# abcam

### Product datasheet

## Anti-BACE1 antibody ab10716

6 References 4 Images

Overview

Product name Anti-BACE1 antibody

**Description** Rabbit polyclonal to BACE1

Host species Rabbit

**Specificity** Detects transfected beta-secretase 1.

Tested applications Suitable for: WB, ICC/IF

Species reactivity Reacts with: Mouse, Human

Predicted to work with: Guinea pig, Cow

Immunogen Synthetic peptide corresponding to Human BACE1 aa 485-501 (C terminal).

Sequence:

CLRQQHDDFADDISLLK

(Peptide available as ab7883)

Run BLAST with
Run BLAST with

Positive control WB: U87-MG, HeLa and Mouse brain cell lysate. ICC/IF: SH-SY5Y cells, Neuro-2a cells, 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

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

80°C. Avoid freeze / thaw cycle.

**Storage buffer** Preservative: 0.05% Sodium azide

Constituents: 99% PBS, 0.1% BSA

Purity Immunogen affinity purified

1

**Clonality** Polyclonal

**Isotype** IgG

#### **Applications**

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab10716 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		1/1000. Predicted molecular weight: 56 kDa.
ICC/IF		1/20 - 1/200.

Target	t
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**Function** Responsible for the proteolytic processing of the amyloid precursor protein (APP). Cleaves at the

N-terminus of the A-beta peptide sequence, between residues 671 and 672 of APP, leads to the

generation and extracellular release of beta-cleaved soluble APP, and a corresponding cell-

associated C-terminal fragment which is later released by gamma-secretase.

**Tissue specificity** Expressed at high levels in the brain and pancreas. In the brain, expression is highest in the

substantia nigra, locus coruleus and medulla oblongata.

**Sequence similarities** Belongs to the peptidase A1 family.

**Domain** The transmembrane domain is necessary for its activity. It determines its late Golgi localization

and access to its substrate, APP.

Post-translational

modifications

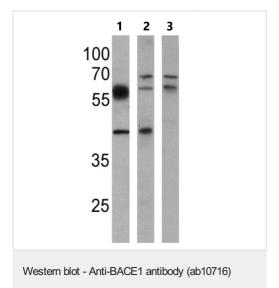
Glycosylated.

**Cellular localization** Membrane. Golgi apparatus > trans-Golgi network. Endoplasmic reticulum. Endosome. Cell

surface. Predominantly localized to the later Golgi/trans-Golgi network (TGN) and minimally detectable in the early Golgi compartments. A small portion is also found in the endoplasmic

reticulum, endosomes and on the cell surface.

#### **Images**



All lanes: Anti-BACE1 antibody (ab10716) at 1/1000 dilution

Lane 1 : U87-MG cell lysate

Lane 2: HeLa cell lysate

Lane 3: Mouse brain cell lysate

Lysates/proteins at 25 µg per lane.

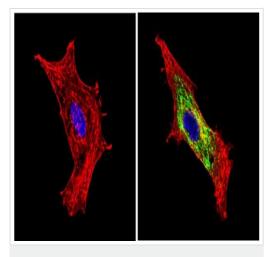
#### Secondary

**All lanes:** HRP-conjugated secondary antibody

Predicted band size: 56 kDa

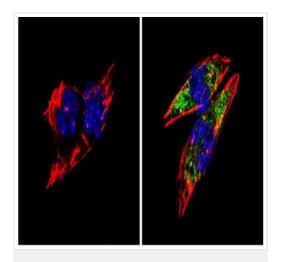
Observed band size: 42,56,70 kDa

Western blot analysis of BACE1 was performed by loading Samples onto an SDS polyacrylamide gel. Proteins were transferred to a PVDF membrane and blocked at 4°C overnight. The membrane was probed with primary antibody overnight at 4°C, washed in TBST, and probed with secondary antibody for 1 hour at room temperature in the dark. Chemiluminescent detection was performed using Pierce ECL Plus Western Blotting Substrate.



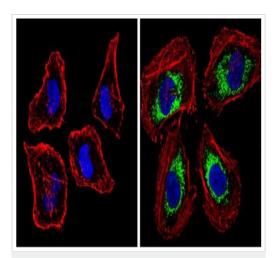
Immunocytochemistry/ Immunofluorescence - Anti-BACE1 antibody (ab10716)

Immunocytochemistry/Immunofluorescent analysis of BACE1 (green) showing staining in the Golgi apparatus of SH-SY5Y cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with a10716 in 3% BSA-PBS at a dilution of 1/100 and incubated overnight at 4°C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight-conjugated secondary antibody in PBS at room temperature in the dark. F-actin (red) was stained with a fluorescent red phalloidin and nuclei (blue) were stained with Hoechst or DAPI. Images were taken at a magnification of 60x.



Immunocytochemistry/ Immunofluorescence - Anti-BACE1 antibody (ab10716)

Immunocytochemistry/Immunofluorescent analysis of BACE1 (green) showing staining in the Golgi apparatus of Neuro-2a cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with ab10716 in 3% BSA-PBS at a dilution of 1/100 and incubated overnight at 4°C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight-conjugated secondary antibody in PBS at room temperature in the dark. F-actin (red) was stained with a fluorescent red phalloidin and nuclei (blue) were stained with Hoechst or DAPI. Images were taken at a magnification of 60x.



Immunocytochemistry/ Immunofluorescence - Anti-BACE1 antibody (ab10716)

Immunocytochemistry/Immunofluorescent analysis of BACE1 (green) showing staining in the Golgi apparatus of HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with ab10716 in 3% BSA-PBS at a dilution of 1/100 and incubated overnight at 4°C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight-conjugated secondary antibody in PBS at room temperature in the dark. F-actin (red) was stained with a fluorescent red phalloidin and nuclei (blue) were stained with Hoechst or DAPI. Images were taken at a magnification of 60x.

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