

## Product datasheet for **TP302576M**

### Aldolase (ALDOA) (NM\_000034) Human Recombinant Protein

#### Product data:

Product Type:	Recombinant Proteins
Description:	Recombinant protein of human aldolase A, fructose-bisphosphate (ALDOA), transcript variant 1, 100 µg
Species:	Human
Expression Host:	HEK293T
Expression cDNA Clone or AA Sequence:	>RC202576 protein sequence <b>Red</b> =Cloning site <b>Green</b> =Tags(s)
	<p>MPYQYPALTPEQKKELSDIAHRIVAPGKGI<del>LA</del>DESTGSI<del>AKRL</del>QSIGTENTEENRRFYRQLLLTADDRV NPCIGGVILFHETLYQKADDGRPFQVIKSKGGVVGIKVDKGVPLAGTNGETTTQGLDGLSERCAQYKK DGADFAKWRCVLKIGEHTPSALAIMENANVLARYASICQQNGIVIVEPEILPDGDHDLKRCQYVTEKVL AAVYKALSDHHIYLEGTLTKPNMVT<del>PGHACT</del>QKFSHEEIAMATVTALRRTVPPAVTGITFLSGGGQSEEEA SINLNAINKCPLLKPWALTF<del>SYGRAL</del>QASALKAWGGKKENL<del>KAAQ</del>EEYVKRALANSLACQ<del>GKYTP</del>SGQAG AAASESLFVSNHAY</p> <p><b>TRTRPLEQKLISEEDLAANDILDYKDDDDKV</b></p>
Tag:	C-Myc/DDK
Predicted MW:	39.2 kDa
Concentration:	>0.05 µg/µL as determined by microplate BCA method
Purity:	> 80% as determined by SDS-PAGE and Coomassie blue staining
Buffer:	25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol
Preparation:	Recombinant protein was captured through anti-DDK affinity column followed by conventional chromatography steps.
Note:	For testing in cell culture applications, please filter before use. Note that you may experience some loss of protein during the filtration process.
Storage:	Store at -80°C.
Stability:	Stable for 12 months from the date of receipt of the product under proper storage and handling conditions. Avoid repeated freeze-thaw cycles.
RefSeq:	<u><a href="#">NP_000025</a></u>



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Locus ID: 226

UniProt ID: [P04075](#), [V9HWN7](#)

RefSeq Size: 2408

Cytogenetics: 16p11.2

RefSeq ORF: 1092

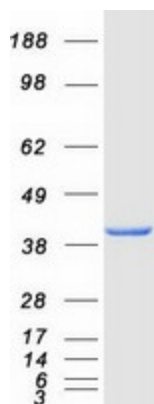
Synonyms: ALDA; GSD12; HEL-S-87p

**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]

**Protein Families:** Druggable Genome

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

### Product images:



Coomassie blue staining of purified ALDOA protein (Cat# [TP302576]). The protein was produced from HEK293T cells transfected with ALDOA cDNA clone (Cat# [RC202576]) using MegaTran 2.0 (Cat# [TT210002]).