

## Product datasheet for RC226309L3V

## OriGene Technologies, Inc.

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## CPS1 (NM\_001122634) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** CPS1 (NM\_001122634) Human Tagged ORF Clone Lentiviral Particle

Symbol: CPS1

Synonyms: CPSASE1; PHN

Mammalian Cell Puromycin

Selection:

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

Tag: Myc-DDK

**ACCN:** NM\_001122634

ORF Size: 3147 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC226309).

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.

**RefSeq:** <u>NM 001122634.2</u>

 RefSeq Size:
 4794 bp

 RefSeq ORF:
 3150 bp

 Locus ID:
 1373

 UniProt ID:
 P31327

 Cytogenetics:
 2q34

**Protein Families:** Druggable Genome





## CPS1 (NM\_001122634) Human Tagged ORF Clone Lentiviral Particle - RC226309L3V

Protein Pathways: Alanine, aspartate and glutamate metabolism, Arginine and proline metabolism, Metabolic

pathways, Nitrogen metabolism

**MW**: 116 kDa

**Gene Summary:** The mitochondrial enzyme encoded by this gene catalyzes synthesis of carbamoyl phosphate

from ammonia and bicarbonate. This reaction is the first committed step of the urea cycle, which is important in the removal of excess urea from cells. The encoded protein may also represent a core mitochondrial nucleoid protein. Three transcript variants encoding different isoforms have been found for this gene. The shortest isoform may not be localized to the mitochondrion. Mutations in this gene have been associated with carbamoyl phosphate synthetase deficiency, susceptibility to persistent pulmonary hypertension, and susceptibility to venoocclusive disease after bone marrow transplantation.[provided by RefSeq, May 2010]