

Product datasheet for RC221273L4V

OriGene Technologies, Inc.

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Synapsin I (SYN1) (NM_006950) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Synapsin I (SYN1) (NM_006950) Human Tagged ORF Clone Lentiviral Particle

Symbol: Synapsin I

Synonyms: EPILX; MRX50; SYN1a; SYN1b; SYNI

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

ACCN: NM_006950 **ORF Size:** 2115 bp

ORF Nucleotide

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Sequence:

The ORF insert of this clone is exactly the same as(RC221273).

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 (o.g. polymorphisms), each with its own valid existence. This

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 006950.2

RefSeq Size:2248 bpRefSeq ORF:2118 bpLocus ID:6853

UniProt ID: P17600

Cytogenetics: Xp11.3-p11.23

Domains: Synapsin MW: 73.9 kDa

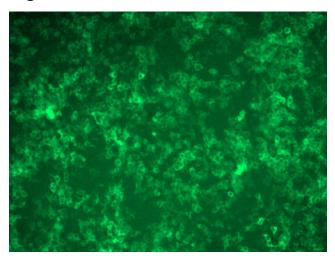




Gene Summary:

This gene is a member of the synapsin gene family. Synapsins encode neuronal phosphoproteins which associate with the cytoplasmic surface of synaptic vesicles. Family members are characterized by common protein domains, and they are implicated in synaptogenesis and the modulation of neurotransmitter release, suggesting a potential role in several neuropsychiatric diseases. This member of the synapsin family plays a role in regulation of axonogenesis and synaptogenesis. The protein encoded serves as a substrate for several different protein kinases and phosphorylation may function in the regulation of this protein in the nerve terminal. Mutations in this gene may be associated with X-linked disorders with primary neuronal degeneration such as Rett syndrome. Alternatively spliced transcript variants encoding different isoforms have been identified. [provided by RefSeq, Jul 2008]

Product images:



[RC221273L4] was used to prepare Lentiviral particles using [TR30037] packaging kit. HEK293T cells were transduced with RC221273L4V particle to overexpress human SYN1-mGFP fusion protein.