

## Product datasheet for RC221240L4V

## OriGene Technologies, Inc.

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## CYP11B1 (NM\_001026213) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** CYP11B1 (NM\_001026213) Human Tagged ORF Clone Lentiviral Particle

Symbol: CYP11B1

Synonyms: CPN1; CYP11B; FHI; P450C11

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_001026213

ORF Size: 1311 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC221240).

Sequence:

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 001026213.1, NP 001021384.1

 RefSeq Size:
 3353 bp

 RefSeq ORF:
 1314 bp

 Locus ID:
 1584

 UniProt ID:
 P15538

 Cytogenetics:
 8q24.3

**Protein Families:** Druggable Genome, P450

**Protein Pathways:** Androgen and estrogen metabolism, C21-Steroid hormone metabolism, Metabolic pathways





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**MW:** 49.7 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 mitochondrial inner membrane and is involved in the conversion of progesterone to cortisol in the adrenal cortex. Mutations in this gene cause congenital adrenal hyperplasia due to 11-beta-hydroxylase deficiency. Transcript variants encoding different isoforms have

been noted for this gene. [provided by RefSeq, Jul 2008]