

Product datasheet for RC222647L3V

OriGene Technologies, Inc.

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CYP7B1 (NM_004820) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: CYP7B1 (NM_004820) Human Tagged ORF Clone Lentiviral Particle

Symbol: CYP7B1

Synonyms: CBAS3; CP7B; SPG5A

Mammalian Cell

Selection:

Puromycin

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

Tag: Myc-DDK
ACCN: NM 004820

ORF Size: 1518 bp

ORF Nucleotide

The OR

OTI Disclaimer:

Sequence:

The ORF insert of this clone is exactly the same as(RC222647).

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.

RefSeg: NM 004820.2

RefSeq Size: 2395 bp
RefSeq ORF: 1521 bp
Locus ID: 9420

 UniProt ID:
 O75881

 Cytogenetics:
 8q12.3

Domains: p450

Protein Families: Druggable Genome, P450, Transmembrane





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Protein Pathways: Primary bile acid biosynthesis

MW: 58.3 kDa

Gene Summary: This gene encodes a member of the cytochrome P450 superfamily of enzymes. The

cytochrome P450 proteins are monooxygenases which catalyze many reactions involved in drug metabolism and synthesis of cholesterol, steroids and other lipids. This endoplasmic reticulum membrane protein catalyzes the first reaction in the cholesterol catabolic pathway of extrahepatic tissues, which converts cholesterol to bile acids. This enzyme likely plays a minor role in total bile acid synthesis, but may also be involved in the development of atherosclerosis, neurosteroid metabolism and sex hormone synthesis. Mutations in this gene

have been associated with hereditary spastic paraplegia (SPG5 or HSP), an autosomal recessive disorder. [provided by RefSeq, Apr 2016]