

Product datasheet for SC204821

OriGene Technologies, Inc.

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Hydroxysteroid (17 beta) Dehydrogenase 4 (HSD17B4) (NM 000414) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: Hydroxysteroid (17 beta) Dehydrogenase 4 (HSD17B4) (NM_000414) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

Symbol: HSD17B4

Synonyms: DBP; MFE-2; MFP-2; MPF-2; PRLTS1; SDR8C1

ACCN: NM_000414

Insert Size: 368 bp

Insert Sequence: >SC204821 3'UTR clone of NM_000414

The sequence shown below is from the reference sequence of NM_000414. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

ATGATTCTTAAAGACTACGCCAAGCTCTGAAGGGCACACTACACTATTAATAAAAATGGAATCATTAAA TACTCTCTTCACCCAAATATGCTTGATTATTCTGCAAAAGTGATTAGAACTAAGATGCAGGGGAAATTG CTTAACATTTTCAGATATCAGATAACTGCAGATTTTCATTTTCTACTAATTTTTCATGTATCATTATTATTATAAGCTAGCACATAATTATCCTTCTGTTCTTAGATCTGTATCTTCATAATAA AAAATTTTGCCCAAGTCCTGTTTCCTTAGAATTTAAATAA

CAATAAAGGCCTTTGATACCTTT

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

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 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

RefSeg: NM 000414.4





Hydroxysteroid (17 beta) Dehydrogenase 4 (HSD17B4) (NM_000414) Human 3' UTR Clone – SC204821

Summary: The protein encoded by this gene is a bifunctional enzyme that is involved in the peroxisomal

beta-oxidation pathway for fatty acids. It also acts as a catalyst for the formation of 3-ketoacyl-CoA intermediates from both straight-chain and 2-methyl-branched-chain fatty acids. Defects in this gene that affect the peroxisomal fatty acid beta-oxidation activity are a cause of D-bifunctional protein deficiency (DBPD). An apparent pseudogene of this gene is present on chromosome 8. Multiple alternatively spliced transcript variants encoding distinct

isoforms have been found for this gene. [provided by RefSeq, May 2014]

Locus ID: 3295 MW: 14.2