniProt ID:
 P36776

 Cytogenetics:
 19p13.3

Domains: LON, AAA, AAA

Protein Families: Druggable Genome, Protease



ORIGENE

MW: 106.5 kDa

Gene Summary:

This gene encodes a mitochondrial matrix protein that belongs to the Lon family of ATP-dependent proteases. This protein mediates the selective degradation of misfolded, unassembled or oxidatively damaged polypeptides in the mitochondrial matrix. It may also have a chaperone function in the assembly of inner membrane protein complexes, and participate in the regulation of mitochondrial gene expression and maintenance of the integrity of the mitochondrial genome. Decreased expression of this gene has been noted in a patient with hereditary spastic paraplegia (PMID:18378094). Alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Feb 2013]