

Product datasheet for TA387550

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H2AC11 Rabbit Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

Applications: ChIP, ICC, IF, WB

Recommended Dilution: Recommended dilution: WB:1:200-1:2000, ICC:1:20-1:200, IF:1:50-1:200

Reactivity: Human
Host: Rabbit
Isotype: IgG

Clonality: Polyclonal

Immunogen: Peptide sequence around site of Crotonyl-Lys (118) derived from Human Histone H2A type 1

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: POCOS8

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

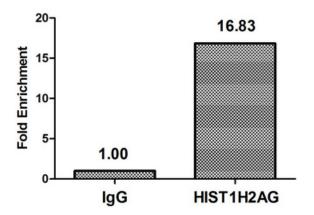
limiting DNA accessibility to the cellular machineries which require DNA as a template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication

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

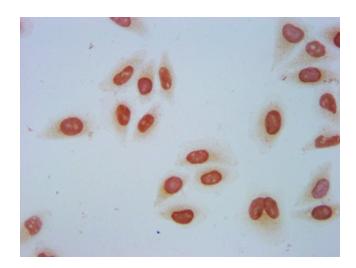
translational modifications of histones, also called histone code, and nucleosome remodeling.



Product images:

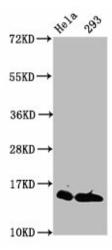


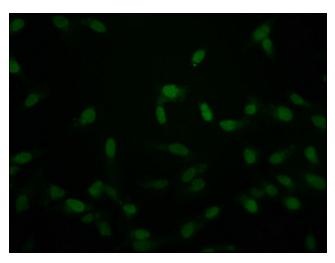
Chromatin Immunoprecipitation Hela (4*10⁶, treated with 30mM sodium crotonylate for 4h) were treated with Micrococcal Nuclease, sonicated, and immunoprecipitated with 5µg anti-HIST1H2AG (TA387550) or a control normal rabbit IgG. The resulting ChIP DNA was quantified using real-time PCR with primers against the β -Globin promoter.



Immunocytochemistry analysis of TA387550 diluted at 1:50 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.







Western Blot

Positive WB detected in: Hela whole cell lysate, 293 whole cell lysate (treated by 30mM sodium crotonylate for 4h)

All lanes: HIST1H2AG antibody at 0.3μg/ml Secondary

Goat polyclonal to rabbit IgG at 1/50000 dilution

Predicted band size: 15 kDa Observed band size: 15 kDa

Immunofluorescence staining of Hela cells with TA387550 at 1:50, 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).