

## Product datasheet for **SC201339**

### ERAB (HSD17B10) (NM\_004493) Human 3' UTR Clone

#### Product data:

Product Type:	3' UTR Clones
Product Name:	ERAB (HSD17B10) (NM_004493) Human 3' UTR Clone
Vector:	pMirTarget (PS100062)
Symbol:	HSD17B10
Synonyms:	17b-HSD10; ABAD; CAMR; DUPXp11.22; ERAB; HADH2; HCD2; HSD10MD; MHBD; MRPP2; MRX17; MRX31; MRXS10; SCHAD; SDR5C1
ACCN:	NM_004493
Insert Size:	173 bp
Insert Sequence:	>SC201339 3'UTR clone of NM_004493 The sequence shown below is from the reference sequence of NM_004493. The complete sequence of this clone may contain minor differences, such as SNPs. <b>Blue</b> =Stop Codon <b>Red</b> =Cloning site  GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAAGCCAAGAAGGGCGGAAAGATCGCCGTG TAACAATTGGCAGAGCTCAGAATTCAA <b>GCGATCGCC</b> CTGGATGGGGCCATTTCGTATGCAGCCT <b>TGA</b> AGGGAGAAGGCAGAGAAAACACACGCTCCTCTGCCCTTC CTTCCCTGGGGTACTACTCTCCAGCTTGGGAGGAAGCCAGTAGCCATTTGTAACTGCCTACCAAGTC GCCCTCTGTGCCTAATAAAGTCTCTTTTCTCACA <b>ACGCGT</b> AAGCGGCCGCGCATCTAGATTGGAAGAAAATGACCGACCAAGCGACGCCAACCTGCCATCA CGAGATTTTCGATTCCACCGCCGCTTCTATGAAAGG
Restriction Sites:	Sgfl-Mlul
OTI Disclaimer:	Our molecular clone sequence data has been matched to the sequence identifier above as a point of reference. Note that the complete sequence of this clone is largely the same as the reference sequence but may contain minor differences, e.g., single nucleotide polymorphisms (SNPs).
Components:	The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 µg dried plasmid DNA. The package also includes 100 pmols of both the corresponding 5' and 3' vector primers in separate vials.
RefSeq:	<u><a href="#">NM_004493.3</a></u>



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**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]

**Locus ID:**

3028

**MW:**

6.4