

Product datasheet for RC223312L1V

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

Thymidine Kinase 2 (TK2) (NM 004614) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Thymidine Kinase 2 (TK2) (NM 004614) Human Tagged ORF Clone Lentiviral Particle

Symbol: Thymidine Kinase 2

Synonyms: MTDPS2; MTTK; PEOB3; SCA31

Mammalian Cell

Selection:

None

Vector: pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK
ACCN: NM 004614

ORF Size: 795 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC223312).

Sequence:

Domains:

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 004614.3

 RefSeq Size:
 3675 bp

 RefSeq ORF:
 798 bp

 Locus ID:
 7084

 UniProt ID:
 000142

 Cytogenetics:
 16q21

Protein Families: Druggable Genome

dNK





Thymidine Kinase 2 (TK2) (NM_004614) Human Tagged ORF Clone Lentiviral Particle – RC223312L1V

Protein Pathways: Drug metabolism - other enzymes, Metabolic pathways, Pyrimidine metabolism

MW: 31 kDa

Gene Summary: This gene encodes a deoxyribonucleoside kinase that specifically phosphorylates thymidine,

deoxycytidine, and deoxyuridine. The encoded enzyme localizes to the mitochondria and is required for mitochondrial DNA synthesis. Mutations in this gene are associated with a myopathic form of mitochondrial DNA depletion syndrome. Alternate splicing results in multiple transcript variants encoding distinct isoforms, some of which lack transit peptide, so

are not localized to mitochondria. [provided by RefSeq, Dec 2012]