

# Butyryl-HIST1H4A (K5) Antibody



PACO58647

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## Product Information

**Size:**

50ul

**Reactivity:**

Human

**Source:**

Rabbit

**Isotype:**

IgG

**Applications:**

ELISA, WB, IF, IP, ChIP

**Recommended dilutions:**

ELISA:1:2000-1:10000, WB:1:500-1:2000,  
IF:1:1-1:10, IP:1:200-1:2000

**Protein Background:**

Core component of nucleosome. Nucleosomes wrap and compact DNA into chromatin, limiting DNA accessibility to the cellular machineries which require DNA as a template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication and chromosomal stability. DNA accessibility is regulated via a complex set of post-translational modifications of histones, also called histone code, and nucleosome remodeling.

**Gene ID:**

HIST1H4A

**Uniprot**

P62805

**Synonyms:**

Histone H4, HIST1H4A; HIST1H4B; HIST1H4C; HIST1H4D; HIST1H4E; HIST1H4F; HIST1H4H; HIST1H4I; HIST1H4J; HIST1H4K; HIST1H4L; HIST2H4A; HIST2H4B; HIST4H4, H4/A H4FA; H4/I H4FI; H4/G H4FG; H4/B H4FB; H4/J H4FJ; H4/C H4FC; H4/H H4FH; H4/M H4FM; H4/E H4FE; H4/D H4FD; H4/K H4FK; H4/N H4F2 H4FN HIST2H4; H4/O H4FO;

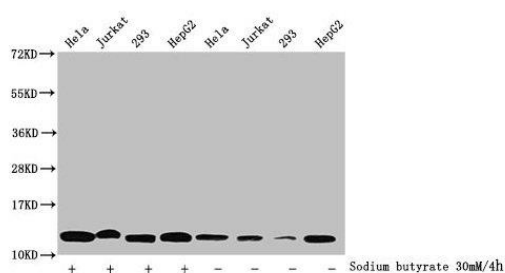
**Immunogen:**

Peptide sequence around site of Butyryl-Lys (5) derived from Human Histone H4.

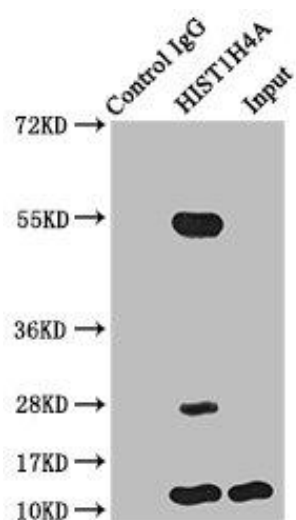
**Storage:**

Preservative: 0.03% Proclin 300. Constituents: 50% Glycerol, 0.01M PBS, pH 7.4

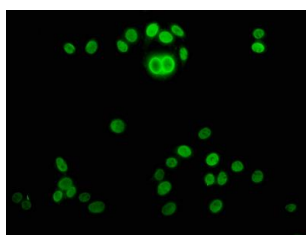
## Product Images



Western Blot. Detected samples: HeLa whole cell lysate, Jurkat whole cell lysate, 293 whole cell lysate, HepG2 whole cell lysate; Untreated (-) or treated (+) with 30mM sodium butyrate for 4h. All lanes: H1ST1H4A antibody at 1:1000. Secondary. Goat polyclonal to rabbit IgG at 1/40000 dilution. Predicted band size: 12 kDa. Observed band size: 12 kDa.



Immunoprecipitating H1ST1H4A in HepG2 whole cell lysate (treated with 30mM sodium butyrate for 4h). Lane 1: Rabbit control IgG instead of PACO58647 in HepG2 whole cell lysate (treated with 30mM sodium butyrate for 4h). For western blotting, a HRP-conjugated Protein G antibody was used as the secondary antibody (1/2000). Lane 2: PACO58647 (5µg) + HepG2 whole cell lysate (treated with 30mM sodium butyrate for 4h) (500µg). Lane 3: HepG2 whole cell lysate (treated with 30mM sodium butyrate for 4h) (20µg).



Immunofluorescence staining of HepG2 cells (treated with 30mM sodium butyrate for 4h) with PACO58647 at 1:5, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-conjugated AffiniPure Goat Anti-Rabbit IgG(H+L).