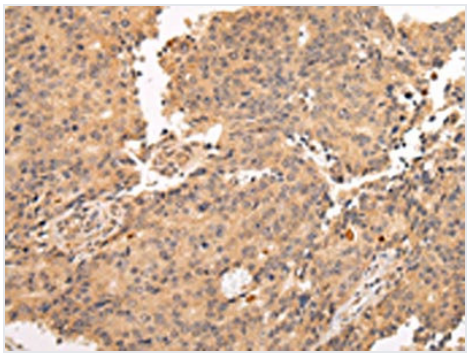
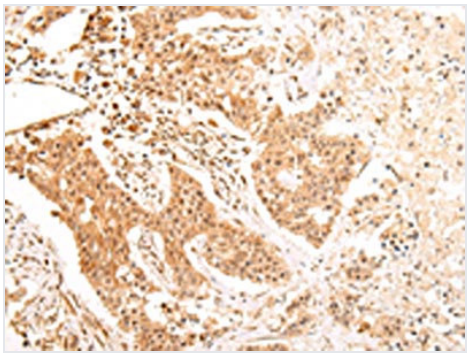
ALDH3A2 Antibody – Rabbit Polyclonal

PRODUCT OVERVIEW

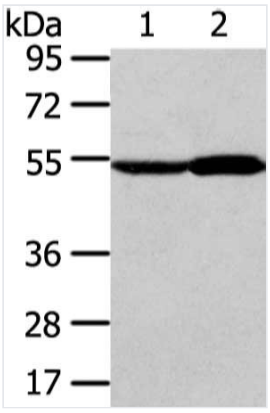
Product Name	ALDH3A2, Polyclonal Antibody
Full Product Name	ALDH3A2 Antibody
Product Gene Name	anti-ALDH3A2 antibody
Product Synonym Names	aldehyde dehydrogenase 3 family; member A2
Clonality	Polyclonal
Isotype	IgG
Host	Rabbit
Species Reactivity	Human
Form/Format	Liquid; In Phosphate Buffered Saline, pH 7.4, 0.05% NaN3, and 40% Glycerol
Purity/Purification	Antigen Affinity Purification
Preparation and Storage	Upon receipt, store at -20 degree C or -80 degree C. Avoid repeated freeze.
Molecular Weight	485
OMIM	phenotype 609523
3D Structure	ModBase 3D Structure for P51648
Immunogen	Fusion protein of human ALDH3A2
Immunogen Species	Human
Conjugate	Non-conjugated
Product Categories/Family for anti-ALDH3A2 antibody	Cardiovascular
ISO Certification	Manufactured in an ISO 9001:2015 Certified Laboratory.
Applications Tested/Suitable for anti-ALDH3A2 antibody	ELISA (EIA), Western Blot (WB), Immunohistochemistry (IHC)
Application Notes for anti-ALDH3A2 antibody	ELISA: 1:2000-1:5000 WB: 1:500-1:2000 IHC: 1:25-1:100
Related Product Information for anti-ALDH3A2 antibody	MBS7126653 is an antibody targeting ALDH3A2. Also known as ALDH3A2 Antibody. Reactive in Human. Host: Rabbit.
Other Notes	Small volumes of anti-ALDH3A2 antibody vial(s) may occasionally become entrapped in the seal of the product vial during shipment and storage. If necessary, briefly centrifuge the vial on a tabletop centrifuge to dislodge any liquid in the container`s cap. Certain products may require to ship with dry ice and additional dry ice fee may apply.



Immunohistochemistry (IHC)-Paraffin of anti-ALDH3A2 antibody
The image on the left is immunohistochemistry of paraffin-embedded Human prostate cancer tissue using MBS7126653(ALDH3A2 Antibody) at dilution 1/20, on the right is treated with fusion protein. (Original magnification: ×200)



Immunohistochemistry (IHC)-Paraffin of anti-ALDH3A2 antibody
The image on the left is immunohistochemistry of paraffin-embedded Human colorectal cancer tissue using MBS7126653(ALDH3A2 Antibody) at dilution 1/20, on the right is treated with fusion protein. (Original magnification: ×200)



SDS-Page of anti-ALDH3A2 antibody
Gel: 6%SDS-PAGE, Lysate: 40 ug, Lane 1-2: Human fetal liver and normal liver tissue, Primary antibody: MBS7126653(ALDH3A2 Antibody) at dilution 1/250 dilution, Secondary antibody: Goat anti rabbit IgG at 1/8000 dilution, Exposure time: 10 seconds

ADDITIONAL DETAILS

NCBI/Uniprot data below describe general gene information for ALDH3A2. It may not necessarily be applicable to this product.

NCBI GI #	1706379
NCBI GeneID	224
NCBI Accession #	P51648.1
UniProt Primary Accession #	P51648
UniProt Secondary Accession #	Q6I9T3; Q93011; Q96J37
UniProt Related Accession #	P51648
NCBI Official Full Name	Fatty aldehyde dehydrogenase
NCBI Official Synonym Full Names	aldehyde dehydrogenase 3 family, member A2
NCBI Official Symbol	ALDH3A2
NCBI Official Synonym Symbols	SLS; FALDH; ALDH10
UniProt Protein Name	Fatty aldehyde dehydrogenase
UniProt Synonym Protein Names	Aldehyde dehydrogenase 10; Aldehyde dehydrogenase family 3 member A2; Microsomal aldehyde dehydrogenase
Protein Family	Fatty aldehyde dehydrogenase
UniProt Gene Name	ALDH3A2

UniProt Synonym Gene Names	ALDH10; FALDH
UniProt Entry Name	AL3A2_HUMAN
NCBI Protein Information	fatty aldehyde dehydrogenase; aldehyde dehydrogenase 10; microsomal aldehyde dehydrogenase; aldehyde dehydrogenase family 3 member A2
NCBI Summary for ALDH3A2	Aldehyde dehydrogenase isozymes are thought to play a major role in the detoxification of aldehydes generated by alcohol metabolism and lipid peroxidation. This gene product catalyzes the oxidation of long-chain aliphatic aldehydes to fatty acid. Mutations in the gene cause Sjogren-Larsson syndrome. Alternatively spliced transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul 2008]
UniProt Comments for ALDH3A2	ALDH3A2: Catalyzes the oxidation of long-chain aliphatic aldehydes to fatty acids. Active on a variety of saturated and unsaturated aliphatic aldehydes between 6 and 24 carbons in length. Responsible for conversion of the sphingosine 1-phosphate (S1P) degradation product hexadecenal to hexadecenoic acid. Defects in ALDH3A2 are the cause of Sjoegren-Larsson syndrome (SLS). SLS is an autosomal recessive neurocutaneous disorder characterized by a combination of severe mental retardation, spastic di- or tetraplegia and congenital ichthyosis (increased keratinization). Ichthyosis is usually evident at birth, neurologic symptoms appear in the first or second year of life. Most patients have an IQ of less than 60. Additional clinical features include glistening white spots on the retina, seizures, short stature and speech defects. Belongs to the aldehyde dehydrogenase family. 2 isoforms of the human protein are produced by alternative splicing. Protein type: Carbohydrate Metabolism - propanoate; Amino Acid Metabolism - histidine; Membrane protein, integral; Oxidoreductase; EC 1.2.1.3; Amino Acid Metabolism - lysine degradation; Carbohydrate Metabolism - ascorbate and aldarate; Carbohydrate Metabolism - pyruvate; Carbohydrate Metabolism - butanoate; Other Amino Acids Metabolism - beta-alanine; Lipid Metabolism - fatty acid; Amino Acid Metabolism - arginine and proline; Secondary Metabolites Metabolism - limonene and pinene degradation; Lipid Metabolism - glycerolipid; Carbohydrate Metabolism - glycolysis and gluconeogenesis; Amino Acid Metabolism - tryptophan; Amino Acid Metabolism - valine, leucine and isoleucine degradation; Mitochondrial Chromosomal Location of Human Ortholog: 17p11.2 Cellular Component: endoplasmic reticulum membrane; intracellular membrane-bound organelle; mitochondrial inner membrane; integral to membrane; peroxisome Molecular Function: long-chain-alcohol oxidase activity; aldehyde dehydrogenase (NAD) activity; aldehyde dehydrogenase [NAD(P)+] activity; 3-chloroallyl aldehyde dehydrogenase activity; long-chain-aldehyde dehydrogenase activity Biological Process: phytol metabolic process; epidermis development; central nervous system development; aldehyde metabolic process; sesquiterpenoid metabolic process; peripheral nervous system development Disease: Sjogren-larsson Syndrome
Research Articles on ALDH3A2	**1.** variation in the neurologic phenotype of Sjogren-Larsson syndrome is not strictly determined by the ALDH3A2 mutation or a biochemical defect, but by unidentified epigenetic/environmental factors, gene modifiers, or other mechanisms.
Precautions	All of MyBioSource's Products are for scientific laboratory research purposes and are not for diagnostic, therapeutics, prophylactic or in vivo use. Through your purchase, you expressly represent and warrant to MyBioSource that you will properly test and use any Products purchased from MyBioSource in accordance with industry standards. MyBioSource and its authorized distributors reserve the right to refuse to process any order where we reasonably believe that the intended use will fall outside of our acceptable guidelines.
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Product Note	Select online data sheet information is drawn from bioinformatics databases, occasionally resulting in ambiguous or non-relevant product information. It is the responsibility of the customer to review, verify, and evaluate the information to make sure it matches their requirements before purchasing the kit. Our ELISA Kit assays are dynamic research tools and sometimes they may be updated and improved. If the format of this assay is important to you then please request the current manual or contact our technical support team with a presales inquiry before placing an order. We will confirm the current details of the assay. We cannot guarantee the sample manual posted online is the most current manual, it is intended to serve as an example only. Please refer to the instructions For Use provided with the assay kit for precise details.

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