

## **Product datasheet for TA387652**

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## **H2BC4** Rabbit Polyclonal Antibody

**Product data:** 

**Product Type:** Primary Antibodies

Applications: ICC, IF, WB

Recommended Dilution: Recommended dilution: WB:1:500-1:2000, ICC:1:10-1:100, IF:1:1-1:10

Reactivity: Human
Host: Rabbit
Isotype: IgG

Clonality: Polyclonal

**Immunogen:** Peptide sequence around site of 2-hydroxyisobutyryl-Lys (12) derived from Human Histone

H2B type 1-C/E/F/G/I

**Formulation:** Preservative: 0.03% Proclin 300

Constituents: 50% Glycerol, 0.01M PBS, pH 7.4

**Concentration:** lot specific

**Purification:** Antigen Affinity Purified

Conjugation: Unconjugated

**Storage:** Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.

**Stability:** 1 year from dispatch.

Database Link: P62807

**Background:** Core component of nucleosome. Nucleosomes wrap and compact DNA into chromatin,

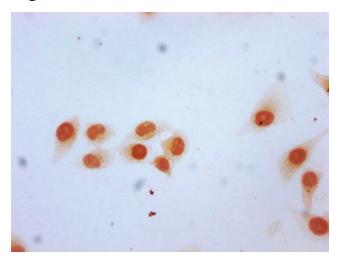
limiting DNA accessibility to the cellular machineries which require DNA as a template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication

and chromosomal stability. DNA accessibility is regulated via a complex set of post-

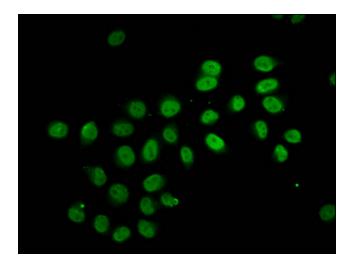
translational modifications of histones, also called histone code, and nucleosome remodeling.



## **Product images:**

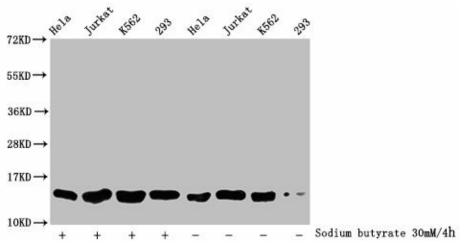


Immunocytochemistry analysis of TA387652 diluted at 1:15 and staining in Hela cells (treated with 30mM sodium butyrate for 4h) performed on a Leica BondTM system. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked with 10% normal goat serum 30min at RT. Then primary 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 butyrate for 4h) with TA387652 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 IgG(H+L).





Western Blot

Detected samples: Hela whole cell lysate, Jurkat whole cell lysate, K562 whole cell lysate, 293 whole cell lysate; Untreated (-) or treated (+) with

30mM sodium butyrate for 4h

All lanes: HIST1H2BC antibody at 1:500

Secondary

Goat polyclonal to rabbit IgG at 1/40000 dilution

Predicted band size: 14 kDa Observed band size: 14 kDa