

Product datasheet for SC331029

OriGene Technologies, Inc.

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Thymidine Kinase 2 (TK2) (NM_001272050) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: Thymidine Kinase 2 (TK2) (NM 001272050) Human Untagged Clone

Tag: Tag Free

Symbol: TK2

Synonyms: MTDPS2; MTTK; PEOB3; SCA31

Vector: pCMV6-Entry (PS100001)

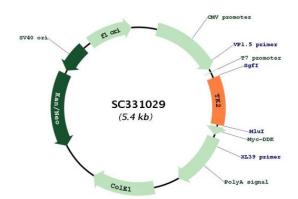
Fully Sequenced ORF: >SC331029 representing NM_001272050.

Blue=Insert sequence Red=Cloning site Green=Tag(s)

GAGAATCGGAAGCATTGCCCATAG

Restriction Sites: Sgfl-Mlul

Plasmid Map:





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ACCN: NM_001272050

Insert Size: 507 bp

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of

shipping when stored at -20°C.

RefSeq: <u>NM 001272050.1</u>

 RefSeq Size:
 4907 bp

 RefSeq ORF:
 507 bp

 Locus ID:
 7084

 UniProt ID:
 000142

 Cytogenetics:
 16q21

Protein Families: Druggable Genome

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

MW: 20.3 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]

Transcript Variant: This variant (9) contains an alternate 5' exon and initiates translation at a downstream start codon, compared to variant 1. The encoded isoform (7) has a shorter N-terminus, compared to isoform 1, but is predicted to be a mitochondrial protein. Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used

for the transcript record were based on transcript alignments.