

## **Product datasheet for SC203708**

## PYGM (NM 005609) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

**Product Name:** PYGM (NM 005609) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: PYGM Synonyms: GSD5

**ACCN:** NM\_005609

**Insert Size:** 291 bp

Insert Sequence: >SC203708 3'UTR clone of NM\_005609

The sequence shown below is from the reference sequence of NM\_005609. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

CGCCTGCCAGCCCCGGATGAGGCCATCTGAGCCTCCAGACCAGACCCCAAACCAGCCCTTGAGTCTGTC ACACTCTCTTGGGCCAGCCCCAGCACCTCATGCAGAGGGTGGGGTACTGGAGTTAGATCTCTAAGCCCC TCCTGGAACCCTCATTTTCCCCACTCTCAATGTCCCAGTGTCCAGCGTGACTAAGGACACGGGCCCCCT TCCGTCCTCGGGCTCCCGGTCCCCTATTTATGGGGTCTGACCAACTGCACCCACTCCCTAATAAAT

TCATTCTCCTTGGGA

**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.

**RefSeg:** NM 005609.4



**OriGene Technologies, Inc.** 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



## PYGM (NM\_005609) Human 3' UTR Clone - SC203708

**Summary:** This gene encodes a muscle enzyme involved in glycogenolysis. Highly similar enzymes

encoded by different genes are found in liver and brain. Mutations in this gene are associated with McArdle disease (myophosphorylase deficiency), a glycogen storage disease of muscle. Alternative splicing results in multiple transcript variants.[provided by RefSeq, Sep 2009]

**Locus ID:** 5837 **MW:** 10.4