abcam

Product datasheet

Anti-ASPA antibody ab97454

1 References 2 Images

Overview

Product name Anti-ASPA antibody

Description Rabbit polyclonal to ASPA

Host species Rabbit

Tested applications Suitable for: WB, IHC-P

Species reactivity Reacts with: Human

Predicted to work with: Mouse, Cow, Pig

Immunogen Recombinant fragment, corresponding to a region within amino acids 39-270 of Human ASPA

(AAH29128).

General notesThe 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

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

Storage buffer pH: 7.00

Preservative: 0.025% Proclin 300

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

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab97454 in the following tested applications.

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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: 36 kDa.
IHC-P		1/100 - 1/500.

Target

Function

Catalyzes the deacetylation of N-acetylaspartic acid (NAA) to produce acetate and L-aspartate.

NAA occurs in high concentration in brain and its hydrolysis NAA plays a significant part in the maintenance of intact white matter. In other tissues it act as a scavenger of NAA from body fluids.

Tissue specificity Brain white matter, skeletal muscle, kidney, adrenal glands, lung and liver.

Involvement in disease Defects in ASPA are the cause of Canavan disease (CAND) [MIM:271900]; also known as spongy degeneration of the brain. CAND is a rare neurodegenerative condition of infancy or childhood characterized by white matter vacualization and demovlination that gives rise to a

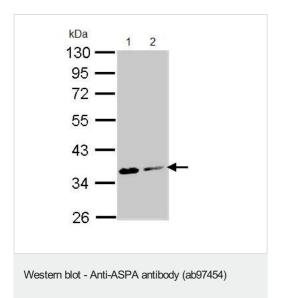
childhood characterized by white matter vacuolization and demeylination that gives rise to a spongy appearance. The clinical features are onset in early infancy, atonia of neck muscles, hypotonia, hyperextension of legs and flexion of arms, blindness, severe mental defect,

megalocephaly, and death by 18 months on the average.

Sequence similaritiesBelongs to the AspA/AstE family. Aspartoacylase subfamily.

Cellular localization Cytoplasm. Nucleus.

Images



All lanes: Anti-ASPA antibody (ab97454) at 1/1000 dilution

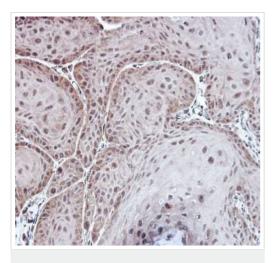
Lane 1 : MOLT4 whole cell lysate

Lane 2: Raji whole cell lysate

Lysates/proteins at 30 µg per lane.

Predicted band size: 36 kDa

10% SDS PAGE



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ASPA antibody (ab97454)

ab97454 at 1/100 dilution staining ASPA in paraffin-embedded Cal27 xenograft by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections).

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

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