abcam

Product datasheet

Anti-Histone H3 (tri methyl K4) antibody [mAbcam12209] - ChIP Grade ab12209

★★★★★ 7 Abreviews 44 References 画像数 7

製品の概要

免疫原

製品名 Anti-Histone H3 (tri methyl K4) antibody [mAbcam12209] - ChIP Grade

製品の詳細 Mouse monoclonal [mAbcam12209] to Histone H3 (tri methyl K4) - ChIP Grade

由来種 Mouse

特異性 ab12209 is strongly blocked in Western blotting on histones by tri methyl K4, weakly by di methyl

> K4 and very weakly by mono methyl K4 peptides. It is not blocked by non-modified peptides. By ELISA the antibody binds to the tri methyl K4 peptide and at high antibody concentrations to di and mono methyl K4 peptides. It does not bind to unmodified, mono, di or tri methyl K9 or di or tri methyl K27 peptides. Not suitable for blocking with milk in Western blot (see Application notes).

> Our internal testing has revealed that the supernatant contains a very small fraction of IgG2b. IgG1

remains the main isotype.

適用あり: WB, ICC/IF, ChIP, ELISA, Flow Cyt (Intra) アプリケーション

種交差性 交差種: Rat, Cow, Human

> 交差が予測される動物種: Mouse, Saccharomyces cerevisiae, Xenopus laevis, Arabidopsis thaliana, Caenorhabditis elegans, Drosophila melanogaster, Schizosaccharomyces pombe,

Zebrafish, Mammals, Neurospora crassa

Synthetic peptide within Human Histone H3 aa 1-100 (tri methyl K4) conjugated to keyhole limpet

haemocyanin. The exact sequence is proprietary.

(Peptide available as ab1342)

ポジティブ・コントロール Flow Cyt (Intra): HeLa cells; ICC/IF: HeLa cells; WB: Calf Thymus Histone Preparation Nuclear

Lysate; ChIP: U2OS cells.

特記事項 This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or

conjugation for your experiments, please contact orders@abcam.com.

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

製品の特性

製品の状態 Liquid

保存方法 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.

Sp2/0-Ag14

バッファー pH: 7.40

Preservative: 0.02% Sodium azide Constituents: PBS, 6.97% L-Arginine

 精製度
 IgG fraction

 ポリ/モノ
 モノクローナル

 クローン名
 mAbcam12209

アイソタイプ lgG1 **軽鎖の種類** kappa

アプリケーション

ミエローマ

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab12209の使用に適用されますアプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
WB	★ ★ ★ ★ ★ (2)	Use a concentration of 1 - 5 µg/ml. Detects a band of approximately 17 kDa (predicted molecular weight: 15 kDa).Can be blocked with <u>Human Histone H3 (tri methyl K4) peptide</u> (ab1342). NOT SUITABLE for blocking with milk. Block in 5% BSA for 1 hour. Our labs have investigated the blocking conditions for this antibody and found that milk significantly decreases the signal and is therefore not a suitable blocking
ICC/IF	★★★☆☆ (1)	Use a concentration of 5 µg/ml.
ChIP	★★★★ <u>(3)</u>	Use 2-5 µg for 25 µg of chromatin.
ELISA		Use at an assay dependent concentration.
Flow Cyt (Intra)		Use 1µg for 10 ⁶ cells. ab170190 - Mouse monoclonal lgG1, is suitable for use as an isotype control with this antibody.

ターゲット情報

機能

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.

配列類似性

発生段階

翻訳後修飾

Belongs to the histone H3 family.

Expressed during S phase, then expression strongly decreases as cell division slows down during the process of differentiation.

Acetylation is generally linked to gene activation. Acetylation on Lys-10 (H3K9ac) impairs methylation at Arg-9 (H3R8me2s). Acetylation on Lys-19 (H3K18ac) and Lys-24 (H3K24ac) favors methylation at Arg-18 (H3R17me).

Citrullination at Arg-9 (H3R8ci) and/or Arg-18 (H3R17ci) by PAD4 impairs methylation and represses transcription.

Asymmetric dimethylation at Arg-18 (H3R17me2a) by CARM1 is linked to gene activation. Symmetric dimethylation at Arg-9 (H3R8me2s) by PRMT5 is linked to gene repression. Asymmetric dimethylation at Arg-3 (H3R2me2a) by PRMT6 is linked to gene repression and is mutually exclusive with H3 Lys-5 methylation (H3K4me2 and H3K4me3). H3R2me2a is present at the 3' of genes regardless of their transcription state and is enriched on inactive promoters, while it is absent on active promoters.

Methylation at Lys-5 (H3K4me), Lys-37 (H3K36me) and Lys-80 (H3K79me) are linked to gene activation. Methylation at Lys-5 (H3K4me) facilitates subsequent acetylation of H3 and H4. Methylation at Lys-80 (H3K79me) is associated with DNA double-strand break (DSB) responses and is a specific target for TP53BP1. Methylation at Lys-10 (H3K9me) and Lys-28 (H3K27me) are linked to gene repression. Methylation at Lys-10 (H3K9me) is a specific target for HP1 proteins (CBX1, CBX3 and CBX5) and prevents subsequent phosphorylation at Ser-11 (H3S10ph) and acetylation of H3 and H4. Methylation at Lys-5 (H3K4me) and Lys-80 (H3K79me) require preliminary monoubiquitination of H2B at 'Lys-120'. Methylation at Lys-10 (H3K9me) and Lys-28 (H3K27me) are enriched in inactive X chromosome chromatin.

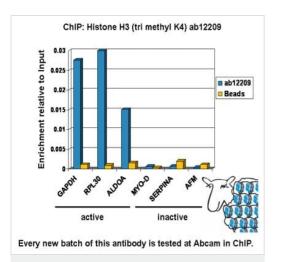
Phosphorylated at Thr-4 (H3T3ph) by GSG2/haspin during prophase and dephosphorylated during anaphase. Phosphorylation at Ser-11 (H3S10ph) by AURKB is crucial for chromosome condensation and cell-cycle progression during mitosis and meiosis. In addition phosphorylation at Ser-11 (H3S10ph) by RPS6KA4 and RPS6KA5 is important during interphase because it enables the transcription of genes following external stimulation, like mitogens, stress, growth factors or UV irradiation and result in the activation of genes, such as c-fos and c-jun. Phosphorylation at Ser-11 (H3S10ph), which is linked to gene activation, prevents methylation at Lys-10 (H3K9me) but facilitates acetylation of H3 and H4. Phosphorylation at Ser-11 (H3S10ph) by AURKB mediates the dissociation of HP1 proteins (CBX1, CBX3 and CBX5) from heterochromatin. Phosphorylation at Ser-11 (H3S10ph) is also an essential regulatory mechanism for neoplastic cell transformation. Phosphorylated at Ser-29 (H3S28ph) by MLTK isoform 1, RPS6KA5 or AURKB during mitosis or upon ultraviolet B irradiation. Phosphorylation at Thr-7 (H3T6ph) by PRKCBB is a specific tag for epigenetic transcriptional activation that prevents demethylation of Lys-5 (H3K4me) by LSD1/KDM1A. At centromeres, specifically phosphorylated at Thr-12 (H3T11ph) from prophase to early anaphase, by DAPK3 and PKN1. Phosphorylation at Thr-12 (H3T11ph) by PKN1 is a specific tag for epigenetic transcriptional activation that promotes demethylation of Lys-10 (H3K9me) by KDM4C/JMJD2C. Phosphorylation at Tyr-42 (H3Y41ph) by JAK2 promotes exclusion of CBX5 (HP1 alpha) from chromatin.

Monoubiquitinated by RAG1 in lymphoid cells, monoubiquitination is required for V(D)J recombination (By similarity). Ubiquitinated by the CUL4-DDB-RBX1 complex in response to ultraviolet irradiation. This may weaken the interaction between histones and DNA and facilitate

細胞内局在

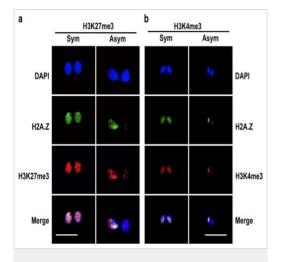
Nucleus. Chromosome.

DNA accessibility to repair proteins.



ChIP - Anti-Histone H3 (tri methyl K4) antibody [mAbcam12209] - ChIP Grade (ab12209)

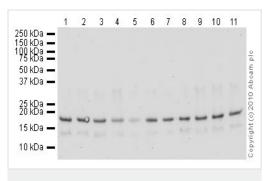
Chromatin was prepared from U2OS cells according to the Abcam X-ChIP protocol. Cells were fixed with formaldehyde for 10min. The ChIP was performed with 25 μg of chromatin, 2 μg of ab12209 (blue), and 20 μl of Protein A/G sepharose beads. No antibody was added to the beads control (yellow). The immunoprecipitated DNA was quantified by real time PCR (Taqman approach). Primers and probes are located in the first kb of the transcribed region.



Ab12209 staining Histone H3 (Tri Methyl K4) in Mouse hair follicle DSCs by ICC/IF (Immunocytochemistry/Immunofluorescence). Cells were fixed with 3.7% paraformaldehyde, permeabilised with 0.2% Triton X-100 and blocked with 10% normal goat serum in PBS. Samples were incubated with primary antibody at 1:200 dilution. An Alexa Fluor [®] 568 conjugated goat anti-mouse IgG was used as a secondary antibody.

Immunocytochemistry/ Immunofluorescence - Anti-Histone H3 (tri methyl K4) antibody [mAbcam12209] - ChIP Grade (ab12209)

Image from Huh YH et al., Cell Death Dis 4(5). Doi: 10.1038/cddis.2014.522. Reproduced under the Creative Commons license http://creativecommons.org/licenses/by/3.0



Western blot - Anti-Histone H3 (tri methyl K4) antibody [mAbcam12209] - ChIP Grade (ab12209)

All lanes : Anti-Histone H3 (tri methyl K4) antibody [mAbcam12209] - ChIP Grade (ab12209) at 2 µg/ml

Lane 1 : Calf Thymus Histone Preparation Nuclear Lysate **Lane 2 :** Calf Thymus Histone Preparation Nuclear Lysate with Human Histone H3 (unmodified) peptide (<u>ab7228</u>) at $0.25~\mu g/ml$ **Lane 3 :** Calf Thymus Histone Preparation Nuclear Lysate with Human Histone H3 (mono methyl K4) peptide (<u>ab1340</u>) at $0.25~\mu g/ml$

Lane 4: Calf Thymus Histone Preparation Nuclear Lysate with Human Histone H3 (di methyl K4) peptide (ab7768) at 0.25 µg/ml

Lane 5: Calf Thymus Histone Preparation Nuclear Lysate with Human Histone H3 (tri methyl K4) peptide (ab1342) at 0.25 µg/ml

Lane 6: Calf Thymus Histone Preparation Nuclear Lysate with

Human Histone H3 (mono methyl K9) peptide (ab1771) at 0.25

µg/ml

Lane 7: Calf Thymus Histone Preparation Nuclear Lysate with Human Histone H3 (di methyl K9) peptide (ab1772) at 0.25 μg/ml

Lane 8: Calf Thymus Histone Preparation Nuclear Lysate with

Human Histone H3 (tri methyl K9) peptide (ab1773) at 0.25 μg/ml

Lane 9: Calf Thymus Histone Preparation Nuclear Lysate with

Human Histone H3 (mono methyl K27) peptide (ab1780) at 0.25

µg/ml

Lane 10: Calf Thymus Histone Preparation Nuclear Lysate with Human Histone H3 (di methyl K27) peptide (ab1781) at 0.25 μg/ml

Lane 11: Calf Thymus Histone Preparation Nuclear Lysate with

Human Histone H3 (tri methyl K27) peptide (ab1782) at 0.25 μg/ml

Lysates/proteins at 0.5 µg per lane.

Secondary

All lanes: Goat polyclonal to Mouse IgG - H&L - Pre-Adsorbed

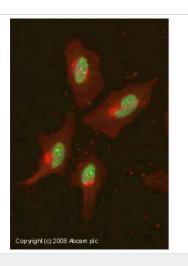
(HRP) at 1/3000 dilution

Performed under reducing conditions.

Predicted band size: 15 kDa

Observed band size: 17 kDa

Exposure time: 16 minutes



Immunocytochemistry/ Immunofluorescence - Anti-Histone H3 (tri methyl K4) antibody [mAbcam12209] - ChIP Grade (ab12209)

ICC/IF image of ab12209 stained human HeLa cells. The cells were 4% PFA fixed (10 min), permabilised in 0.1% PBS-Tween (20 min) and incubated with the antibody (ab12209, 5µg/ml) for 1h at room temperature. 1%BSA / 10% normal goat serum / 0.3M glycine was used to block non-specific protein-protein interactions. The secondary antibody (green) was Alexa Fluor® 488 goat anti-mouse lgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red). DAPI was used to stain the cell nuclei (blue).

ELISA using ab12209 at varying antibody concentrations.

Curve SPL4 indicates binding to the tri methyl K4 peptide

mono methyl K4 peptide ab1340 (Curve_SPL2).

Binding to the following peptides was not seen:

K27, SPL10 Histone H3 tri methyl K27.

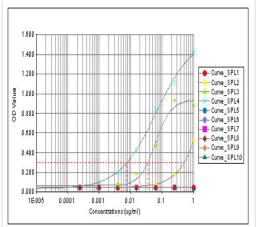
ab1342. In addition, SPL3 indicates partial binding to the di methyl

K4 peptide ab7768. There is very weak cross-reactivity with the

SPL1 unmodified Histone H3, SPL5 Histone H3 mono methyl K9,

SPL6 Histone H3 di methyl K9, SPL7 Histone H3 tri methyl K9,

SPL8 Histone H3 mono methyl K27, SPL9 Histone H3 di methyl



ELISA - Anti-Histone H3 (tri methyl K4) antibody [mAbcam12209] - ChIP Grade (ab12209)

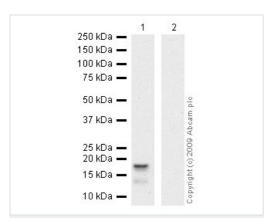


Lane 1: Anti-Histone H3 (tri methyl K4) antibody [mAbcam12209] -ChIP Grade (ab12209) at 1 µg/ml (BLOCKED WITH 5% BSA)



All lanes: Calf Thymus Histone Preparation Nuclear Lysate

Lysates/proteins at 0.5 µg per lane.



Western blot - Anti-Histone H3 (tri methyl K4) antibody [mAbcam12209] - ChIP Grade (ab12209)

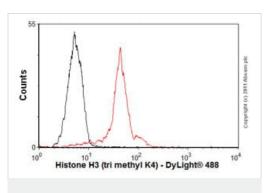
Secondary

All lanes: Goat polyclonal to Mouse IgG - H&L - Pre-Adsorbed (HRP) at 1/3000 dilution

Performed under reducing conditions.

Predicted band size: 15 kDa **Observed band size:** 17 kDa

Exposure time: 12 minutes



Flow Cytometry (Intracellular) - Anti-Histone H3 (tri methyl K4) antibody [mAbcam12209] - ChIP Grade (ab12209)

Overlay histogram showing HeLa cells stained with ab12209 (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab12209, 1 μ g/1x10⁶ cells) for 30 min at 22°C. The secondary antibody used was a goat **anti-mouse DyLight® 488** (lgG; H+L) (**ab96879**) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse lgG1 [ICIGG1] (**ab91353**, 2 μ g/1x10⁶ cells) used under the same conditions. Acquisition of >5,000 events was performed.

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