

Product datasheet for RC200698L3V

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

PRPS1 (NM_002764) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: PRPS1 (NM_002764) Human Tagged ORF Clone Lentiviral Particle

Symbol: PRPS1

Synonyms: ARTS; CMTX5; DFN2; DFNX1; PPRibP; PRS-I; PRSI

Mammalian Cell

Selection:

Puromycin

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

Tag: Myc-DDK
ACCN: NM 002764

ORF Size: 954 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC200698).

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 002764.2

 RefSeq Size:
 2156 bp

 RefSeq ORF:
 957 bp

 Locus ID:
 5631

 UniProt ID:
 P60891

 Cytogenetics:
 Xq22.3

Domains: Pribosyltran

Protein Families: Druggable Genome





PRPS1 (NM_002764) Human Tagged ORF Clone Lentiviral Particle - RC200698L3V

Protein Pathways: Metabolic pathways, Pentose phosphate pathway, Purine metabolism

MW: 34.8 kDa

Gene Summary: This gene encodes an enzyme that catalyzes the phosphoribosylation of ribose 5-phosphate

to 5-phosphoribosyl-1-pyrophosphate, which is necessary for purine metabolism and nucleotide biosynthesis. Defects in this gene are a cause of phosphoribosylpyrophosphate synthetase superactivity, Charcot-Marie-Tooth disease X-linked recessive type 5 and Arts Syndrome. Two transcript variants encoding different isoforms have been found for this

gene. [provided by RefSeq, Feb 2011]