HIST1H3A (Ab-128) Antibody



PACO60631

Reactivity:

Product Information

Size:

50ul Core component of nucleosome. Nucleosomes wrap and compact DNA into chromatin,

Protein Background:

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

Human, Mouse, Rat replication and chromosomal stability. DNA accessibility is regulated via a cor of post-translational modifications of histones, also called histone code, and

Source: nucleosome remodeling.

Rabbit Gene ID:

Isotype: HIST1H3A

lgG Uniprot

Applications: P68431

ELISA, WB, IHC, IP Synonyms:

Recommended dilutions:

ELISA:1:2000-1:10000, WB:1:200-1:2000, IHC:1:10-1:100, IP:1:200-1:2000

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; HIST1H3B; HIST1H3C; HIST1H3D; HIST1H3E; HIST1H3F; HIST1H3G; HIST1H3H; HIST1H3I; HIST1H3J, H3FA; H3FL; H3FC; H3FB; H3FD; H3FI; H3FH; H3FK; H3FF; H3FJ

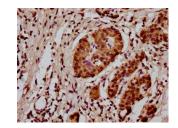
Immunogen:

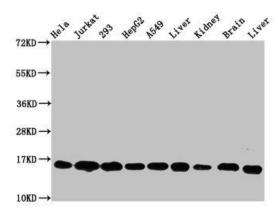
Peptide sequence around site of Arg (128) derived from Human Histone H3.1.

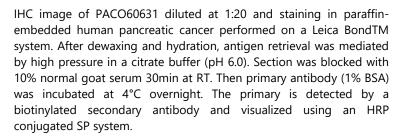
Storage:

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

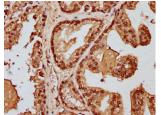
Product Images







Western Blot. Positive WB detected in: Hela whole cell lysate, Jurkat whole cell lysate, 293 whole cell lysate, HepG2 whole cell lysate, A549 whole cell lysate, Rat liver tissue, Rat kidney tissue, Mouse brain tissue, Mouse liver tissue. All lanes: HIST1H3A antibody at 0.53µg/ml. Secondary. Goat polyclonal to rabbit lgG at 1/50000 dilution. Predicted band size: 16 kDa. Observed band size: 16 kDa.



IHC image of PACO60631 diluted at 1:20 and staining in paraffinembedded human prostate cancer 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.