

Product datasheet for SC312282

OriGene Technologies, Inc.

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SERAC1 (BC028594) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: SERAC1 (BC028594) Human Untagged Clone

Tag: Tag Free
Symbol: SERAC1

Synonyms: 4930511N22Rik; D17Ertd141e; FLJ14917; OTTMUSP00000028541; serine active site containing

1

Vector: pCMV6 series

Fully Sequenced ORF: >NCBI ORF sequence for BC028594, the custom clone sequence may differ by one or more

 ${\tt nucleotides}$

TGGCATGGTAATGAGACTACC

Restriction Sites: Please inquire ACCN: BC028594

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).

OTI Annotation: This TrueClone is provided through our Custom Cloning Process that includes sub-cloning

into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.

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).



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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>BC028594.1</u>, <u>AAH28594.1</u>

RefSeq Size: 4705 bp
RefSeq ORF: 504 bp
Locus ID: 84947
Cytogenetics: 6q25.3

Gene Summary: The protein encoded by this gene is a phosphatidylglycerol remodeling protein found at the

interface of mitochondria and endoplasmic reticula, where it mediates phospholipid

exchange. The encoded protein plays a major role in mitochondrial function and intracellular cholesterol trafficking. Defects in this gene are a cause of 3-methylglutaconic aciduria with deafness, encephalopathy, and Leigh-like syndrome (MEGDEL). Two transcript variants, one protein-coding and the other non-protein coding, have been found for this gene. [provided by

RefSeq, Aug 2012]