

Product datasheet for SC336853

Artemis (DCLRE1C) (NM_001289076) Human Untagged Clone

Product data:

Product Type: Expression Plasmids
Product Name: Artemis (DCLRE1C) (NM_001289076) Human Untagged Clone
Tag: Tag Free
Symbol: DCLRE1C
Synonyms: A-SCID; DCLREC1C; RS-SCID; SCIDA; SNM1C
Vector: pCMV6-Entry (PS100001)
Fully Sequenced ORF: >SC336853 representing NM_001289076.
 Blue=Insert sequence Red=Cloning site Green=Tag(s)

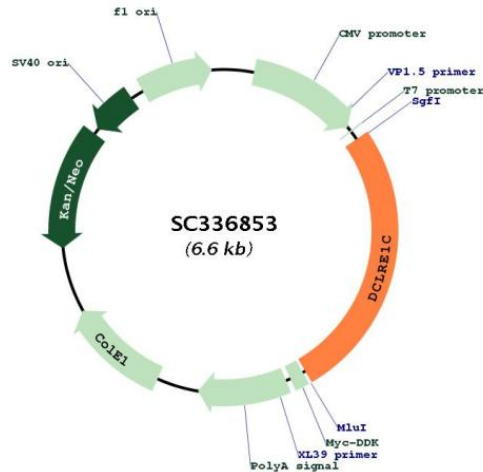
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Restriction Sites: SgfI-MluI



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Plasmid Map:


ACCN: NM_001289076

Insert Size: 1734 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_001289076.1](#)

RefSeq Size: 6157 bp

RefSeq ORF: 1734 bp

Locus ID: 64421

UniProt ID: [Q96SD1](#)

Cytogenetics: 10p13

Protein Families: Druggable Genome

Protein Pathways: Cell cycle, Non-homologous end-joining, Primary immunodeficiency

MW: 65.3 kDa

Gene Summary: This gene encodes a nuclear protein that is involved in V(D)J recombination and DNA repair. The encoded protein has single-strand-specific 5'-3' exonuclease activity; it also exhibits endonuclease activity on 5' and 3' overhangs and hairpins. The protein also functions in the regulation of the cell cycle in response to DNA damage. Mutations in this gene can cause Athabaskan-type severe combined immunodeficiency (SCIDA) and Omenn syndrome. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jan 2014]
Transcript Variant: This variant (e) lacks two exons in the 5' coding region and initiates translation at an alternate downstream start codon, compared to variant a. It encodes isoform b, which has a shorter and distinct N-terminus compared to variant a. Variants b, e, and g encode the same isoform (b). Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments.