

Acetyl Lysine (Biotin) Antibody

Rabbit Polyclonal Antibody Catalog # ABV11119

Specification

Acetyl Lysine (Biotin) Antibody - Product Information

Application WB, IF, E, IP
Host Rabbit
Clonality Polyclonal
Isotype Rabbit IgG

Acetyl Lysine (Biotin) Antibody - Additional Information

Application & Usage
Other Names
Acetyl Lysine

Western blot, IP, ELISA and IF.

Target/Specificity

Acetyl Lysine

Antibody Form

Liquid

AppearanceColorless liquid

Formulation

50 μg of antibody in 200 μl PBS, containing 0.01% sodium azide.

Handling

The antibody solution should be gently mixed before use.

Reconstitution & Storage

-20 °C

Background Descriptions

Precautions

Acetyl Lysine (Biotin) Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

Acetyl Lysine (Biotin) Antibody - Protein Information

Acetyl Lysine (Biotin) Antibody - Protocols





Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- <u>Immunohistochemistry</u>
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

Acetyl Lysine (Biotin) Antibody - Images

Acetyl Lysine (Biotin) Antibody - Background

Post-translational modifications of proteins play critical roles in the regulation and function of many known biological processes. Proteins can be post-translationally modified in many different ways, and a common post-transcriptional modification of lysine involves acetylation. The conserved amino-terminal domains of the four core histones (H2A, H2B, H3, and H4) contain lysines that are acetylated by histone acetyltransferases (HATs) and deacetylated by histone deacetylases (HDACs). Protein post-translational reversible lysine Nε-acetylation and deacetylation have been recognized as an emerging intracellular signaling mechanism that plays critical roles in regulating gene transcription, cell-cycle progression, apoptosis, DNA repair, and cytoskeletal organization. The regulation of protein acetylation status is impaired in the pathologies of cancer and polyglutamine diseases, and HDACs have become promising targets for anti-cancer dr µgs currently in development.