

Product datasheet for RC201507L3V

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

ACAD8 (NM 014384) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: ACAD8 (NM_014384) Human Tagged ORF Clone Lentiviral Particle

Symbol:

ACAD-8; ARC42; IBDH Synonyms:

Mammalian Cell

Selection:

Puromycin

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

Tag: Myc-DDK NM 014384 ACCN:

ORF Size: 1245 bp

ORF Nucleotide

OTI Disclaimer:

Sequence:

The ORF insert of this clone is exactly the same as(RC201507).

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: NM 014384.2

RefSeq Size: 2216 bp RefSeq ORF: 1248 bp Locus ID: 27034 **UniProt ID:** Q9UKU7 Cytogenetics: 11q25

Domains: Acyl-CoA_dh, Acyl-CoA_dh_M, Acyl-CoA_dh_N

Protein Families: Transcription Factors





ACAD8 (NM_014384) Human Tagged ORF Clone Lentiviral Particle - RC201507L3V

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

MW: 45.1 kDa

Gene Summary: This gene encodes a member of the acyl-CoA dehydrogenase family of enzymes that catalyze

the dehydrogenation of acyl-CoA derivatives in the metabolism of fatty acids or branch chained amino acids. The encoded protein is a mitochondrial enzyme that functions in catabolism of the branched-chain amino acid valine. Defects in this gene are the cause of

isobutyryl-CoA dehydrogenase deficiency.[provided by RefSeq, Nov 2009]