

Product datasheet for SC335371

C2 (NM_001282459) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: C2 (NM_001282459) Human Untagged Clone

Tag: Tag Free

Symbol: C2

Synonyms: ARMD14; CO2

Mammalian Cell Neomycin

Selection:

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

Fully Sequenced ORF: >SC335371 representing NM_001282459.

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

GATCCGGTACCGAGGAGATCTGCCGCCGCGATCGCC

ACGCGTACGCGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGAT

TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC

Restriction Sites: Sgfl-Mlul



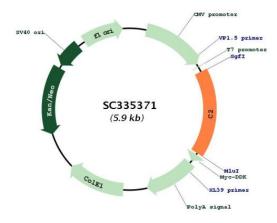
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Plasmid Map:



ACCN: NM_001282459

Insert Size: 987 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.

RefSeg: NM 001282459.1

RefSeq Size: 1916 bp
RefSeq ORF: 987 bp
Locus ID: 717

Cytogenetics: 6p21.33

Protein Families: Druggable Genome, Protease, Secreted Protein



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Protein Pathways: Complement and coagulation cascades, Systemic lupus erythematosus

MW: 35.5 kDa

Gene Summary: Component C2 is a serum glycoprotein that functions as part of the classical pathway of the

complement system. Activated C1 cleaves C2 into C2a and C2b. The serine proteinase C2a then combines with complement factor 4b to create the C3 or C5 convertase. Deficiency of C2 has been reported to associated with certain autoimmune diseases and SNPs in this gene have been associated with altered susceptibility to age-related macular degeneration. This gene localizes within the class III region of the MHC on the short arm of chromosome 6. Alternative splicing results in multiple transcript variants encoding distinct isoforms. Additional transcript variants have been described in publications but their full-length

sequence has not been determined.[provided by RefSeq, Mar 2009]

Transcript Variant: This variant (6) uses an alternate 3' exon structure, and thus differs in the 3' coding region and 3' UTR compared to variant 1. It encodes isoform 6 which is shorter and

has a distinct C-terminus, compared to isoform 1.