

# Product datasheet for RC210009L4V

### OriGene Technologies, Inc.

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## **HGD (NM\_000187) Human Tagged ORF Clone Lentiviral Particle**

#### **Product data:**

Product Type: Lentiviral Particles

**Product Name:** HGD (NM\_000187) Human Tagged ORF Clone Lentiviral Particle

Symbol: HGD

Synonyms: AKU; HGO

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_000187 **ORF Size:** 1335 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC210009).

Sequence:
OTI Disclaimer:

**Domains:** 

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 000187.2</u>

 RefSeq Size:
 2012 bp

 RefSeq ORF:
 1338 bp

 Locus ID:
 3081

 UniProt ID:
 Q93099

 Cytogenetics:
 3q13.33

**Protein Families:** Druggable Genome

HgmA





## HGD (NM\_000187) Human Tagged ORF Clone Lentiviral Particle - RC210009L4V

**Protein Pathways:** Metabolic pathways, Tyrosine metabolism

**MW:** 50 kDa

**Gene Summary:** This gene encodes the enzyme homogentisate 1,2 dioxygenase. This enzyme is involved in

the catabolism of the amino acids tyrosine and phenylalanine. Mutations in this gene are the cause of the autosomal recessive metabolism disorder alkaptonuria.[provided by RefSeq,

May 2010]