

Product datasheet for RC209042L4V

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Cytochrome P450 17A1 (CYP17A1) (NM 000102) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Cytochrome P450 17A1 (CYP17A1) (NM_000102) Human Tagged ORF Clone Lentiviral Particle

Symbol: Cytochrome P450 17A1

Synonyms: CPT7; CYP17; P450C17; S17AH

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

ACCN: NM_000102 **ORF Size:** 1524 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC209042).

OTI Disclaimer:

Sequence:

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 000102.2

 RefSeq Size:
 1755 bp

 RefSeq ORF:
 1527 bp

 Locus ID:
 1586

 UniProt ID:
 P05093

 Cytogenetics:
 10q24.32

Domains: p450

Protein Families: Druggable Genome, P450





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Protein Pathways: C21-Steroid hormone metabolism, Metabolic pathways

MW: 57.2 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 protein localizes to the endoplasmic reticulum. It has both 17alpha-hydroxylase and 17,20-lyase activities and is a key enzyme in the steroidogenic pathway that produces progestins, mineralocorticoids, glucocorticoids, androgens, and estrogens. Mutations in this gene are associated with isolated steroid-17 alpha-hydroxylase deficiency, 17-alpha-hydroxylase/17,20-lyase

deficiency, pseudohermaphroditism, and adrenal hyperplasia. [provided by RefSeq, Jul 2008]