

Product datasheet for RC203764L3V

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

OXCT1 (NM_000436) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: OXCT1 (NM 000436) Human Tagged ORF Clone Lentiviral Particle

Symbol: OXCT1

Synonyms: OXCT; SCOT

Mammalian Cell Pu

Selection:

ACCN:

Puromycin

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

NM 000436

Tag: Myc-DDK

ORF Size: 1560 bp

ORF Nucleotide

5p13.1

Sequence:

Cytogenetics:

The ORF insert of this clone is exactly the same as(RC203764).

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 000436.3</u>

RefSeq Size: 3572 bp
RefSeq ORF: 1563 bp
Locus ID: 5019

UniProt ID: P55809

Domains: CoA_trans



OXCT1 (NM_000436) Human Tagged ORF Clone Lentiviral Particle - RC203764L3V

Protein Pathways: Butanoate metabolism, Synthesis and degradation of ketone bodies, Valine, leucine and

isoleucine degradation

MW: 56.2 kDa

Gene Summary: This gene encodes a member of the 3-oxoacid CoA-transferase gene family. The encoded

protein is a homodimeric mitochondrial matrix enzyme that plays a central role in extrahepatic ketone body catabolism by catalyzing the reversible transfer of coenzyme A

from succinyl-CoA to acetoacetate. Mutations in this gene are associated with succinyl CoA:3-

oxoacid CoA transferase deficiency. [provided by RefSeq, Jul 2008]