

Product datasheet for RC227499L4V

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

Choline Acetyltransferase (CHAT) (NM_001142933) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Choline Acetyltransferase (CHAT) (NM_001142933) 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_001142933

ORF Size: 1998 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC227499).

OTI Disclaimer:

Sequence:

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: NM 001142933.1, NP 001136405.1

 RefSeq ORF:
 2001 bp

 Locus ID:
 1103

 UniProt ID:
 P28329

 Cytogenetics:
 10q11.23

Protein Families: Druggable Genome

Protein Pathways: Glycerophospholipid metabolism





Choline Acetyltransferase (CHAT) (NM_001142933) Human Tagged ORF Clone Lentiviral Particle – RC227499L4V

MW: 74.2 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]