

BOSTER BIOLOGICAL TECHNOLOGY

Building C21, 3rd and 4th floors, Optics Valley Biomedical Accelerator, Wuhan East Lake High-tech Development Zone

Web: www.boster.com Phone: 027-67845390 Email: boster@boster.com

Basic Information		
Product Name	Anti-Beta Amyloid protein 42/APP Antibody	
Gene Name	APP	
Source	Rabbit	
Clonality	Polyclonal	
lsotype	IgG	
Species Reactivity	human, mouse, rat	
Tested Application	WB, IHC, ICC/IF	
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.	
Immunogen	A synthetic peptide corresponding to a sequence at the C-terminus of human APP, different from the related mouse and rat sequences by three amino acids.	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	87-120 kDa	
Dilution Ratios	Western blot (WB): Immunohistochemistry (IHC): Immunocytochemistry/Immunofluorescence (ICC/IF): (Boiling the paraffin sections in 10mM citrate buffer,pH6.0, for 20 mins is required for the staining of formalin/paraffin dilutions must be determined by end user.	

### **Storage**

12 months from date of receipt, -20°C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

## **Background Information**

beta Amyloid, also called Abeta or Abeta, denotes peptides of 36–43 amino acidsthat are crucially involved in Alzheimer's disease as the main component of the amyloid plaques found in the brains of Alzheimer patients. It is mapped to 19q13.12. Several potential activities have been discovered for beta Amyloid, including activation of kinase enzymes, functioning as atranscription factor, and anti-microbial activity (potentially associated with beta Amyloid's pro-inflammatoryactivity). Moreover, monomeric beta Amyloid is indicated to protect neurons by quenching metal-inducible oxygen radical generation and thereby inhibiting neurotoxicity.

### Reference

#### Product datasheet Anti-Beta Amyloid protein 42/APP Antibody Catalog Number: PB9091



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Anti-Beta Amyloid protein 42/APP Antibody被引用在15文献中。

# **Selected Validation Data**

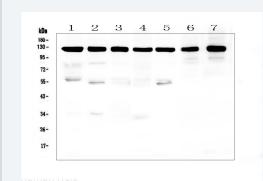


Figure 1. Western blot analysis of Beta Amyloid protein 42/APP using anti-Beta Amyloid protein 42/APP antibody (PB9091). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human Hela whole cell lysates,

Lane 2: human U-87MG whole cell lysates,

Lane 3: human T-47D whole cell lysates,

Lane 4: human A549 whole cell lysates,

Lane 5: human U2OS whole cell lysates,

Lane 6: rat brain tissue lysates,

Lane 7: mouse brain tissue lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-Beta Amyloid protein 42/APP antigen affinity purified polyclonal antibody (PB9091) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for Beta Amyloid protein 42/APP at approximately 87-120 kDa. The expected band size for Beta Amyloid protein 42/APP is at 87 kDa.

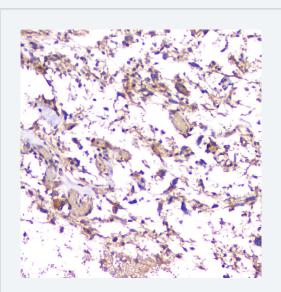


Figure 2. IHC analysis of Beta Amyloid protein 42/APP using anti-Beta Amyloid protein 42/APP antibody (PB9091). Beta Amyloid protein 42/APP was detected in a paraffinembedded section of human glioma tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was incubated with rabbit anti-Beta Amyloid protein 42/APP Antibody (PB9091) at a dilution of 1:200 and developed using Strepavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB (Catalog # AR1022) as the chromogen.

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Figure 7. IF analysis of APP using anti- APP antibody (PB9091). APP was detected in immunocytochemical section of A431 cell. Enzyme antigen retrieval was performed using IHC enzyme antigen retrieval reagent (AR0022) for 15 mins. The cells were blocked with 10% goat serum. And then incubated with 2µg/mL rabbit anti- APP Antibody (PB9091) overnight at 4°C. DyLight488 Conjugated Goat Anti-Rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution and incubated for 30 minutes at 37°C. The section was counterstained with DAPI. Visualize using a fluorescence microscope and filter sets appropriate for the label used.