# abcam

### Product datasheet

## Anti-Glycine decarboxylase antibody ab97625

**5 References** 2 Images

Overview

Product name Anti-Glycine decarboxylase antibody

**Description** Rabbit polyclonal to Glycine decarboxylase

Host species Rabbit

Tested applications Suitable for: WB, ICC/IF

Species reactivity Reacts with: Human

Predicted to work with: Mouse

**Immunogen** Recombinant fragment corresponding to Human Glycine decarboxylase aa 492-693.

Database link: NP\_000161

Positive control HepG2 cells and lysate

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

**Storage instructions** Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw

cycles.

Storage buffer pH: 7.00

Preservative: 0.025% Proclin 300

Constituents: 78% PBS, 1% BSA, 20% Glycerol (glycerin, glycerine)

Purity Immunogen affinity purified

**Clonality** Polyclonal

**Isotype** IgG

**Applications** 

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#### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab97625 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: 113 kDa.
ICC/IF		1/100 - 1/200.

#### **Target**

Function

The glycine cleavage system catalyzes the degradation of glycine. The P protein binds the alphaamino group of glycine through its pyridoxal phosphate cofactor; CO(2) is released and the
remaining methylamine moiety is then transferred to the lipoamide cofactor of the H protein.

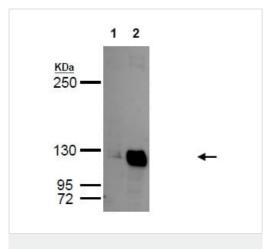
Involvement in disease

Defects in GLDC are a cause of non-ketotic hyperglycinemia (NKH) [MIM:605899]; also known as
glycine encephalopathy (GCE). NKH is an autosomal recessive disease characterized by
accumulation of a large amount of glycine in body fluid and by severe neurological symptoms.

**Sequence similarities** Belongs to the gcvP family.

Cellular localization Mitochondrion.

#### **Images**



Western blot - Anti-Glycine decarboxylase antibody (ab97625)

**All lanes :** Anti-Glycine decarboxylase antibody (ab97625) at 1/500 dilution

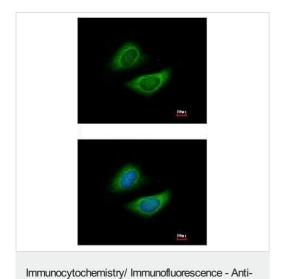
Lane 1 : HeLa whole cell lysate

Lane 2 : HepG2 whole cell lysate

Lysates/proteins at 30 µg per lane.

Predicted band size: 113 kDa

5% SDS-PAGE



Glycine decarboxylase antibody (ab97625)

ab97625, at a 1/200 dilution, staining Glycine decarboxylase in methanol fixed HeLa cells by Immunofluorescence analysis. Lower image was co-stained with Hoechst 33342.

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

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- Replacement or refund for products not performing as stated on the datasheet
- Valid for 12 months from date of delivery
- 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 <a href="https://www.abcam.com/abpromise">https://www.abcam.com/abpromise</a> or contact our technical team.

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