

### Product datasheet for RC224986L3V

#### OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## Small EDRK rich factor 1 (SERF1A) (NM 022968) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** Small EDRK rich factor 1 (SERF1A) (NM\_022968) Human Tagged ORF Clone Lentiviral Particle

Symbol: Small EDRK rich factor 1

4F5; FAM2A; H4F5; SERF1; SMAM1 Synonyms:

**Mammalian Cell** 

Selection:

Puromycin

Vector: pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK NM 022968 ACCN:

**ORF Size:** 186 bp

**ORF Nucleotide** 

Sequence:

The ORF insert of this clone is exactly the same as(RC224986).

The molecular sequence of this clone aligns with the gene accession number as a point of OTI Disclaimer: reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This

clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

**OTI Annotation:** This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

RefSeq: NM 022968.1

RefSeq ORF: 189 bp Locus ID: 8293 **UniProt ID:** 075920

Cytogenetics: 5q13.2

**Protein Families:** Transmembrane

7.2 kDa MW:





# Small EDRK rich factor 1 (SERF1A) (NM\_022968) Human Tagged ORF Clone Lentiviral Particle – RC224986L3V

#### **Gene Summary:**

This gene is part of a 500 kb inverted duplication on chromosome 5q13. This duplicated region contains at least four genes and repetitive elements which make it prone to rearrangements and deletions. The repetitiveness and complexity of the sequence have also caused difficulty in determining the organization of this genomic region. The duplication region includes both a telomeric and a centromeric copy of this gene. Deletions of this gene, the telomeric copy, often accompany deletions of the neighboring SMN1 gene in spinal muscular atrophy (SMA) patients, and so it is thought that this gene may be a modifier of the SMA phenotype. The function of this protein is not known; however, it bears low-level homology with the RNA-binding domain of matrin-cyclophilin, a protein which colocalizes with small nuclear ribonucleoproteins (snRNPs) and the SMN1 gene product. Alternatively spliced transcripts have been documented but it is unclear whether alternative splicing occurs for both the centromeric and telomeric copies of the gene. [provided by RefSeq, Jul 2008]