

Product datasheet for RC201972L1V

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

Methylmalonyl Coenzyme A mutase (MUT) (NM_000255) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Methylmalonyl Coenzyme A mutase (MUT) (NM_000255) Human Tagged ORF Clone Lentiviral

Particle

Symbol: Methylmalonyl Coenzyme A mutase

Synonyms: MCM; MUT

Mammalian Cell

Selection:

None

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

 Tag:
 Myc-DDK

 ACCN:
 NM_000255

ORF Size: 2250 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC201972).

OTI Disclaimer:

Sequence:

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 000255.1</u>

 RefSeq Size:
 3886 bp

 RefSeq ORF:
 2253 bp

 Locus ID:
 4594

 UniProt ID:
 P22033

Cytogenetics: 6p12.3

Domains: MM_CoA_mutase, B12-binding





Methylmalonyl Coenzyme A mutase (MUT) (NM_000255) Human Tagged ORF Clone Lentiviral Particle – RC201972L1V

Protein Families: Druggable Genome

Protein Pathways: Metabolic pathways, Propanoate metabolism, Valine, leucine and isoleucine degradation

MW: 83.2 kDa

Gene Summary: This gene encodes the mitochondrial enzyme methylmalonyl Coenzyme A mutase. In

humans, the product of this gene is a vitamin B12-dependent enzyme which catalyzes the isomerization of methylmalonyl-CoA to succinyl-CoA, while in other species this enzyme may have different functions. Mutations in this gene may lead to various types of methylmalonic

aciduria. [provided by RefSeq, Jul 2008]