

Product datasheet for SC328540

CYB5R3 (NM_001171660) Human Untagged Clone

Product data:

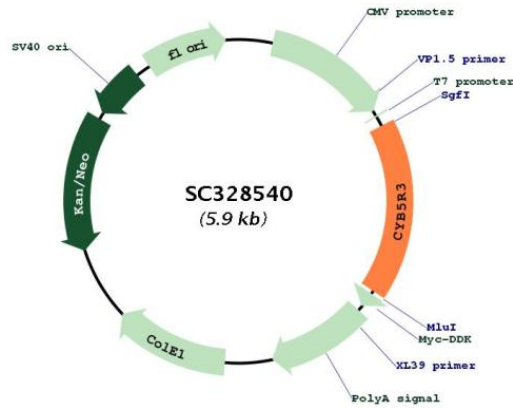
Product Type:	Expression Plasmids
Product Name:	CYB5R3 (NM_001171660) Human Untagged Clone
Tag:	Tag Free
Symbol:	CYB5R3
Synonyms:	B5R; DIA1
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
Fully Sequenced ORF:	>SC328540 representing NM_001171660. Blue=Insert sequence Red=Cloning site Green=Tag(s)

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GCTCGTTTGTGAACCGTCAGAATTTGTAAACGACTACTATAGGGCGCCGGGAATTCGTCGACTG
GATCCGGTACCGAGGAGATCTGCCGCCGCGATCGCC
ATGAATAGGAGTTTGCTGGTTGGATGCATGCAGAGCAAGGACATTTGGGGCAGAGAGGAGAGCATATGC
GAACGCCTGAAGCAAGATGGGCTTGATGTTGAAAGAGCAGAGAGCTGGGAGTTGGGCCATATGGTGCTC
TTCCCAGTCTGGTTCCTGTACAGTCTGCTCATGAAGCTGTTCCAGCGCTCCACGCCAGCCATCACCTC
GAGAGCCCGGACATCAAGTACCCGCTGCGGCTCATCGACCGGAGATCATCAGCCATGACACCCGGCGC
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CAGTACCTGGAGAGCATGCAGATTGGAGACACCATTGAGTTCCGGGGCCCAAGTGGGCTGCTGGTCTAC
CAGGGCAAAGGAAGTTCGCCATCCGACCTGACAAAAAGTCCAACCCATCATCAGGACAGTGAAGTCT
GTGGGCATGATCGCGGGAGGGACAGGCATACCCCGATGCTGCAGGTGATCCGCGCCATCATGAAGGAC
CCTGATGACCACACTGTGTGCCACCTGCTTTGCCAACCGAGCCGAGAAGGACATCCTGCTGCGACCT
GAGCTGGAGGAACCTCAGGAACAAACATTCTGCACGCTTCAAGCTCTGGTACACGCTGGACAGAGCCCT
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GAGGAGCCGCTGGTCTGATGTGTGGCCCCACCCATGATCCAGTACGCTGCCCTCCCAACCTGGAC
CAGTGGGCCACCCACGGAGCGCTGCTTCGTTCTGA
ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGAT
TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC
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Restriction Sites: Sgfl-Mlul



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Plasmid Map:


ACCN: NM_001171660

Insert Size: 1005 bp

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative RNA splicing form or single nucleotide polymorphism (SNP).

OTI Annotation: This TrueClone is provided through our Custom Cloning Process that includes sub-cloning into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method:

1. Centrifuge at 5,000xg for 5min.
2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.
3. Close the tube and incubate for 10 minutes at room temperature.
4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.
5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.

RefSeq: [NM_001171660.1](#)

RefSeq Size: 3063 bp

RefSeq ORF: 1005 bp

Locus ID: 1727

UniProt ID:	P00387
Cytogenetics:	22q13.2
Protein Families:	Druggable Genome
Protein Pathways:	Amino sugar and nucleotide sugar metabolism
MW:	38.2 kDa

Gene Summary: This gene encodes cytochrome b5 reductase, which includes a membrane-bound form in somatic cells (anchored in the endoplasmic reticulum, mitochondrial and other membranes) and a soluble form in erythrocytes. The membrane-bound form exists mainly on the cytoplasmic side of the endoplasmic reticulum and functions in desaturation and elongation of fatty acids, in cholesterol biosynthesis, and in drug metabolism. The erythrocyte form is located in a soluble fraction of circulating erythrocytes and is involved in methemoglobin reduction. The membrane-bound form has both membrane-binding and catalytic domains, while the soluble form has only the catalytic domain. Alternate splicing results in multiple transcript variants. Mutations in this gene cause methemoglobinemias. [provided by RefSeq, Jan 2010]

Transcript Variant: This variant (5) differs in the 5' UTR and 5' coding region and initiates translation at an alternate start codon, compared to variant 1. The encoded isoform (3) is longer and has a distinct N-terminus, compared to isoform 1. Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments.