

Product datasheet for TA387614

H2BC4 Rabbit Polyclonal Antibody

Product data:

OriGene Technologies, Inc.

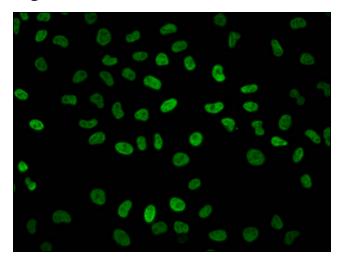
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Primary Antibodies
IF, IHC, WB
Recommended dilution: WB:1:100-1:1000, IHC:1:10-1:100, IF:1:1-1:10
Human
Rabbit
lgG
Polyclonal
Peptide sequence around site of Lys (23) derived from Human Histone H2B type 1-C/E/F/G/I
Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, pH 7.4
lot specific
Antigen Affinity Purified
Unconjugated
Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
1 year from dispatch.
<u>P62807</u>
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.

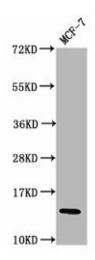


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Product images:

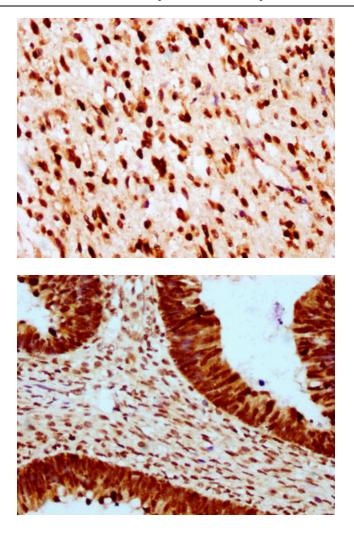


Immunofluorescence staining of Hela cells with TA387614 at 1:1, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).



Western Blot Positive WB detected in: MCF-7 whole cell lysate All lanes: HIST1H2BC antibody at 0.04µg/ml Secondary Goat polyclonal to rabbit IgG at 1/50000 dilution Predicted band size: 14 kDa Observed band size: 14 kDa

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IHC image of TA387614 diluted at 1:10 and staining in paraffin-embedded human glioma 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.

IHC image of TA387614 diluted at 1:10 and staining in paraffin-embedded human ovarian 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.

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