abcam

Product datasheet

Anti-NUBPL antibody ab229670

2 Images

Overview

Product name Anti-NUBPL antibody

Description Rabbit polyclonal to NUBPL

Host species Rabbit

Tested applications Suitable for: WB

Species reactivity Reacts with: Mouse, Human

Predicted to work with: Rat, Cow

Immunogen Recombinant fragment within Human NUBPL (internal sequence). The exact sequence is

proprietary.

Database link: Q8TB37

Positive control Mouse liver tissue extract; HepG2 whole cell extract.

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. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°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

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Applications

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab229670 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: 34 kDa.

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Function	Required for the assembly of the mitochondrial membrane respiratory chain	NADH

dehydrogenase (Complex I). May deliver of one or more Fe-S clusters to complex I subunits.

Tissue specificity

Highest expression in liver and kidney. expressed at significant levels in small intestine and brain

(at protein level).

Involvement in disease

Defects in NUBPL are a cause of mitochondrial complex I deficiency (MT-C1D) [MIM:252010]. A disorder of the mitochondrial respiratory chain that causes a wide range of clinical manifestations from lethal neonatal disease to adult-onset neurodegenerative disorders. Phenotypes include macrocephaly with progressive leukodystrophy, non-specific encephalopathy, cardiomyopathy, myopathy, liver disease, Leigh syndrome, Leber hereditary optic neuropathy, and some forms of

Parkinson disease.

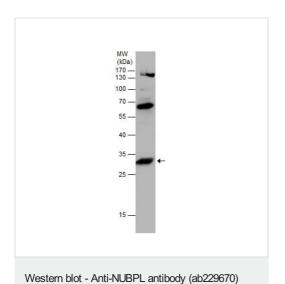
Sequence similarities

Belongs to the Mrp/NBP35 ATP-binding proteins family.

Cellular localization

Mitochondrion.

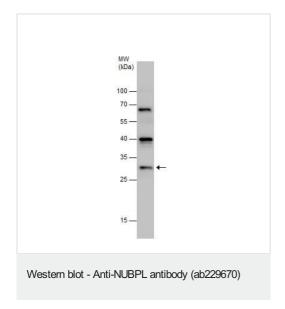
Images



Anti-NUBPL antibody (ab229670) at 1/500 dilution + HepG2 (human liver hepatocellular carcinoma cell line) whole cell extract at $30~\mu g$

Predicted band size: 34 kDa

12% SDS-PAGE gel.



Anti-NUBPL antibody (ab229670) at 1/1000 dilution + Mouse liver tissue extract at 50 μg

Predicted band size: 34 kDa

12% SDS-PAGE gel.

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

Our Abpromise to you: Quality guaranteed and expert technical support

- Replacement or refund for products not performing as stated on the datasheet
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- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- · We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit **https://www.abcam.com/abpromise** or contact our technical team.

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