

## Product datasheet for RC203908L3V

## OriGene Technologies, Inc.

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## Aldolase (ALDOA) (NM\_184043) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** Aldolase (ALDOA) (NM\_184043) Human Tagged ORF Clone Lentiviral Particle

Symbol: Aldolase

**Synonyms:** ALDA; GSD12; HEL-S-87p

Mammalian Cell

Selection:

Puromycin

**Vector:** pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK
ACCN: NM 184043

ORF Size: 1092 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC203908).

Sequence:

**UniProt ID:** 

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.

**RefSeg:** NM 184043.1

RefSeq Size: 1569 bp
RefSeq ORF: 1095 bp
Locus ID: 226

Cytogenetics: 16p11.2

**Protein Families:** Druggable Genome

P04075





## Aldolase (ALDOA) (NM\_184043) Human Tagged ORF Clone Lentiviral Particle - RC203908L3V

**Protein Pathways:** Fructose and mannose metabolism, Glycolysis / Gluconeogenesis, Metabolic pathways,

Pentose phosphate pathway

**MW:** 39.4 kDa

**Gene Summary:** This gene encodes a member of the class I fructose-bisphosphate aldolase protein family.

The encoded protein is a glycolytic enzyme that catalyzes the reversible conversion of fructose-1,6-bisphosphate to glyceraldehyde 3-phosphate and dihydroxyacetone phosphate. Three aldolase isozymes (A, B, and C), encoded by three different genes, are differentially expressed during development. Mutations in this gene have been associated with Glycogen Storage Disease XII, an autosomal recessive disorder associated with hemolytic anemia. Disruption of this gene also plays a role in the progression of multiple types of cancers. Related pseudogenes have been identified on chromosomes 3 and 10. [provided by RefSeq,

Sep 2017]