For Research Use Only

Acetyl-Histone H2A (Lys9) Recombinant antibody, PBS Only

Antibodies | ELISA kits | Proteins www.ptglab.com

Purification Method:

Protein A purfication

CloneNo.:

1017

Catalog Number:82823-2-PBS

Basic Information

Catalog Number:

82823-2-PBS

Concentration: 1 mg/ml

Source: Rabbit Isotype:

IgG

GenBank Accession Number:

BC062211

GeneID (NCBI): 221613

UNIPROT ID: Q96QV6 Full Name:

histone cluster 1, H2aa

Observed MW: 16 kDa

Applications

Tested Applications:

WB, Dot Blot, Indirect ELISA, ChIP-qPCR

Species Specificity:

human

Background Information

Storage

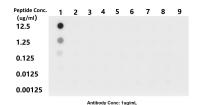
Storage:

Store at -80°C.

The product is shipped with ice packs. Upon receipt, store it immediately at -80°C

Storage Buffer: PBS only, pH7.3

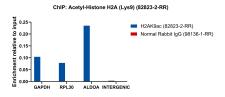
Selected Validation Data



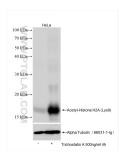
Dot blot analysis was used to confirm the specificity of Histone HZAK9ac antibody. Acetylated peptides were spotted onto NC and probed with antibody at 1 µg/ml.The amount of peptide (µg/mL) spotted is indicated next to each row.

Column 1: H2AK9ac. Column 2: Unmodified H2AK9. Column 3: H2AK5ac. Column 4: Unmodified H2AK Column 5: H2AK13Ac. Column 6: Unmodified H2AK13. Column 7: H2AK15ac. Column 8:Unmodified H2AK15. Column 9: Blank(PBS).

Dot blot analysis was used to confirm the specificity of Acetyl-Histone H2A (Lys9) antibody. Acetylated peptides were spotted onto NC and probed with antibody at 1 µg/ml. The amount of peptide (µ g/ml.) spotted is indicated next to each row. Column 1: H2AK9ac. Column 2: Unmodified H2AK9. Column 3: H2AK5ac. Column 4: Unmodified H2AK5. Column 5: H2AK13Ac. Column 6: Unmodified H2AK13. Column 7: H2AK15ac. Column 8:Unmodified H2AK15. Column 9:



Chromatin was prepared from HeLa cells. Cells were fixed with formaldehyde for 10 minutes. The ChIP was performed with 15 µg of cross-linked chromatin, 5 µg of Acetyl-Histone H2A (Lys9) (82823-2-RR) or 5 µg of Normal Rabbit IgG (98136-1-RR), and 20 µl of Protein A Magarose Beads. The immunoprecipitated DNA was quantified by real-time PCR. This data was developed using the same antibody clone with 82823-2-PBS in a different storage buffer formulation.



Trichostain A treated and untreated HeLa cell lysates were subjected to SDS PAGE followed by western blot with 82823-2-RR (Acetyl-Histone H2A (Lys9) antibody) at dilution of 1:2500 incubated at room temperature for 1.5 hours. The membrane was stripped and reblotted with Alpha Tubulin Monoclonal antibody (66031-1-lg) as loading control. This data was developed using the same antibody clone with 82823-2-PBS in a different storage buffer formulation.