

Product datasheet for SC204529

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Glucosidase 2 subunit beta (PRKCSH) (NM_002743) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: Glucosidase 2 subunit beta (PRKCSH) (NM_002743) Human 3' UTR Clone

Symbol: Glucosidase 2 subunit beta

Synonyms: AGE-R2; G19P1; GIIB; PCLD; PCLD1; PKCSH; PLD1; VASAP-60

Mammalian Cell

Selection:

Neomycin

Vector: pMirTarget (PS100062)

ACCN: NM_002743

Insert Size: 383 bp

Insert Sequence: >SC204529 3'UTR clone of NM_002743

The sequence shown below is from the reference sequence of NM_002743. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

AATAAATGTGATCCCCCACCCAAAAAAAAAAAAAAAAA

ACGCGTAAGCGGCCGCGCATCTAGATTCGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA

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.





Glucosidase 2 subunit beta (PRKCSH) (NM_002743) Human 3' UTR Clone - SC204529

RefSeq: <u>NM 002743.3</u>

Summary: This gene encodes the beta-subunit of glucosidase II, an N-linked glycan-processing enzyme

in the endoplasmic reticulum. The encoded protein is an acidic phosphoprotein known to be a substrate for protein kinase C. Mutations in this gene have been associated with the autosomal dominant polycystic liver disease. Alternative splicing results in multiple transcript

variants. [provided by RefSeq, Jan 2014]

Locus ID: 5589 **MW:** 13.3