

# Anti-Histone H2B (glcnac S112) antibody ab130951

[4 References](#) [4 Images](#)

## Overview

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<b>Product name</b>	Anti-Histone H2B (glcnac S112) antibody
<b>Description</b>	Rabbit polyclonal to Histone H2B (glcnac S112)
<b>Host species</b>	Rabbit
<b>Specificity</b>	<p>ab130951 has been tested for specificity in peptide array. This product shows &lt;20% cross reactivity in peptide array with the unmodified peptide and does not cross react with GlcNAc S36, GlcNAc T101 or GlcNAc S47.</p> <p>From Mar 2024, QC testing of replenishment batches of this polyclonal changed. All tested and expected application and reactive species combinations are still covered by our Abcam product promise. However, we no longer test all applications. For more information on a specific batch, please contact our Scientific Support who will be happy to help.</p>
<b>Tested applications</b>	<b>Suitable for:</b> IHC-P, WB, ICC/IF, PepArr
<b>Species reactivity</b>	<b>Reacts with:</b> Cow, Human <b>Predicted to work with:</b> Mouse 
<b>Immunogen</b>	<p>Synthetic peptide corresponding to Human Histone H2B aa 100 to the C-terminus (glcnac S112) conjugated to keyhole limpet haemocyanin.</p> <p>Database link: <a href="#">Q16778</a> (Peptide available as <a href="#">ab166685</a>)</p>
<b>Positive control</b>	WB: Calf thymus histone preparation nuclear lysate. ICC/IF: MCF-7 cells. IHC-P: Human normal colon tissue.
<b>General notes</b>	<p>The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.</p> <p>If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&amp;As</p>

## Properties

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<b>Form</b>	Liquid
<b>Storage instructions</b>	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

#### Storage buffer

pH: 7.40  
Preservative: 0.02% Sodium azide  
Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising agent. If you would like information about the formulation of a specific lot, please contact our scientific support team who will be happy to help.

#### Purity

Immunogen affinity purified

#### Clonality

Polyclonal

#### Isotype

IgG

### Applications

#### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab130951 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IHC-P		Use a concentration of 1 µg/ml. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
WB		Use a concentration of 1 µg/ml. Detects a band of approximately 18 kDa (predicted molecular weight: 14 kDa). Can be blocked with <b>Human Histone H2B (glcnac S112) peptide (ab166685)</b> .
ICC/IF		Use a concentration of 1 µg/ml.
PepArr		Use a concentration of 0.2 - 0.02 µg/ml.

### Target

#### Relevance

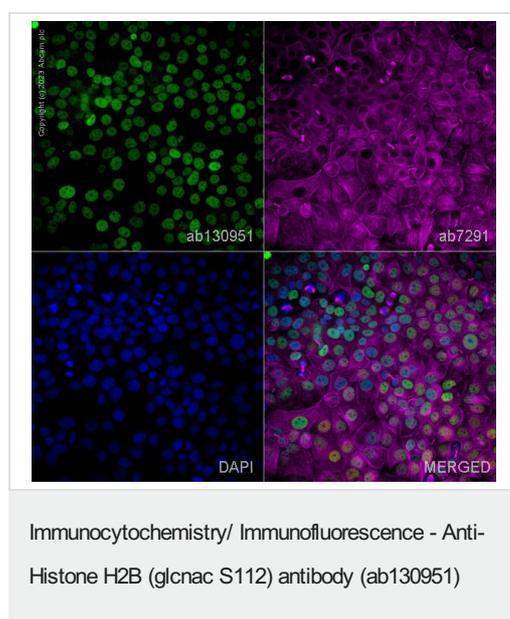
Core component of nucleosome. Nucleosomes wrap and compact DNA into chromatin, limiting DNA accessibility to the cellular machineries which require DNA as a template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication and chromosomal stability. DNA accessibility is regulated via a complex set of post-translational modifications of histones, also called histone code, and nucleosome remodeling. Subunit structure The nucleosome is a histone octamer containing two molecules each of H2A, H2B, H3 and H4 assembled in one H3-H4 heterotetramer and two H2A-H2B heterodimers. The octamer wraps approximately 147 bp of DNA. Post-translational modification Monoubiquitination at Lys-35 (H2BK34Ub) by the MSL1/MSL2 dimer is required for histone H3 'Lys-4' (H3K4me) and 'Lys-79' (H3K79me) methylation and transcription activation at specific gene loci, such as HOXA9 and MEIS1 loci. Similarly, monoubiquitination at Lys-121 (H2BK120Ub) by the RNF20/40 complex gives a specific tag for epigenetic transcriptional activation and is also prerequisite for histone H3 'Lys-4' and 'Lys-79' methylation. It also functions cooperatively with the FACT dimer to stimulate elongation by RNA polymerase II. H2BK120Ub also acts as a regulator of mRNA splicing: deubiquitination by USP49 is required for efficient cotranscriptional splicing of a large set of exons. Phosphorylation at Ser-37 (H2BS36ph) by AMPK in response to stress promotes

transcription. Phosphorylated on Ser-15 (H2BS14ph) by STK4/MST1 during apoptosis; which facilitates apoptotic chromatin condensation. Also phosphorylated on Ser-15 in response to DNA double strand breaks (DSBs), and in correlation with somatic hypermutation and immunoglobulin class-switch recombination. GlcNAcylation at Ser-113 promotes monoubiquitination of Lys-121. It fluctuates in response to extracellular glucose, and associates with transcribed genes. Crotonylation (Kcr) is specifically present in male germ cells and marks testis-specific genes in post-meiotic cells, including X-linked genes that escape sex chromosome inactivation in haploid cells. Crotonylation marks active promoters and enhancers and confers resistance to transcriptional repressors. It is also associated with post-meiotically activated genes on autosomes.

## Cellular localization

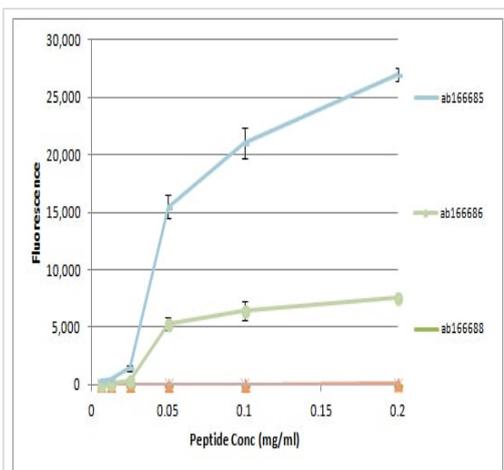
Nuclear

## Images



ab130951 staining Histone H2B (glcna S112) in MCF7 cells. The cells were fixed with 4% paraformaldehyde (10 min), permeabilized with 0.1% PBS-Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated overnight at 4°C with ab130951 at 1µg/ml and [ab7291](#), Mouse monoclonal [DM1A] to alpha Tubulin - Loading Control. Cells were then incubated with [ab150081](#), Goat polyclonal Secondary Antibody to Rabbit IgG - H&L (Alexa Fluor® 488), pre-adsorbed at 1/1000 dilution (shown in green) and [ab150120](#), Goat polyclonal Secondary Antibody to Mouse IgG - H&L (Alexa Fluor® 594), pre-adsorbed at 1/1000 dilution (shown in pseudocolour magenta). Nuclear DNA was labelled with DAPI (shown in blue).

Image was acquired with a high-content analyser (Operetta CLS, Perkin Elmer) and a maximum intensity projection of confocal sections is shown.



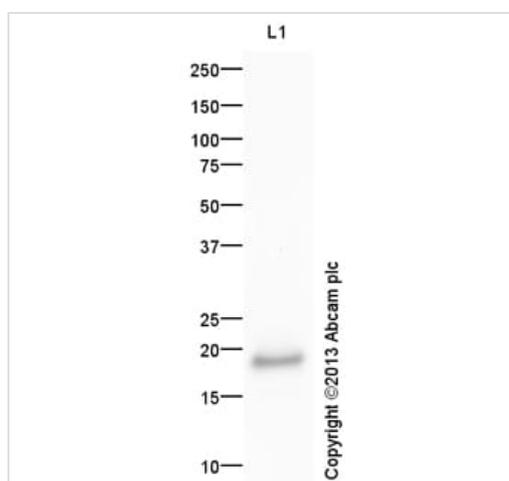
Peptide Array - Anti-Histone H2B (glcnac S112) antibody (ab130951)

All batches of ab130951 are tested in Peptide Array against peptides to different Histone H2B modifications. Six dilutions of each peptide are printed on to the Peptide Array in triplicate and results are averaged before being plotted on to a graph. Results show strong binding to Histone H2B - GlcNAc S112 peptide (**ab166685**), indicating that this antibody recognises the Histone H2B - GlcNAc S112 modification. This product shows <20% cross reactivity in peptide array with the unmodified peptide and does not cross react with GlcNAc S36, GlcNAc T101 or GlcNAc S47.

**ab166685** - Histone H2B - GlcNAc S112

**ab166686** - Histone H2B - unmodified

**ab166688** - Histone H2B - GlcNAc S36



Western blot - Anti-Histone H2B (glcnac S112) antibody (ab130951)

Anti-Histone H2B (glcnac S112) antibody (ab130951) at 1 µg/ml + Calf Thymus Histone Preparation Lysate at 0.25 µg

### Secondary

Goat Anti-Rabbit IgG H&L (HRP) (**ab97051**) at 1/10000 dilution

Developed using the ECL technique.

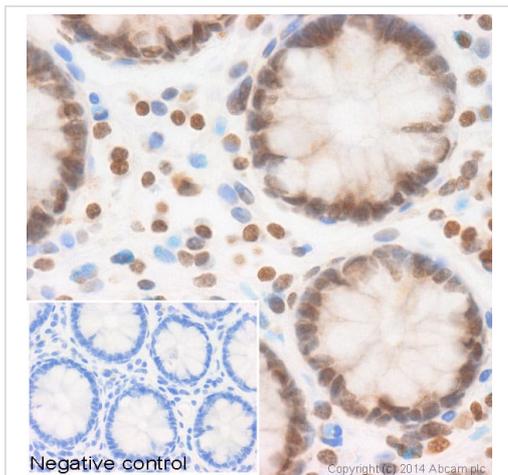
Performed under reducing conditions.

**Predicted band size:** 14 kDa

**Observed band size:** 18 kDa

**Exposure time:** 4 minutes

This blot was produced using a 10% Bis-tris gel under the MES buffer system. The gel was run at 200V for 35 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 5% Bovine Serum Albumin before being incubated with ab130951 overnight at 4°C. Antibody binding was detected using an anti-rabbit antibody conjugated to HRP, and visualised using ECL development solution.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Histone H2B (glcnac S112) antibody (ab130951)

IHC image of ab130951 staining Histone H2B (glcnac S112) in human colon formalin fixed paraffin embedded tissue sections, performed on a Leica Bond. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab130951, 1µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX. No primary antibody was used in the negative control (shown on the inset).

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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