



Aldehyde dehydrogenase 10 mouse mAb

Catalog No	YP-Ab-02360
Isotype	IgG
Reactivity	Human
Applications	WB
Gene Name	aldh3a2
Immunogen	Purified recombinant human Aldehyde dehydrogenase 10 protein fragments expressed in E.coli.
Specificity	This antibody detects endogenous levels of Aldehyde dehydrogenase 10 and does not cross-react with related proteins.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse
Purification	The antibody was affinity-purified from mouse ascites by affinity-chromatography using epitope-specific immunogen.
Dilution	wb 1:1000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	Aldehyde dehydrogenase 10;ALDH10;EC:1.2.1.3;FALDH;SLS;aldh3a2; Aldehyde dehydrogenase;Aldehyde dehydrogenase 3 family member A2; DKFZp686E23276;Fatty aldehyde dehydrogenase;FLJ20851;Microsomal aldehyde dehydrogenase;OTTHUMP00000065799;OTTHUMP00000065800; OTTHUMP00000065801;SLS.
Observed Band	55kD
Calculated Molecular Weight	55kD
Cell Pathway	Microsome membrane ; Single-pass membrane protein . Endoplasmic reticulum membrane ; Single-pass membrane protein ; Cytoplasmic side .
Tissue Specificity	Detected in liver (at protein level).
Function	catalytic activity:An aldehyde + NAD(+) + H(2)O = an acid + NADH., disease:Defects in ALDH3A2 are the cause of Sjogren-Larsson syndrome (SLS) [MIM:270200]. 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.,function: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.,similarity:Belongs to the aldehyde dehydrogenase family.,

Background

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],

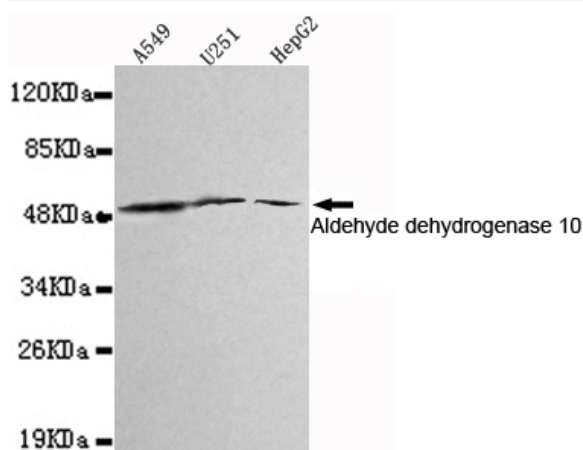
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western blot detection of Aldehyde dehydrogenase 10 in A549,U251 and HepG2 cell lysates using Aldehyde dehydrogenase 10 mouse mAb (1:1000 diluted).Predicted band size:55KDa.Observed band size:55KDa.