

Product datasheet for SC333696

PGAP2 (NM 001283040) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: PGAP2 (NM_001283040) Human Untagged Clone

Tag: Tag Free Symbol: PGAP2

Synonyms: CWH43-N; FRAG1; HPMRS3; MRT17; MRT21

Mammalian Cell

Selection:

Neomycin

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

Fully Sequenced ORF: >SC333696 representing NM_001283040.

Blue=Insert sequence Red=Cloning site Green=Tag(s)

GATCCGGTACCGAGGAGATCTGCCGCCGCGATCGCC

ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGAT

TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC

Restriction Sites: Sgfl-Mlul

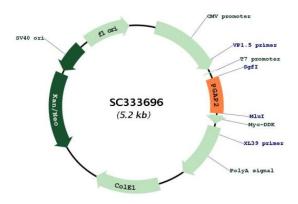
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Plasmid Map:



ACCN: NM_001283040

Insert Size: 369 bp

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).

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 001283040.1</u>



PGAP2 (NM_001283040) Human Untagged Clone - SC333696

RefSeq Size: 1460 bp
RefSeq ORF: 369 bp
Locus ID: 27315
UniProt ID: Q9UHJ9
Cytogenetics: 11p15.4

Protein Families: Druggable Genome, Transmembrane

MW: 13.1 kDa

Gene Summary: The protein encoded by this gene plays a role in the maturation of

glycosylphosphatidylinositol (GPI) anchors on GPI-anchored proteins. Mutations in this gene are associated with an autosomal recessive syndrome characterized by hyperphosphatasia

and intellectual disability. [provided by RefSeq, Jul 2017]

Transcript Variant: This variant (19) uses an alternate exon structure in the 5' region

compared to variant 1. The encoded isoform (11) is distinct and shorter, compared to isoform

1, but it shares regions in common with isoforms 5 and 7.