

Product datasheet for RC209853L3V

OriGene Technologies, Inc.

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HADHB (NM_000183) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: HADHB (NM_000183) Human Tagged ORF Clone Lentiviral Particle

Symbol: HADHE

Synonyms: ECHB; MSTP029; MTPB; TP-BETA

Mammalian Cell

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Selection:

Puromycin

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

Tag: Myc-DDK
ACCN: NM_000183

ORF Size: 1428 bp

ORF Nucleotide

F Nucleatide The OR

Sequence:
OTI Disclaimer:

The ORF insert of this clone is exactly the same as(RC209853).

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.

RefSeg: NM 000183.2

 RefSeq Size:
 2196 bp

 RefSeq ORF:
 1425 bp

 Locus ID:
 3032

 UniProt ID:
 P55084

 Cytogenetics:
 2p23.3

Domains: thiolase





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Protein Pathways: Fatty acid elongation in mitochondria, Fatty acid metabolism, Metabolic pathways, Valine,

leucine and isoleucine degradation

MW: 51.4 kDa

Gene Summary: This gene encodes the beta subunit of the mitochondrial trifunctional protein, which catalyzes

the last three steps of mitochondrial beta-oxidation of long chain fatty acids. The

mitochondrial membrane-bound heterocomplex is composed of four alpha and four beta subunits, with the beta subunit catalyzing the 3-ketoacyl-CoA thiolase activity. The encoded protein can also bind RNA and decreases the stability of some mRNAs. The genes of the alpha and beta subunits of the mitochondrial trifunctional protein are located adjacent to each other in the human genome in a head-to-head orientation. Mutations in this gene result in trifunctional protein deficiency. Alternatively spliced transcript variants encoding different

isoforms have been described. [provided by RefSeq, Jul 2013]