

## Product datasheet for RC221871L4V

## OriGene Technologies, Inc.

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## KIAA1970 (EARS2) (NM 001083614) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** KIAA1970 (EARS2) (NM\_001083614) Human Tagged ORF Clone Lentiviral Particle

Symbol: KIAA1970

**Synonyms:** COXPD12; gluRS; MSE1; mtGlnRS

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_001083614

ORF Size: 1569 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC221871).

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

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 001083614.1

 RefSeq Size:
 3979 bp

 RefSeq ORF:
 1572 bp

 Locus ID:
 124454

 UniProt ID:
 Q5|PH6

 Cytogenetics:
 16p12.2

Protein Families: Druggable Genome

**Protein Pathways:** Aminoacyl-tRNA biosynthesis, Metabolic pathways, Porphyrin and chlorophyll metabolism





**MW:** 58.5 kDa

**Gene Summary:** 

This gene encodes a member of the class I family of aminoacyl-tRNA synthetases. These enzymes play a critical role in protein biosynthesis by charging tRNAs with their cognate amino acids. This protein is encoded by the nuclear genome but is likely to be imported to the mitochondrion where it is thought to catalyze the ligation of glutamate to tRNA molecules. Mutations in this gene have been associated with combined oxidative phosphorylation deficiency 12 (COXPD12). Alternative splicing results in multiple transcript variants. [provided by RefSeq, Mar 2015]