

Datasheet for ABIN1393379

anti-ALDH3A2 antibody (AA 101-200) (AbBy Fluor® 555)



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3.01.11011		
Quantity:	100 μL	
Target:	ALDH3A2	
Binding Specificity:	AA 101-200	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This ALDH3A2 antibody is conjugated to AbBy Fluor® 555	
Application:	Western Blotting (WB), Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p))	
Product Details		
Immunogen:	KLH conjugated synthetic peptide derived from human ALDH3A2	
Isotype:	IgG	
Predicted Reactivity:	Human,Mouse,Rat	
Purification:	Purified by Protein A.	
Target Details		
Target:	ALDH3A2	
Alternative Name:	ALDH3A2 (ALDH3A2 Products)	
Background:	Synonyms: Ahd 3, Ahd 3r, Ahd3, Ahd3 r, AL3A2_HUMAN, Aldehyde dehydrogenase 10, Aldehyde	

dehydrogenase 3, Aldehyde dehydrogenase 3 family, member A2, Aldehyde dehydrogenase family 3 member A2, Aldehyde dehydrogenase family 3, subfamily A2, Aldehyde dehydrogenase, family 3, subfamily A, member 2, ALDH10, Aldh3, ALDH3A2, Aldh4, Aldh4 r, Aldh4r, FALDH, Fatty aldehyde dehydrogenase, FLJ20851, Microsomal aldehyde dehydrogenase, msALDH, SLS.

Background: Aldehyde dehydrogenases (ALDHs) mediate the NADP+-dependent oxidation of aldehydes into acids and play an important role in the detoxification of alcohol-derived acetaldehyde, as well as in lipid peroxidation and in the metabolism of corticosteroids, biogenic amines and neurotransmitters. ALDH3A2 (aldehyde dehydrogenase 3 family, member A2), also known as SLS, FALDH or ALDH10, is a 485 amino acid single-pass membrane protein that localizes to the cytoplasmic side of the endoplasmic reticulum and belongs to the aldehyde dehydrogenase family. Expressed in a variety of tissues, including liver, heart, lung, brain, kidney and placenta, ALDH3A2 catalyzes the NAD+-dependent oxidation of long-chain aliphatic aldehydes to fatty acids, a process that is necessary for detoxification and lipid metabolism. Defects in the gene encoding ALDH3A2 are the cause of Sjoegren-Larsson syndrome (SLS), an autosomal recessive neurocutaneous disorder characterized by severe mental retardation, seizures and speech defects. Multiple isoforms of ALDH3A2 exist due to alternative splicing events.

Application Details

Application Notes:	IF(IHC-P) 1:50-200
	IF(IHC-F) 1:50-200
	IF(ICC) 1:50-200
Restrictions:	For Research Use only

Handling

папишту	
Format:	Liquid
Concentration:	1 μg/μL
Buffer:	Aqueous buffered solution containing 0.01M TBS (pH 7.4) with 1 % BSA, 0.03 % Proclin300 and 50 % Glycerol.
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be handled by trained staff only.

Handling

Storage:	-20 °C
Storage Comment:	Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles.
Expiry Date:	12 months