

Product datasheet for RC203539L3V

OriGene Technologies, Inc.

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ERAB (HSD17B10) (NM 001037811) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: ERAB (HSD17B10) (NM_001037811) Human Tagged ORF Clone Lentiviral Particle

Symbol: ERAE

Synonyms: 17b-HSD10; ABAD; CAMR; DUPXp11.22; ERAB; HADH2; HCD2; HSD10MD; MHBD; MRPP2;

MRX17; MRX31; MRXS10; SCHAD; SDR5C1

Mammalian Cell

Selection:

Puromycin

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

Tag: Myc-DDK

ACCN: NM_001037811

ORF Size: 756 bp

ORF Nucleotide

The

Sequence:

Cytogenetics:

The ORF insert of this clone is exactly the same as(RC203539).

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 001037811.2</u>, <u>NP 001032900.1</u>

Xp11.22

 RefSeq Size:
 936 bp

 RefSeq ORF:
 759 bp

 Locus ID:
 3028

 UniProt ID:
 Q99714

Protein Families: Druggable Genome





ERAB (HSD17B10) (NM_001037811) Human Tagged ORF Clone Lentiviral Particle - RC203539L3V

Protein Pathways: Alzheimer's disease, Metabolic pathways, Valine, leucine and isoleucine degradation

MW: 26 kDa

Gene Summary: This gene encodes 3-hydroxyacyl-CoA dehydrogenase type II, a member of the short-chain

dehydrogenase/reductase superfamily. The gene product is a mitochondrial protein that catalyzes the oxidation of a wide variety of fatty acids and steroids, and is a subunit of mitochondrial ribonuclease P, which is involved in tRNA maturation. The protein has been implicated in the development of Alzheimer disease, and mutations in the gene are the cause of 17beta-hydroxysteroid dehydrogenase type 10 (HSD10) deficiency. Several alternatively spliced transcript variants have been identified, but the full-length nature of only two

transcript variants has been determined. [provided by RefSeq, Aug 2014]