

## Product datasheet for RC224118L4V

## 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

## SLC6A17 (NM\_001010898) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: SLC6A17 (NM 001010898) Human Tagged ORF Clone Lentiviral Particle

Symbol: SLC6A17

Synonyms: MRT48; NTT4

Mammalian Cell Puromycin

Selection:

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

Tag: mGFP

**ACCN:** NM 001010898

ORF Size: 2181 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC224118).

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 001010898.1

 RefSeq Size:
 6420 bp

 RefSeq ORF:
 2184 bp

 Locus ID:
 388662

 UniProt ID:
 Q9H1V8

 Cytogenetics:
 1p13.3

**Protein Families:** Druggable Genome, Transmembrane

MW: 80.8 kDa







## **Gene Summary:**

The protein encoded by this gene is a member of the SLC6 family of transporters, which are responsible for the presynaptic uptake of most neurotransmitters. The encoded vesicular transporter is selective for proline, glycine, leucine and alanine. In mouse, the strongest expression of this gene was in cortical and hippocampal tissues where expression increased during embryonic brain development and peaked postnatally. Defects in this gene cause a form of autosomal recessive intellectual disability. [provided by RefSeq, Jul 2017]