

## Product datasheet for RC227219L3V

## OriGene Technologies, Inc.

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## Choline Acetyltransferase (CHAT) (NM\_001142929) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Symbol:** Choline Acetyltransferase

**Synonyms:** CHOACTASE; CMS1A; CMS1A2; CMS6

Mammalian Cell Puromycin

Selection:

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

Tag: Myc-DDK

ACCN: NM\_001142929

ORF Size: 1890 bp

ORF Nucleotide Sequence: The ORF insert of this clone is exactly the same as(RC227219).

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:** <u>NM\_001142929.1</u>, <u>NP\_001136401.1</u>

RefSeq ORF: 1893 bp

Locus ID: 1103

UniProt ID: P28329

Cytogenetics: 10q11.23

**Protein Families:** Druggable Genome





## Choline Acetyltransferase (CHAT) (NM\_001142929) Human Tagged ORF Clone Lentiviral Particle | RC227219L3V

**Protein Pathways:** Glycerophospholipid metabolism

**MW:** 70.2 kDa

Gene Summary: This gene encodes an enzyme which catalyzes the biosynthesis of the neurotransmitter

acetylcholine. This gene product is a characteristic feature of cholinergic neurons, and changes in these neurons may explain some of the symptoms of Alzheimer's disease.

Polymorphisms in this gene have been associated with Alzheimer's disease and mild cognitive

impairment. Mutations in this gene are associated with congenital myasthenic syndrome associated with episodic apnea. Multiple transcript variants encoding different isoforms have been found for this gene, and some of these variants have been shown to encode more than

one isoform. [provided by RefSeq, May 2010]