For Research Use Only

Na N4-acetylcytidine Monoclonal antibody, PBS Only

Catalog Number:68498-1-PBS

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Antibodies | ELISA kits | Proteins
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Basic Information

Catalog Number:
68498-1-PBS

Size:
1 mg/ml

Source:
Mouse

Isotype:
IgG1

GenBank Accession Number:

GeneID (NCBI):

Full Name:

Purification Method:
Protein G purification

CloneNo.:
3C2C2

Applications

Tested Applications:
Indirect ELISA,ELISA,Dot Blot

Species Specificity:
Human, mouse, rat

Background Information

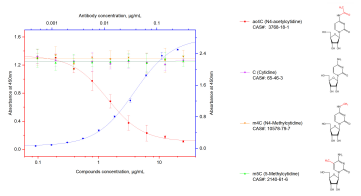
N4-Acetylcytidine, CasNo. 3768-18-1, is a modified nucleoside and endogenous urinary nucleoside product of the degradation of tRNA, 18s rRNA and mRNA. N4-Acetylcytidine is a biological marker for various cancers with elevated concentrations present in urine. N4-Acetylcytidine is also a partially protected cytidine and therefore can be used as a synthetic building block to prepare further derivatized nucleosides such as 2',3'-dideoxycytidine. NAT10 catalyzes the formation of N4-acetylcytidine (ac4C) modification on mRNAs, 18S rRNA and tRNAs.

Storage

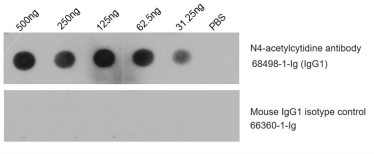
Storage:
Store at -80°C.
The product is shipped with ice packs. Upon receipt, store it immediately at -80°C
Storage Buffer:
PBS Only

For technical support and original validation data for this product please contact:
T: 4006900926 E: Proteintech-CN@ptglab.com W: ptgcn.com

Selected Validation Data



Indirect ELISA and competitive ELISA results show that this antibody is specific to ac4C (N4-acetylcytidine). Indirect ELISA (blue curve, refer to top X-right Y axis) was performed by coating BSA conjugated ac4C at 50ng/well followed by blocking with 5% non fat milk. Serial diluted primary antibody was added to the plates and incubated at 37℃. HRP-goat anti-mouse was used for detection. Competitive ELISA was performed similarly except that different concentration of Ac4C or



Total RNA was isolated from HeLa cell line and was dotted to NC membrane at different amount as indicated above the dots. The membrane was blocked with 5% milk and blotted with N4-acetylcytidine antibody 68498-1-Ig at 1:2000 followed by incubation of HRP-goat anti-mouse secondary antibody. Signal was developed by ECL substrate. A parallel dot blot was performed using Mouse IgG1 isotype control Monoclonal antibody 66360-1-Ig at the same dose. This data was