

## Product datasheet for RC222531L4V

## OriGene Technologies, Inc.

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## Choline Acetyltransferase (CHAT) (NM\_020549) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** Choline Acetyltransferase (CHAT) (NM\_020549) Human Tagged ORF Clone Lentiviral Particle

**Symbol:** Choline Acetyltransferase

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

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_020549 **ORF Size:** 2244 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC222531).

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 020549.2, NP 065574.3

 RefSeq Size:
 2485 bp

 RefSeq ORF:
 2247 bp

 Locus ID:
 1103

 UniProt ID:
 P28329

 Cytogenetics:
 10q11.23

**Protein Families:** Druggable Genome

**Protein Pathways:** Glycerophospholipid metabolism





## Choline Acetyltransferase (CHAT) (NM\_020549) Human Tagged ORF Clone Lentiviral Particle – RC222531L4V

MW: 82.4 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]