

## Product datasheet for RC217112L4V

## 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

## C20orf7 (NDUFAF5) (NM\_024120) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: C20orf7 (NDUFAF5) (NM\_024120) Human Tagged ORF Clone Lentiviral Particle

Symbol: C20orf7

**Synonyms:** bA526K24.2; C20orf7; dJ842G6.1; MC1DN16

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_024120 **ORF Size:** 1035 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC217112).

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 024120.2

 RefSeq Size:
 1650 bp

 RefSeq ORF:
 1038 bp

 Locus ID:
 79133

 UniProt ID:
 Q5TEU4

 Cytogenetics:
 20p12.1

**Protein Families:** Druggable Genome

MW: 38.7 kDa







## **Gene Summary:**

The NADH-ubiquinone oxidoreductase complex (complex I) of the mitochondrial respiratory chain catalyzes the transfer of electrons from NADH to ubiquinone, and consists of at least 43 subunits. The complex is located in the inner mitochondrial membrane. This gene encodes a mitochondrial protein that is associated with the matrix face of the mitochondrial inner membrane and is required for complex I assembly. A mutation in this gene results in mitochondrial complex I deficiency. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Oct 2009]