## abcam

### **Product datasheet**

# Anti-Histone H2B (mono methyl K5) antibody - ChIP Grade ab12929

★★★★★ <u>5 Abreviews</u> <u>4 References</u> 5 图像

概述			
产品名称	Anti-Histone H2B (mono methyl K5) <b>抗体-</b> ChIP Grade		
描述	<b>兔多克隆抗体</b> to Histone H2B (mono methyl K5) - ChIP Grade		
宿主	Rabbit		
特异性	Mono methylation of Histone H2B K5 is a putative modification site. <b>ab22512</b> in ELISA specifically recognises mono-methyl K5 histone H2B peptide but not the corresponding unmodified histone H2B peptide. From Jan 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.		
经测试应 <b>用</b>	适用于: IHC-P, IP, WB, ChIP		
<b>种属反应性</b>	与反应: Cow, Human		
	预测可用于: Mouse, Rat, Chicken, Xenopus laevis 🛛 🔺		
免疫原	Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.		
<b>常</b> 规说 <b>明</b>	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.		
	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&As		

性能	
形式	Liquid
存 <b>放</b> 说明	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or - 80°C. Avoid freeze / thaw cycle.
存储溶液	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<br/>agent. If you would like information about the formulation of a specific lot, please contact our<br/>scientific support team who will be happy to help.纯度Immunogen affinity purified克隆多克隆同种型IgG

应用

## The Abpromise guarantee Abpromise™</u>承诺保证使用ab12929于以下的经测试应用 "应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
IHC-P		Use a concentration of 0.1 $\mu$ g/ml. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.
IP		Use at an assay dependent concentration.
WB	★ ★ ★ ★ ★ <u>(3)</u>	Use a concentration of 1 µg/ml.
ChIP	★ ★ ★ ★ ★ <u>(1)</u>	Use 2-5 µg for 25 µg of chromatin.

靶标

相关性

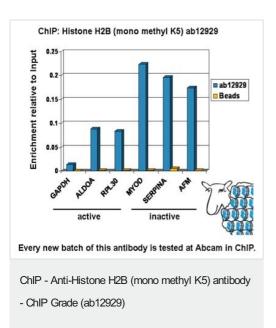
Core component of nucleosome. Nucleosomes wrap and compact DNA into chromatin, limiting DNA accessibility to the cellular machineries which require DNA as a template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication and chromosomal stability. DNA accessibility 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 monoubiguitination 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.

#### Nuclear

#### 图片

细胞定位



1 2 3

- d

(c) 2007 Aboam

150 kDa — 100 kDa — 75 kDa —

50 kDa 🗕

37 kDa -

25 kDa 🗕

20 kDa =

15 kDa

antibody - ChIP Grade (ab12929)

Western blot - Anti-Histone H2B (mono methyl K5)

Chromatin was prepared from Hela cells according to the Abcam X-ChIP protocol. Cells were fixed with formaldehyde for 10 min. The ChIP was performed with 25 µg of chromatin, 2 µg of ab12929 (blue), and 20 µl of Protein A/G sepharose beads. No antibody was added to the beads control (yellow). The immunoprecipitated DNA was quantified by real time PCR (Taqman approach). Primers and probes are located in the first kb of the transcribed region.

**All lanes :** Anti-Histone H2B (mono methyl K5) antibody - ChIP Grade (ab12929) at 1 µg/ml

Lane 1 : Calf Thymus Histone Preparation Nuclear Lysate (<u>ab121</u>)
Lane 2 : Calf Thymus Histone Preparation Nuclear Lysate (<u>ab121</u>)
with Human Histone H2B (mono methyl K5) peptide (<u>ab13211</u>)
Lane 3 : Calf Thymus Histone Preparation Nuclear Lysate (<u>ab121</u>)
with Human Histone H2B peptide (<u>ab13212</u>)

Lysates/proteins at 20 µg per lane.

#### Secondary

**All lanes :** IRDye 680 Conjugated Goat Anti-Rabbit lgG (H+L) at 1/10000 dilution

Performed under reducing conditions.

Predicted band size: 14 kDa



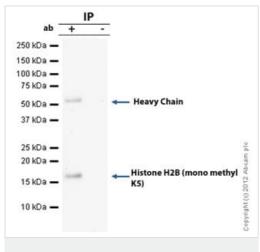
ab12929 at a 1/600 dilution for ChIP analysis of mouse dorsal skin epidermis whole tissue lysate, incubated for 15 hours at 4°C with ChIP dilution buffer. Cross-linking (X-ChIP) using 1% formaldehyde for 10 minutes.

Detection step: Semiquantitative PCR. Negative control: Rabbit lgG. Cells untreated.

ChIP - Anti-Histone H2B (mono methyl K5) antibody

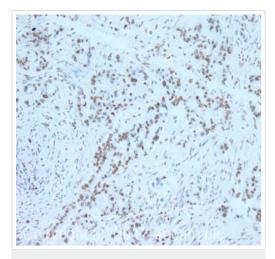
#### - ChIP Grade (ab12929)

This image is courtesy of an anonymous abreview.



Immunoprecipitation - Anti-Histone H2B (mono methyl K5) antibody - ChIP Grade (ab12929) Histone H2B (mono methyl K5) was immunoprecipitated using 0.5mg Hela whole cell extract, 5µg of Rabbit polyclonal to Histone H2B (mono methyl K5) - ChIP Grade and 50µl of protein G magnetic beads (+). No antibody was added to the control (-). The antibody was incubated under agitation with Protein G beads for 10min, Hela whole cell extract lysate diluted in RIPA buffer was added to each sample and incubated for a further 10min under agitation.

Proteins were eluted by addition of 40µl SDS loading buffer and incubated for 10min at 70°C; 10µl of each sample was separated on a SDS PAGE gel, transferred to a nitrocellulose membrane, blocked with 5% BSA and probed with ab12929. Secondary: Clean blot (HRP conjugate) at 1/1000 dilution. Band: 17kDa: Histone H2B (mono methyl K5).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Histone H2B (mono methyl K5) antibody - ChIP Grade (ab12929) IHC image of Histone H2B (mono methyl K5) staining in human pancreatic adenocarcinoma formalin fixed paraffin embedded tissue section, performed on a Leica Bond <sup>™</sup> system using the standard protocol F. 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 ab12929, 0.1µg/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.

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.

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