

## Product datasheet for RC215206L2V

## 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

## IMPDH1 (NM\_000883) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** IMPDH1 (NM\_000883) Human Tagged ORF Clone Lentiviral Particle

Symbol: IMPDH<sup>\*</sup>

Synonyms: IMPD; IMPD1; IMPDH-I; LCA11; RP10; sWSS2608

Mammalian Cell

Selection:

None

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

Tag: mGFP

**ACCN:** NM\_000883 **ORF Size:** 1797 bp

**ORF Nucleotide** 

- - -

Sequence:

The ORF insert of this clone is exactly the same as(RC215206).

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

variants is recommended prior to use. <u>infore info</u>

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 000883.2</u>

 RefSeq Size:
 2880 bp

 RefSeq ORF:
 1800 bp

 Locus ID:
 3614

 UniProt ID:
 P20839

 Cytogenetics:
 7q32.1

Domains: CBS, IMPDH

**Protein Families:** Druggable Genome





## IMPDH1 (NM\_000883) Human Tagged ORF Clone Lentiviral Particle - RC215206L2V

**Protein Pathways:** Drug metabolism - other enzymes, Metabolic pathways, Purine metabolism

**MW:** 64.1 kDa

**Gene Summary:** The protein encoded by this gene acts as a homotetramer to regulate cell growth. The

encoded protein is an enzyme that catalyzes the synthesis of xanthine monophosphate (XMP) from inosine-5'-monophosphate (IMP). This is the rate-limiting step in the de novo synthesis of guanine nucleotides. Defects in this gene are a cause of retinitis pigmentosa type 10 (RP10). Several transcript variants encoding different isoforms have been found for this gene.

[provided by RefSeq, Dec 2008]