

Product datasheet for PH303908

Aldolase (ALDOA) (NM_184043) Human Mass Spec Standard

Product data:

| | |
|---------------------------------------|---|
| Product Type: | Mass Spec Standards |
| Description: | ALDOA MS Standard C13 and N15-labeled recombinant protein (NP_908932) |
| Species: | Human |
| Expression Host: | HEK293 |
| Expression cDNA Clone or AA Sequence: | RC203908 |
| Predicted MW: | 39.4 kDa |
| Protein Sequence: | >RC203908 protein sequence Red=Cloning site Green=Tags(s) |

MPYQYPALTPEQKKELSDIAHRIVAPGKGI LADESTGSI AKRLQSIGTENTEENRRFYRQLLLTADDRV
NPCIGGVILFHETLYQKADDGRPFQVIKSKGGVVGIVKVDKGVVPLAGTNGETTTQGLDGLSERCAQYKK
DGADFAKWRVCVLKIGEHTPSALAIMENANVLARYASICQNGIVPIVEPEILPDGDHDLKRCQYVTEKVL
AAVYKALSDHHIYLEGTLKPNMVTGPHACTQKFSHEEIAMATVTALRRTPPAVTGITFLSGGQSEEEA
SINLNAINKCPLLKPWALTF SYGRALQASALKAWGGKENL KAAQEEYVKRALANSLACQKYPSPGQAG
AAASESLFVSNHAY

TRTRPLEQKLI SEEDLAANDILDYKDDDDKV

| | |
|------------------|--|
| Tag: | C-Myc/DDK |
| Purity: | > 80% as determined by SDS-PAGE and Coomassie blue staining |
| Concentration: | >0.05 µg/µL as determined by microplate BCA method |
| Labeling Method: | Labeled with [U- ¹³ C ₆ , ¹⁵ N ₄]-L-Arginine and [U- ¹³ C ₆ , ¹⁵ N ₂]-L-Lysine |
| Buffer: | 25 mM Tris-HCl, 100 mM glycine, pH 7.3 |
| Storage: | Store at -80°C. Avoid repeated freeze-thaw cycles. |
| Stability: | Stable for 3 months from receipt of products under proper storage and handling conditions. |
| RefSeq: | NP_908932 |
| RefSeq Size: | 1569 |
| RefSeq ORF: | 1092 |
| Synonyms: | ALDA; GSD12; HEL-S-87p |
| Locus ID: | 226 |



[View online »](#)

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

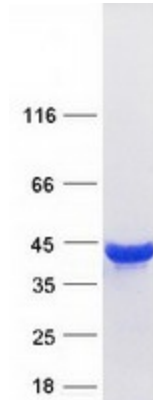
Cytogenetics: 16p11.2

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# [TP303908]). The protein was produced from HEK293T cells transfected with ALDOA cDNA clone (Cat# [RC203908]) using MegaTran 2.0 (Cat# [TT210002]).