

Product datasheet for SC334831

SLC35C2 (NM 001281457) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: SLC35C2 (NM_001281457) Human Untagged Clone

Tag: Tag Free Symbol: SLC35C2

Synonyms: BA394O2.1; C20orf5; CGI-15; OVCOV1

Mammalian Cell

Selection:

Neomycin

Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

Fully Sequenced ORF: >NCBI ORF sequence for NM_001281457, the custom clone sequence may differ by one or

more nucleotides

Restriction Sites: Sgfl-Mlul

ACCN: NM 001281457

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).



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SLC35C2 (NM_001281457) Human Untagged Clone - SC334831

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: <u>NM 001281457.1</u>, <u>NP 001268386.1</u>

 RefSeq Size:
 2254 bp

 RefSeq ORF:
 756 bp

 Locus ID:
 51006

 UniProt ID:
 Q9NQQ7

 Cytogenetics:
 20q13.12

Protein Families: Druggable Genome, Transmembrane

Gene Summary: This gene encodes a member of the triose-phosphate transporter protein family. This gene is

regulated by oxygen tension, is induced in hypoxic trophoblast cells, and is overexpressed in ovarian cancer. Alternative splicing results in multiple transcript variants. A pseudogene of

this gene has been defined on the X chromosome. [provided by RefSeq, Jul 2013]
Transcript Variant: This variant (4) differs in the 5' UTR and initiates translation at a

downstream start codon, compared to variant 1. The encoded isoform (c) has a shorter N-

terminus, compared to isoform a.