

# Product datasheet for MR200941L3V

### OriGene Technologies, Inc.

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## Ap4s1 (NM\_021710) Mouse Tagged ORF Clone Lentiviral Particle

#### **Product data:**

**Product Type:** Lentiviral Particles

**Product Name:** Ap4s1 (NM\_021710) Mouse Tagged ORF Clone Lentiviral Particle

Symbol: Ap4s1

AI314282 Synonyms:

**Mammalian Cell** Puromycin

Selection:

**ORF Size:** 

Sequence:

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

Tag: Myc-DDK

ACCN: NM 021710

435 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(MR200941).

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.

RefSeq: NM 021710.2

RefSeq Size: 1113 bp RefSeq ORF: 435 bp Locus ID: 11782 **UniProt ID:** Q9WVL1 Cytogenetics: 12 B3







#### **Gene Summary:**

This gene encodes the sigma subunit of the adaptor-related protein complex 4 which mediates intracellular membrane trafficking along the endocytic and secretory transport pathways. This complex contains four subunits, beta, epsilon, mu, and sigma, and belongs to a family of five adapter protein complexes, including three clathrin-associated complexes and two non clathrin-associated complexes, that localize to different intracellular compartments and mediate membrane vesicle trafficking using distinct pathways. In humans, loss-of-function mutations in this gene have been linked to specific adapter complex 4 deficiency disorders including hereditary spastic paraplegia. Alternate splicing results in multiple transcript variants. [provided by RefSeq, Jul 2016]