

## Product datasheet for RC204160L4V

## 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

## Prostaglandin dehydrogenase 1 (HPGD) (NM\_000860) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** Prostaglandin dehydrogenase 1 (HPGD) (NM\_000860) Human Tagged ORF Clone Lentiviral

Particle

Symbol: Prostaglandin dehydrogenase 1

Synonyms: 15-PGDH; PGDH; PGDH1; PHOAR1; SDR36C1

**Mammalian Cell** 

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_000860

ORF Size: 798 bp

**ORF Nucleotide** 

Sequence:

The ORF insert of this clone is exactly the same as(RC204160).

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 000860.3</u>

 RefSeq Size:
 3044 bp

 RefSeq ORF:
 801 bp

 Locus ID:
 3248

 UniProt ID:
 P15428

 Cytogenetics:
 4q34.1

**Domains:** adh\_short





## Prostaglandin dehydrogenase 1 (HPGD) (NM\_000860) Human Tagged ORF Clone Lentiviral Particle – RC204160L4V

**Protein Families:** Druggable Genome

MW: 29 kDa

**Gene Summary:** This gene encodes a member of the short-chain nonmetalloenzyme alcohol dehydrogenase

protein family. The encoded enzyme is responsible for the metabolism of prostaglandins, which function in a variety of physiologic and cellular processes such as inflammation. Mutations in this gene result in primary autosomal recessive hypertrophic osteoarthropathy and cranioosteoarthropathy. Multiple transcript variants encoding different isoforms have

been found for this gene. [provided by RefSeq, Mar 2009]