

## Product datasheet for RC209091L4V

## 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

## SLC25A4 (NM 001151) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** SLC25A4 (NM\_001151) Human Tagged ORF Clone Lentiviral Particle

Symbol:

AAC1; ANT; ANT 1; ANT1; MTDPS12; MTDPS12A; PEO2; PEO3; PEOA2; T1 Synonyms:

**Mammalian Cell** 

Selection:

Puromycin

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

mGFP Tag:

NM 001151 ACCN:

**ORF Size:** 894 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC209091).

Sequence:

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 001151.2

RefSeq Size: 4420 bp RefSeq ORF: 897 bp Locus ID: 291 **UniProt ID:** P12235

Cytogenetics: 4q35.1 **Domains:** 

mito\_carr

**Protein Families:** Druggable Genome, Transmembrane





## SLC25A4 (NM\_001151) Human Tagged ORF Clone Lentiviral Particle - RC209091L4V

**Protein Pathways:** Calcium signaling pathway, Huntington's disease, Parkinson's disease

MW: 33.1 kDa

**Gene Summary:** This gene is a member of the mitochondrial carrier subfamily of solute carrier protein genes.

The product of this gene functions as a gated pore that translocates ADP from the cytoplasm into the mitochondrial matrix and ATP from the mitochondrial matrix into the cytoplasm. The protein forms a homodimer embedded in the inner mitochondria membrane. Mutations in

this gene have been shown to result in autosomal dominant progressive external

opthalmoplegia and familial hypertrophic cardiomyopathy. [provided by RefSeq, Jun 2013]