

## Product datasheet for RC219336L4V

## 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

## IDUA (NM\_000203) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

Product Name: IDUA (NM 000203) Human Tagged ORF Clone Lentiviral Particle

Symbol: IDUA

Synonyms: IDA; MPS1; MPSI

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_000203 **ORF Size:** 1959 bp

**ORF Nucleotide** 

. . . . . .

Sequence:
OTI Disclaimer:

The ORF insert of this clone is exactly the same as(RC219336).

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.

**RefSeg:** NM 000203.2

 RefSeq Size:
 2197 bp

 RefSeq ORF:
 1962 bp

 Locus ID:
 3425

 UniProt ID:
 P35475

 Cytogenetics:
 4p16.3

**Protein Families:** Druggable Genome

**Protein Pathways:** Glycosaminoglycan degradation, Lysosome, Metabolic pathways



## IDUA (NM\_000203) Human Tagged ORF Clone Lentiviral Particle - RC219336L4V

**MW:** 72.67 kDa

**Gene Summary:** This gene encodes an enzyme that hydrolyzes the terminal alpha-L-iduronic acid residues of

two glycosaminoglycans, dermatan sulfate and heparan sulfate. This hydrolysis is required for the lysosomal degradation of these glycosaminoglycans. Mutations in this gene that result in enzymatic deficiency lead to the autosomal recessive disease mucopolysaccharidosis type I

(MPS I). [provided by RefSeq, Jul 2008]