

Product datasheet for SC330236

FRMD3 (NM 001244962) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: FRMD3 (NM_001244962) Human Untagged Clone

Tag: Tag Free
Symbol: FRMD3

Synonyms: 4.10; EPB41L40; EPB41L0; P410

Vector: pCMV6-Entry (PS100001)

Fully Sequenced ORF: >SC330236 representing NM_001244962.

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

GAGTCAGTCTCCATGCAGTAA

Restriction Sites: Sgfl-Mlul

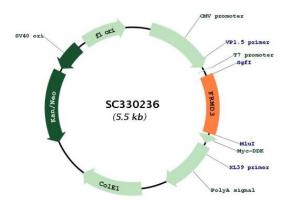
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Plasmid Map:



ACCN: NM_001244962

Insert Size: 642 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 001244962.1</u>

 RefSeq Size:
 982 bp

 RefSeq ORF:
 642 bp

 Locus ID:
 257019

 UniProt ID:
 A2A2Y4

 Cytogenetics:
 9q21.32

Protein Families: Transmembrane

MW: 23.9 kDa

Gene Summary: The protein encoded by this gene is a single pass membrane protein primarily found in

ovaries. A similar protein in erythrocytes helps determine the shape of red blood cells, but the function of the encoded protein has not been determined. There is some evidence that this is a tumor suppressor gene, and there is also evidence linking defects in this gene to susceptibility to diabetic nephropathy in type 1 diabetes. Several transcript variants encoding

different isoforms have been found for this gene. [provided by RefSeq, Oct 2011]

Transcript Variant: This variant (5) lacks the first exon compared to variant 1 and has an alternate, downstream first exon instead. It also uses an alternate splice junction at the 3' end of an exon and has an alternate 3' terminal exon compared to variant 1. The resulting isoform

(5) has shorter and distinct N and C-terminus compared to isoform 1.