

## Product datasheet for TR506868

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com

OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

EU: info-de@origene.com CN: techsupport@origene.cn

## Aldob Mouse shRNA Plasmid (Locus ID 230163)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Aldob Mouse shRNA Plasmid (Locus ID 230163)

Locus ID:

Al; Ald; Aldo-2; Aldo2; BC016435 Synonyms:

Vector: pRS (TR20003)

E. coli Selection: Ampicillin Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Aldob - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = Components:

230163). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

BC016435, BC022113, BC024056, BC024112, BC026577, BC030724, BC030725, BC034169, RefSeq:

BC034171, BC034172, BC034173, BC036130, BC036131, BC036132, BC036133, NM 144903,

NM 144903.1, NM 144903.2, NM 144903.3, BC036139, BC036142

UniProt ID: Q91Y97

Summary: This gene encodes a subunit of the homotetrameric enzyme aldolase B, an isozyme of the

> class I fructose 1,6-bisphosphate aldolase enzyme. This enzyme catalyzes the conversion of fructose 1,6-bisphosphate to dihydroxyacetone phosphate and glyceraldehyde 3-phosphate. Homozygous knockout mice for this gene exhibit liver damage and death following fructose ingestion. A pseudogene of this gene has been identified in the genome. [provided by RefSeq,

Aug 2015]

These shRNA constructs were designed against multiple splice variants at this gene locus. To shRNA Design:

> be certain that your variant of interest is targeted, please contact <a href="techsupport@origene.com">techsupport@origene.com</a>. If you need a special design or shRNA sequence, please utilize our custom shRNA service.





## Performance Guaranteed:

OriGene guarantees that the sequences in the shRNA expression cassettes are verified to correspond to the target gene with 100% identity. One of the four constructs at minimum are guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must be used in comparison with the target-specific shRNA transfected samples.

For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).