

## Product datasheet for RC219187L1V

## OriGene Technologies, Inc.

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## Iduronate 2 sulfatase (IDS) (NM 000202) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** Iduronate 2 sulfatase (IDS) (NM\_000202) Human Tagged ORF Clone Lentiviral Particle

Symbol: Iduronate 2 sulfatase

ID2S: MPS2: SIDS Synonyms:

**Mammalian Cell** 

Selection:

ACCN:

None

Vector: pLenti-C-Myc-DDK (PS100064)

Myc-DDK Tag: NM 000202

**ORF Size:** 1650 bp

**ORF Nucleotide** 

Sequence:

The ORF insert of this clone is exactly the same as(RC219187).

The molecular sequence of this clone aligns with the gene accession number as a point of OTI Disclaimer: 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: NM 000202.2

RefSeq Size: 2504 bp RefSeq ORF: 1653 bp Locus ID: 3423 **UniProt ID:** P22304

Cytogenetics: Xq28

**Domains:** Sulfatase

**Protein Families:** Druggable Genome





## Iduronate 2 sulfatase (IDS) (NM\_000202) Human Tagged ORF Clone Lentiviral Particle – RC219187L1V

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

MW: 61.87 kDa

**Gene Summary:** This gene encodes a member of the sulfatase family of proteins. The encoded preproprotein

is proteolytically processed to generate two polypeptide chains. This enzyme is involved in the lysosomal degradation of heparan sulfate and dermatan sulfate. Mutations in this gene are associated with the X-linked lysosomal storage disease mucopolysaccharidosis type II, also known as Hunter syndrome. Alternative splicing results in multiple transcript variants, at least one of which encodes a preproprotein that is proteolytically processed. [provided by

RefSeq, Jan 2016]