# abcam

# Product datasheet

# Anti-BTD antibody ab 166934

# 1 Image

#### Overview

Product name Anti-BTD antibody

**Description** Goat polyclonal to BTD

Host species Goat

Tested applications Suitable for: WB

Species reactivity Reacts with: Human

Predicted to work with: Mouse, Rat

Immunogen Synthetic peptide, corresponding to internal sequence amino acids 372-385 of Human BTD

(NP\_000051.1).

Run BLAST with EXPASY M Run BLAST with S NCBI

Positive control Human Liver lysates.

**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. Avoid repeated freeze / thaw cycles.

Storage buffer pH: 7.30

Preservative: 0.02% Sodium azide

Constituents: 99% Tris buffered saline, 0.5% BSA

Purity Immunogen affinity purified

Purification notes ab166934 is purified from goat serum by ammonium sulphate precipitation followed by antigen

affinity chromatography using the immunizing peptide.

**Clonality** Polyclonal

**Isotype** IgG

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#### **Applications**

#### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab166934 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		Use a concentration of 0.5 - 1.5 μg/ml. Predicted molecular weight: 61 kDa.  1 hour primary incubation is recommended for this product.

#### **Target**

<b>Function</b> Catalytic release of biotin from biocytin, the product of biotin-dependent carboxylas
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degradation.

# **Involvement in disease**Defects in BTD are the cause of biotinidase deficiency (BTD deficiency) [MIM:253260]; also

called late-onset multiple carboxylase deficiency. BTD deficiency is a juvenile form of multiple carboxylase deficiency, an autosomal recessive disorder of biotin metabolism, characterized by ketoacidosis, hyperammonemia, excretion of abnormal organic acid metabolites, and dermatitis. BTD deficiency is characterized by seizures, hypotonia, skin rash, alopecia, ataxia, hearing loss, and optic atrophy. If untreated, symptoms usually become progressively worse, and coma and

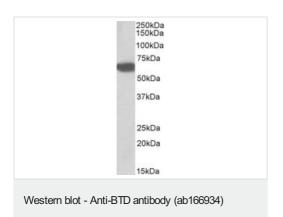
death may occur.

Sequence similarities Belongs to the CN hydrolase family. BTD/VNN subfamily.

Contains 1 CN hydrolase domain.

**Cellular localization** Secreted > extracellular space.

#### **Images**



Anti-BTD antibody (ab166934) at 0.5  $\mu g/ml$  + Human liver lysate (in

RIPA buffer) at 35 µg

Developed using the ECL technique.

Predicted band size: 61 kDa

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 <a href="https://www.abcam.com/abpromise">https://www.abcam.com/abpromise</a> or contact our technical team.

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