

## **Product datasheet for SC203639**

## HMBS (NM 000190) Human 3' UTR Clone

## **Product data:**

**Product Type:** 3' UTR Clones

Product Name: HMBS (NM\_000190) Human 3' UTR Clone

Symbol: HMBS

Synonyms: PBG-D; PBGD; PORC; UPS

**Mammalian Cell** 

Selection:

Neomycin

**Vector:** pMirTarget (PS100062)

**ACCN:** NM\_000190

**Insert Size:** 296 bp

Insert Sequence: >SC203639 3'UTR clone of NM\_000190

The sequence shown below is from the reference sequence of NM\_000190. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

GTTGCACGGCAGCTTAACGATGCCCATTAACTGGTTTGTGGGGCACAGATGCCTGGGTTGCTGCTCCAGTGCCTACATCCCGGGCCTCAGTGCCCCATTCTCACTGCTATCTGGGGAGTGATTACCCCGGGAGACTGAACTGCAGGGTTCAAGCCTTCCAGGGATTTGCCTCACCTTGGGGCCTTGATGACTGCCTTGCCTCCTCAGTATGTGGGGGCTTCATCTCTTTAGAGAAGTCCAAGCAACAGCCTTTGAATGTAACCAATCCTACTAA

TAAACCAGTTCTGAAGGTGT

**ACGCGT**AAGCGGCCGCGCATCTAGATTCGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA

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.

RefSeq: <u>NM 000190.4</u>



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## HMBS (NM\_000190) Human 3' UTR Clone - SC203639

Summary: This gene encodes a member of the hydroxymethylbilane synthase superfamily. The encoded

protein is the third enzyme of the heme biosynthetic pathway and catalyzes the head to tail condensation of four porphobilinogen molecules into the linear hydroxymethylbilane. Mutations in this gene are associated with the autosomal dominant disease acute

intermittent porphyria. Alternatively spliced transcript variants encoding different isoforms

have been described. [provided by RefSeq, Jul 2008]

**Locus ID:** 3145 **MW:** 10.5