Crotonyl-HIST1H4A (K8) Antibody



PACO58663

Reactivity:

Human

Product Information

Size: Protein Background:

50ul 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

Source: nucleosome remodeling.

Rabbit Gene ID:

Isotype: HIST1H4A

lgG **Uniprot**

Applications: P62805

ELISA, WB, ICC, IF, ChIP Synonyms:

Recommended dilutions:

ELISA:1:2000-1:10000, WB:1:100-1:1000, ICC:1:10-1:100, IF:1:1-1:10

Histone H4, HIST1H4A; HIST1H4B; HIST1H4C; HIST1H4D; HIST1H4E; HIST1H4F; HIST1H4H; HIST1H4H; HIST1H4H; HIST1H4H; HIST1H4H; HIST1H4H; HIST2H4A; HIST2H4B; HIST4H4A, 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;

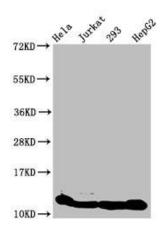
Immunogen:

Peptide sequence around site of Crotonyl-Lys (8) derived from Human Histone H4.

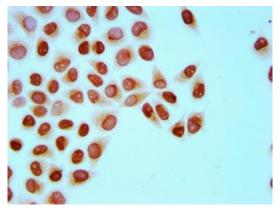
Storage:

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

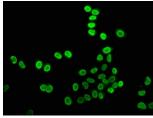
Product Images



Western Blot. Positive WB detected in: Hela whole cell lysate, Jurkat whole cell lysate, 293 whole cell lysate, HepG2 whole cell lysate (treated by 30mM sodium crotonylate for 4h). All lanes: HIST1H4A antibody at $1.6\mu g/ml$. Secondary. Goat polyclonal to rabbit lgG at 1/50000 dilution. Predicted band size: 12 kDa. Observed band size: 12 kDa.



Immunocytochemistry analysis of PACO58663 diluted at 1:15 and staining in Hela cells (treated with 30mM sodium crotonylate for 4h) performed on a Leica BondTM system. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



Immunofluorescence staining of Hela cells (treated with 30mM sodium crotonylate for 4h) with PACO58663 at 1:7.5, 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-congugated AffiniPure Goat Anti-Rabbit IqG(H+L).