

Product datasheet for SC206853

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CYP21A2 (NM_001128590) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: CYP21A2 (NM_001128590) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

Symbol: CYP21A2

Synonyms: CA21H; CAH1; CPS1; CYP21B; P450c21B

ACCN: NM_001128590

Insert Size: 538 bp

Insert Sequence: >SC206853 3'UTR clone of NM_001128590

The sequence shown below is from the reference sequence of NM_001128590. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

GGGGCCCACAGCCCGGGCCAGAGCCAGTGATGGGGCAGGACCGATGCCAGCCGGGTACCTCAGTTTCTC CTTTATTGCTCCCGTACGAACCCCTCCCCCCCTGTAAACACAGTGCTGCGAGATCGCTGGCAGAG AAGGCTTCCTCCAGCGGCTGGTGGAGAGGACCCTTCCTCTCGGGGCGACCCCTCAGTGCTCG GCAGTCATACTGGGGTGCGAGAGAGGTGGGCAGCAGCACCCTCAGCCTCCCCCCGCTGGGGAGCGAAAGTTTC TTGGTCTCAGCTTCATTTCCGTGAAGGGCACCGAGAACTCGAAGCCCTTCCAGTGGTACCAGCTCACTC CCTGGGAAAAGGGGTTGTCAAGAGAGAGTCAAAGCCGGATGTCCCATCTGCTCTTCCCGTTCCCCTTAAG GAGGTAGCTCCCACCACCACCAACCCTCCCCCGCAGAGCTCCCTTCCTGACCCTCCGCTGCAGAGGATT

GAGGCTTAATTCTGAGCTGGCCCTTTCCAGCCAATAAATCAACTCCAGCTCCCTC

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.

RefSeg: NM 001128590.4







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 and hydroxylates steroids at the 21 position. Its activity is required for the synthesis of steroid hormones including cortisol and aldosterone. Mutations in this gene cause congenital adrenal hyperplasia. A related pseudogene is located near this gene; gene conversion events involving the functional gene and the pseudogene are thought to account for many cases of steroid 21-hydroxylase deficiency. Two transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul 2008]

Locus ID: 1589 **MW:** 18.7