

Product datasheet for SC204121

FATP2 (SLC27A2) (NM_001159629) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones **Product Name:** FATP2 (SLC27A2) (NM_001159629) Human 3' UTR Clone Symbol: FATP2 ACSVL1; FACVL1; FATP2; hFACVL1; HsT17226; VLACS; VLCS Synonyms: **Mammalian Cell** Neomycin Selection: pMirTarget (PS100062) Vector: ACCN: NM 001159629 Insert Size: 319 bp **Insert Sequence:** >SC204121 3'UTR clone of NM_001159629 The sequence shown below is from the reference sequence of NM_001159629. The complete sequence of this clone may contain minor differences, such as SNPs. Blue=Stop Codon Red=Cloning site GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC AATTTGCATACCCGTAAAGGGAGACTTTTTTAAATAACAGTTGAGTCTTTGCAAGTAAAAAGATTTAGA GATTATTATTTTCAGTGTGCACCTACTGTTTGTATTTGCAAACTGAGCTTGTTGGAGGGAAGGCATTA TTTTTTAAAATACTTAGTAAATTAAATGAACACCAACATGTGA 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. NM 001159629.2 RefSeq:



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	FATP2 (SLC27A2) (NM_001159629) Human 3' UTR Clone – SC204121
Summary:	The protein encoded by this gene is an isozyme of long-chain fatty-acid-coenzyme A ligase family. Although differing in substrate specificity, subcellular localization, and tissue distribution, all isozymes of this family convert free long-chain fatty acids into fatty acyl-CoA esters, and thereby play a key role in lipid biosynthesis and fatty acid degradation. This isozyme activates long-chain, branched-chain and very-long-chain fatty acids containing 22 or more carbons to their CoA derivatives. It is expressed primarily in liver and kidney, and is present in both endoplasmic reticulum and peroxisomes, but not in mitochondria. Its decreased peroxisomal enzyme activity is in part responsible for the biochemical pathology in X-linked adrenoleukodystrophy. Alternatively spliced transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Apr 2009]
Locus ID:	11001
MW:	12.2

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