Pan Acetyl-Lysine Rabbit pAb

Catalog No.: A2391 34 Publications



Basic Information

Observed MW Refer to figures

Calculated MW

Category **Polyclonal Antibody**

Applications WB, IF/ICC, ELISA

Cross-Reactivity Human, Mouse, Rat, Other (Wide Range Predicted)

Background

Acetylation of lysine, like phosphorylation of serine, threonine or tyrosine, is an important reversible modification controlling protein activity. The conserved aminoterminal 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) . Signaling resulting in acetylation/deacetylation of histones, transcription factors, and other proteins affects a diverse array of cellular processes including chromatin structure and gene activity, cell growth, differentiation, and apoptosis . Recent proteomic surveys suggest that acetylation of lysine residues may be a widespread and important form of post-translational protein modification that affects thousands of proteins involved in control of cell cycle and metabolism, longevity, actin polymerization, and nuclear transport . The regulation of protein acetylation status is impaired in cancer and polyglutamine diseases, and HDACs have become promising targets for anti-cancer drugs currently in development .

Recommended Dilutions

WB	1:500 - 1:1000
IF/ICC	1:50 - 1:200
ELISA	Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

Immunogen Information

1:500 - 1:1000	Gene ID	Swiss Prot
1:50 - 1:200 Recommended starting concentration is 1 μg/mL. Please optimize the concentration based on your specific assay requirements.	Immunogen Synthetic peptide. This Synonyms	information is considered to be commercially sensitive.
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Product Information

G www.abclonal.com Source Rabbit

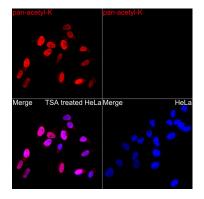
Isotype IgG

Purification Affinity purification

Storage

Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.09% sodium azide, 50% glycerol, pH7.3.

Validation Data



Immunofluorescence analysis of HeLa TSA and HeLa cells using Pan Acetyl-Lysine Rabbit pAb (A2391) at dilution of 1:50 (40x lens). Secondary antibody: Cy3-conjugated Goat anti-Rabbit IgG (H+L) (AS007) at 1:500 dilution. Blue: DAPI for nuclear staining.