

Product datasheet for SC205246

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Glypican 3 (GPC3) (NM 004484) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: Glypican 3 (GPC3) (NM 004484) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

GPC3 Symbol:

DGSX; GTR2-2; MXR7; OCI-5; SDYS; SGB; SGBS; SGBS1 Synonyms:

ACCN: NM 004484

Insert Size: 409 bp

>SC205246 3'UTR clone of NM_004484 **Insert Sequence:**

The sequence shown below is from the reference sequence of NM_004484. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

GTGGTGTGCTTCTTCCTGGTGCACTGACTGCCTGGTGCCCAGCACATGTGCTGCCCTACAGCACCC TACCTCCTCCAGCCATGAAGTAGAGGACTAACCATGTGTTATGTTTTCGAAAATCAAATGGTATCTTTT

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.

NM 004484.4 RefSeq:





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Summary:

Cell surface heparan sulfate proteoglycans are composed of a membrane-associated protein core substituted with a variable number of heparan sulfate chains. Members of the glypican-related integral membrane proteoglycan family (GRIPS) contain a core protein anchored to the cytoplasmic membrane via a glycosyl phosphatidylinositol linkage. These proteins may play a role in the control of cell division and growth regulation. The protein encoded by this gene can bind to and inhibit the dipeptidyl peptidase activity of CD26, and it can induce apoptosis in certain cell types. Deletion mutations in this gene are associated with Simpson-Golabi-Behmel syndrome, also known as Simpson dysmorphia syndrome. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Sep 2009]

Locus ID: 2719 MW: 15.8