

## **Product datasheet for SC335303**

### OriGene Technologies, Inc.

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# Apolipoprotein E (APOE) (NM\_001302690) Human Untagged Clone

### **Product data:**

**Product Type:** Expression Plasmids

**Product Name:** Apolipoprotein E (APOE) (NM\_001302690) Human Untagged Clone

Tag: Tag Free Symbol: APOE

Synonyms: AD2; APO-E; ApoE4; LDLCQ5; LPG

**Vector:** pCMV6-Entry (PS100001)

Fully Sequenced ORF: >SC335303 representing NM\_001302690.

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

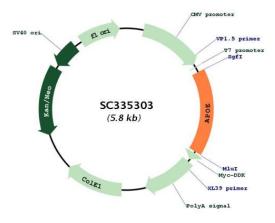
AAGGTGCAGGCTGCCGTGGGCACCAGCGCCCCCTGTGCCCAGCGACAATCAC<mark>TGA</mark>

**Restriction Sites:** Sgfl-Mlul





#### Plasmid Map:



**ACCN:** NM\_001302690

**Insert Size:** 954 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 001302690.1</u>

 RefSeq Size:
 1265 bp

 RefSeq ORF:
 954 bp

 Locus ID:
 348

 UniProt ID:
 P02649

 Cytogenetics:
 19q13.32

**Protein Families:** Adult stem cells, Druggable Genome, Secreted Protein, Stem cell - Pluripotency

**Protein Pathways:** Alzheimer's disease

**MW:** 36.2 kDa

**Gene Summary:** The protein encoded by this gene is a major apoprotein of the chylomicron. It binds to a

specific liver and peripheral cell receptor, and is essential for the normal catabolism of

triglyceride-rich lipoprotein constituents. This gene maps to chromosome 19 in a cluster with

the related apolipoprotein C1 and C2 genes. Mutations in this gene result in familial

dysbetalipoproteinemia, or type III hyperlipoproteinemia (HLP III), in which increased plasma cholesterol and triglycerides are the consequence of impaired clearance of chylomicron and

VLDL remnants. [provided by RefSeq, Jun 2016]

Transcript Variant: This variant (4) contains an alternate 5' terminal exon, and it thus differs in the 5' UTR and initiates translation at a downstream in-frame start codon, compared to

variant 1. The encoded isoform (b) is shorter at the N-terminus, compared to isoform a.

Variants 2, 3, 4 and 5 all encode isoform b.