

# Product datasheet for RC214832L2V

### 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

## PCCA (NM\_000282) Human Tagged ORF Clone Lentiviral Particle

#### **Product data:**

Product Type: Lentiviral Particles

**Product Name:** PCCA (NM\_000282) Human Tagged ORF Clone Lentiviral Particle

Symbol: PCCA

Mammalian Cell

Selection:

None

**Vector:** pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM\_000282

ORF Size: 2184 bp

**ORF Nucleotide** 

Sequence:

The ORF insert of this clone is exactly the same as(RC214832).

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 000282.2</u>

 RefSeq Size:
 2518 bp

 RefSeq ORF:
 2187 bp

 Locus ID:
 5095

 UniProt ID:
 P05165

 Cytogenetics:
 13q32.3

**Domains:** biotin\_lipoyl, CPSase\_L\_D2, CPSase\_L\_chain, Biotin\_carb\_C

**Protein Families:** Druggable Genome

**Protein Pathways:** Metabolic pathways, Propanoate metabolism, Valine, leucine and isoleucine degradation





## PCCA (NM\_000282) Human Tagged ORF Clone Lentiviral Particle - RC214832L2V

**MW:** 80.06 kDa

**Gene Summary:** The protein encoded by this gene is the alpha subunit of the heterodimeric mitochondrial

enzyme Propionyl-CoA carboxylase. PCCA encodes the biotin-binding region of this enzyme. Mutations in either PCCA or PCCB (encoding the beta subunit) lead to an enzyme deficiency resulting in propionic acidemia. Multiple transcript variants encoding different isoforms have

been found for this gene.[provided by RefSeq, May 2010]