abcam

Product datasheet

Anti-PDHA1 antibody ab97352

1 References 3 Images

Overview

Product name Anti-PDHA1 antibody

Description Rabbit polyclonal to PDHA1

Host species Rabbit

Suitable for: WB, ICC/IF **Tested applications** Species reactivity Reacts with: Human

Predicted to work with: Mouse, Cow, Zebrafish

Immunogen Recombinant fragment within Human PDHA1 aa 39-295. The exact sequence is proprietary.

(NP_000275).

Positive control WB: 293T, A431, H1299, HeLa S3, HepG2, MOLT4 and Raji ICC/IF: HeLa cells

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Shipped at 4°C. Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle. Storage instructions

pH: 7.00 Storage buffer

Preservative: 0.025% Proclin 300

Constituents: 79% PBS, 20% Glycerol (glycerin, glycerine)

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype lqG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab97352 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		1/500 - 1/3000. Predicted molecular weight: 43 kDa.
ICC/IF		1/100 - 1/200.

Target

Function

The pyruvate dehydrogenase complex catalyzes the overall conversion of pyruvate to acetyl-CoA and CO(2). It contains multiple copies of three enzymatic components: pyruvate dehydrogenase (E1), dihydrolipoamide acetyltransferase (E2) and lipoamide dehydrogenase (E3).

Tissue specificity

Involvement in disease

Ubiquitous.

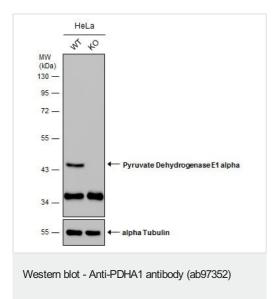
Defects in PDHA1 are a cause of pyruvate decarboxylase E1 component deficiency (PDHE1 deficiency) [MIM:312170]. PDHE1 deficiency is the most common enzyme defect in patients with primary lactic acidosis. It is associated with variable clinical phenotypes ranging from neonatal death to prolonged survival complicated by developmental delay, seizures, ataxia, apnea, and in some cases to an X-linked form of Leigh syndrome (X-LS).

Defects in PDHA1 are the cause of X-linked Leigh syndrome (X-LS) [MIM:308930]. X-LS is an early-onset progressive neurodegenerative disorder with a characteristic neuropathology consisting of focal, bilateral lesions in one or more areas of the central nervous system, including the brainstem, thalamus, basal ganglia, cerebellum, and spinal cord. The lesions are areas of demyelination, gliosis, necrosis, spongiosis, or capillary proliferation. Clinical symptoms depend on which areas of the central nervous system are involved. The most common underlying cause is a defect in oxidative phosphorylation. LS may be a feature of a deficiency of any of the mitochondrial respiratory chain complexes.

Cellular localization

Mitochondrion matrix.

Images



All lanes: Anti-PDHA1 antibody (ab97352) at 1/1000 dilution

Lane 1: WT HeLa cell extract

Lane 2: Pyruvate Dehydrogenase E1-alpha subunit Knockout

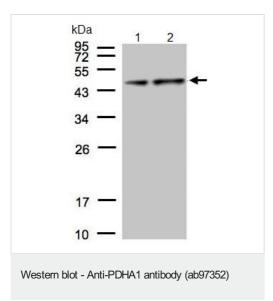
HeLa cell extracts

Lysates/proteins at 30 µg per lane.

Secondary

All lanes: HRP-conjugated anti-rabbit lgG

Predicted band size: 43 kDa



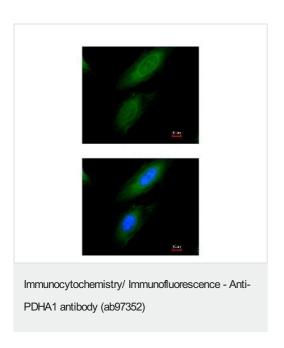
All lanes: Anti-PDHA1 antibody (ab97352) at 1/1000 dilution

Lane 1 : HepG2 whole cell lysate
Lane 2 : MOLT4 whole cell lysate

Lysates/proteins at 30 µg per lane.

Predicted band size: 43 kDa

12% SDS PAGE



ab97352, at a 1/200 dilution, staining PDHA1 in paraformaldehyde fixed HeLa cells by Immunofluorescence. Lower image was merged with DNA probe.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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