

Product datasheet for **SC330054**

DEPDC5 (NM_001242897) Human Untagged Clone

Product data:

Product Type: Expression Plasmids
Product Name: DEPDC5 (NM_001242897) Human Untagged Clone
Tag: Tag Free
Symbol: DEPDC5
Synonyms: DEP.5; FFEVF; FFEVF1
Vector: pCMV6-Entry (PS100001)
Fully Sequenced ORF: >SC330054 representing NM_001242897.
 Blue=Insert sequence Red=Cloning site Green=Tag(s)

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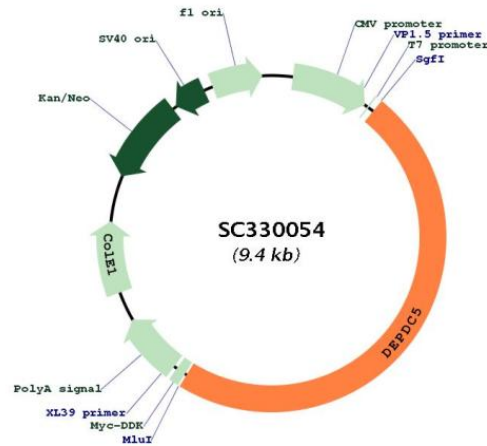


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Restriction Sites:

SgfI-MluI

Plasmid Map:


ACCN: NM_001242897

Insert Size: 4512 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: [NM_001242897.1](#)

RefSeq Size: 5155 bp

RefSeq ORF: 4512 bp

Locus ID: 9681

UniProt ID: [O75140](#)

Cytogenetics: 22q12.2-q12.3

MW: 170.5 kDa

Gene Summary: This gene encodes a member of the IML1 family of proteins involved in G-protein signaling pathways. The mechanistic target of rapamycin complex 1 (mTORC1) pathway regulates cell growth by sensing the availability of nutrients. The protein encoded by this gene is a component of the GATOR1 (GAP activity toward Rags) complex which inhibits the amino acid-sensing branch of the mTORC1 pathway. Mutations in this gene are associated with autosomal dominant familial focal epilepsy with variable foci. A single nucleotide polymorphism in an intron of this gene has been associated with an increased risk of hepatocellular carcinoma in individuals with chronic hepatitis C virus infection. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Mar 2014]
Transcript Variant: This variant (5) differs in the 5' UTR and lacks three in-frame exons in the coding region compared to variant 4. The encoded protein (isoform 5) is shorter than isoform 4.