

Product datasheet for RC217919L4V

OriGene Technologies, Inc.

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AMPD1 (NM_000036) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: AMPD1 (NM_000036) Human Tagged ORF Clone Lentiviral Particle

Symbol: AMPD1

Synonyms: MAD; MADA; MMDD

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

ACCN: NM_000036 **ORF Size:** 2241 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC217919).

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 000036.1</u>, <u>NP 000027.1</u>

 RefSeq Size:
 2426 bp

 RefSeq ORF:
 2244 bp

 Locus ID:
 270

 UniProt ID:
 P23109

 Cytogenetics:
 1p13.2

Domains: A deaminase

Protein Families: Druggable Genome





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Protein Pathways: Metabolic pathways, Purine metabolism

MW: 86.5 kDa

Gene Summary: Adenosine monophosphate deaminase 1 catalyzes the deamination of AMP to IMP in skeletal

muscle and plays an important role in the purine nucleotide cycle. Two other genes have been identified, AMPD2 and AMPD3, for the liver- and erythocyte-specific isoforms, respectively. Deficiency of the muscle-specific enzyme is apparently a common cause of exercise-induced myopathy and probably the most common cause of metabolic myopathy in the human. Alternatively spliced transcript variants encoding different isoforms have been

identified in this gene.[provided by RefSeq, Feb 2010]