

Product datasheet for **RC200460L4V**

Hydroxysteroid (17 beta) Dehydrogenase 4 (HSD17B4) (NM_000414) Human Tagged ORF Clone Lentiviral Particle

Product data:

| | |
|---------------------------|--|
| Product Type: | Lentiviral Particles |
| Product Name: | Hydroxysteroid (17 beta) Dehydrogenase 4 (HSD17B4) (NM_000414) Human Tagged ORF Clone Lentiviral Particle |
| Symbol: | HSD17B4 |
| Synonyms: | DBP; MFE-2; MFP-2; MPF-2; PRLTS1; SDR8C1 |
| Mammalian Cell Selection: | Puromycin |
| Vector: | pLenti-C-mGFP-P2A-Puro (PS100093) |
| Tag: | mGFP |
| ACCN: | NM_000414 |
| ORF Size: | 2208 bp |
| ORF Nucleotide Sequence: | The ORF insert of this clone is exactly the same as(RC200460). |
| 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: | NM_000414.1 |
| RefSeq Size: | 2710 bp |
| RefSeq ORF: | 2211 bp |
| Locus ID: | 3295 |
| UniProt ID: | P51659 |
| Cytogenetics: | 5q23.1 |
| Domains: | adh_short, MaoC_dehydratas, SCP2 |



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|--------------------------|--|
| Protein Families: | Druggable Genome |
| Protein Pathways: | Metabolic pathways, Primary bile acid biosynthesis |
| MW: | 79.7 kDa |
| Gene Summary: | <p>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]</p> |