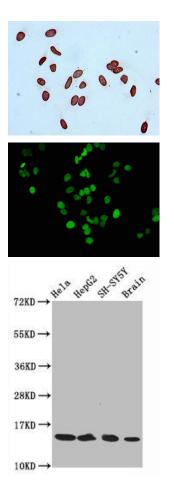
Acetyl-Histone H3.1 (K14) Recombinant Antibody

RACO0005



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 nucleosome remodeling.
Reactivity:	
Human, Rat	
Source:	
Human	Gene ID:
lsotype:	HIST1H3A
Rabbit IgG	Uniprot
Applications:	P68431
ELISA, WB, ICC, IF	Synonyms:
Recommended dilutions:	Histone H3.1, Histone H3/a, Histone H3/b, Histone H3/c, Histone H3/d, Histone H3/f, Histone H3/h, Histone H3/i, Histone H3/j, Histone H3/k, Histone H3/l, HIST1H3A, H3FA, AND, HIST1H3B, H3FL, AND, HIST1H3C, H3FC, AND, HIST1H3D, H3FB, AND, HIST1H3E, H3FD, AND, HIST1H3F, H3FI, AND, HIST1H3G, H3FH, AND, HIST1H3H, H3FK, AND, HIST1H3I, H3FF, AND, HIST1H3J, H3FJ
WB:1:500-1:2000, ICC:1:50-1:500, IF:1:30- 1:200	
	Immunogen:
	A synthesized peptide.
	Storage:
	Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and

Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.



Immunocytochemistry analysis of RACO0005 diluted at 1:100 and staining in Hela cells performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was 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 by 15mM sodium butyrate for 30min) with RACO0005 at 1:46, 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 IgG (H+L).

Western Blot

Positive WB detected in(Hela whole cell lysate) HepG2 whole cell lysate) SH-SY5Y whole cell lysate) Rat brain tissue All lanes: Acetyl-Histone H3.1(K14)antibody at 0.75µg/ml Secondary Goat polyclonal to rabbit IgG at 1:50000 dilution Predicted band size: 15 KDa Observed band size: 15 KDa