

## Product datasheet for RC206020L4V

## OriGene Technologies, Inc.

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## SART2 (DSE) (NM\_013352) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** SART2 (DSE) (NM\_013352) Human Tagged ORF Clone Lentiviral Particle

Symbol: SART2

Synonyms: DS-epi1; DSEP; DSEPI; EDSMC2; SART-2; SART2

Mammalian Cell

Selection:

Puromycin

**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

**ACCN:** NM\_013352 **ORF Size:** 2874 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC206020).

Sequence:

OTI Disclaimer:

The molecular sequence of this clone aligns with the gene accession number as a point of 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.

**RefSeg:** NM 013352.2

 RefSeq Size:
 4077 bp

 RefSeq ORF:
 2877 bp

 Locus ID:
 29940

 UniProt ID:
 Q9UL01

 Cytogenetics:
 6q22.1

**Protein Families:** Transmembrane

**Protein Pathways:** Chondroitin sulfate biosynthesis





**MW:** 109.8 kDa

**Gene Summary:** The protein encoded by this gene is a tumor-rejection antigen. It is localized to the

endoplasmic reticulum and functions to convert D-glucuronic acid to L-iduronic acid during the biosynthesis of dermatan sulfate. This antigen possesses tumor epitopes capable of inducing HLA-A24-restricted and tumor-specific cytotoxic T lymphocytes in cancer patients

and may be useful for specific immunotherapy. Mutations in this gene cause

inmusculocontractural Ehlers-Danlos syndrome. Alternative splicing results in multiple transcript variants. A related pseudogene has been identified on chromosome 9, and a

paralogous gene exists on chromosome 18. [provided by RefSeq, Apr 2016]