abcam

Product datasheet

Anti-Histone H3 (di methyl K4) antibody - ChIP Grade ab7766

★★★★★ 25 Abreviews 304 References 画像数 8

製品の概要

免疫原

製品名 Anti-Histone H3 (di methyl K4) antibody - ChIP Grade

製品の詳細 Rabbit polyclonal to Histone H3 (di methyl K4) - ChIP Grade

由来種 Rabbit

特異性 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. You may also be interested in our

alternative recombinant antibody, ab176878.

アプリケーション 適用あり: ChIP, WB, PepArr, ICC/IF

種交差性 交差種: Mouse, Rat, Cow, Human

交差が予測される動物種: Pig, Saccharomyces cerevisiae, Tetrahymena, Drosophila melanogaster, Schizosaccharomyces pombe, Mammals, Plasmodium falciparum, Common

marmoset, Candida albicans

Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

(Peptide available as ab7768)

ポジティブ・コントロール ChIP: Chromatin prepared from HeLa cells. WB: HeLa, NIH/3T3 and PC12 nuclear lysate (triton

enriched). Calf Thymus Histone Preparation Nuclear Lysate. ICC/IF: HeLa cells.

特記事項 For detection of Histone H3 specifically methylated at position Lys 4. This antibody was used in a

screen by Dover et al, (2002) to isolate yeast mutants that are unable to methylate Lysine 4. In immunofluorescence, this antibody detects foci in the nucleus that are non-colocalising with condensed chromatin. The perinuclear and perinucleolar heterochromatin are not stained with this

antibody.

Learn about ChIP assay kits, other ChIP antibodies, protocols and more in the **ChIP assay**

<u>guide</u>.

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

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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.

バッファー 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.

精製度 Immunogen affinity purified

一次抗体 備考 For detection of Histone H3 specifically methylated at position Lys 4. This antibody was used in a

screen by Dover et al, (2002) to isolate yeast mutants that are unable to methylate Lysine 4. In immunofluorescence, this antibody detects foci in the nucleus that are non-colocalising with condensed chromatin. The perinuclear and perinucleolar heterochromatin are not stained with this

antibody.

ポリ/モノ ポリクローナル

アイソタイプ IgG

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab7766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
ChIP	★★★★☆(3)	Use 2 µg for 25 µg of chromatin. Use ALDOA ChIP primer pair <u>ab269260</u> as positive control.
WB	**** (<u>11)</u>	Use a concentration of 1 µg/ml. Detects a band of approximately 17 kDa (predicted molecular weight: 15 kDa).
PepArr		Use a concentration of 0.2 - 2 µg/ml.
ICC/IF	★★★★☆ (6)	Use a concentration of 5 µg/ml.

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 Belongs to the histone H3 family.

発生段階 Expressed during S phase, then expression strongly decreases as cell division slows down

翻訳後修飾

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.

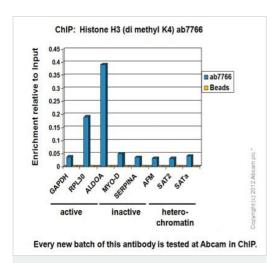
Monoubiquitinated by RAG1 in lymphoid cells, monoubiquitination is required for V(D)J recombination (By similarity). Ubiquitinated by the CUL4-DDB-RBX1 complex in response to ultraviolet irradiation. This may weaken the interaction between histones and DNA and facilitate

DNA accessibility to repair proteins.

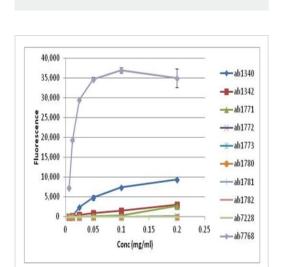
Nucleus. Chromosome.

細胞内局在

画像



ChIP - Anti-Histone H3 (di methyl K4) antibody - ChIP Grade (ab7766)



Peptide Array - Anti-Histone H3 (di methyl K4) antibody - ChIP Grade (ab7766)

Chromatin was prepared from HeLa cells according to the Abcam X-ChIP protocol. Cells were fixed with formaldehyde for 10 minutes. The ChIP was performed with 25µg of chromatin, 2µg of ab7766 (blue), and 20µl 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 for active and inactive loci, Sybr green approach for heterochromatic loci). Primers and probes are located in the first kb of the transcribed region.

All batches of ab7766 are tested in Peptide Array against peptides to different Histone H3 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 H3 - di methyl K4 peptide (ab7768), indicating that this antibody specifically recognises the Histone H3 - di methyl K4 modification.

ab1340 - Histone H3 - mono methyl K4

ab1342 - Histone H3 - tri methyl K4

ab1771 - Histone H3 - mono methyl K9

ab1772 - Histone H3 - di methyl K9

ab1773 - Histone H3 - tri methyl K9

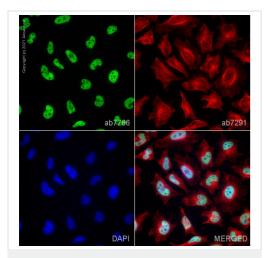
ab1780 - Histone H3 - mono methyl K27

ab1781 - Histone H3 - di methyl K27

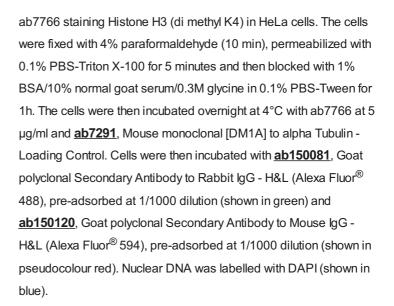
ab1782 - Histone H3 - tri methyl K27

ab7228 - Histone H3 - unmodified

ab7768 - Histone H3 - di methyl K4

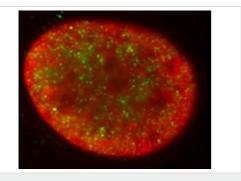


Immunocytochemistry/ Immunofluorescence - Anti-Histone H3 (di methyl K4) antibody - ChIP Grade (ab7766)



Also suitable in cells fixed with 100% methanol (5 min).

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



Immunocytochemistry/ Immunofluorescence - Anti-Histone H3 (di methyl K4) antibody - ChIP Grade (ab7766)

This image is courtesy of Kirk McManus in the lab of Michael Hendzel, Univeristy of Alberta

Dimethylated lysine 4 (green) is found in several hundred small nuclear foci that do not colocalize with condensed regions of chromatin (DAPI stained, red). The perinuclear and perinucleolar heterochromatin do not stain with this antibody.



Western blot - Anti-Histone H3 (di methyl K4) antibody - ChIP Grade (ab7766)

Anti-Histone H3 (di methyl K4) antibody - ChIP Grade (ab7766) at 1 μ g/ml + Calf Thymus Histone Preparation Nuclear Lysate (ab121) at 0.5 μ g

Secondary

Goat Anti-Rabbit IgG H&L (HRP) preadsorbed (ab97080) at 1/5000 dilution

Developed using the ECL technique.

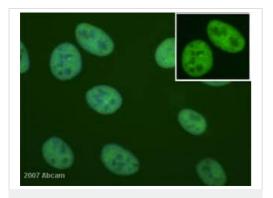
Performed under reducing conditions.

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

Additional bands at: 14 kDa. We are unsure as to the identity of

these extra bands.

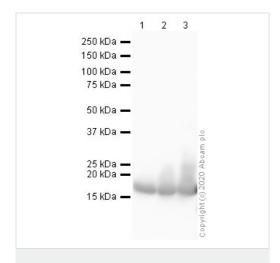
Exposure time: 4 minutes



Immunocytochemistry/ Immunofluorescence - Anti-Histone H3 (di methyl K4) antibody - ChIP Grade (ab7766)

This image is courtesy of an Abreview submitted by Dr Alexander Rapp

ab7766 at 1/200 staining human U2OS (osteosarcoma) cells by ICC/IF. The cells were paraformaldehyde fixed and then stained with the antibody for 1 hour. A Cy2 ® conjugated donkey anti-rabbit antibody was used as the secondary (green). The image shows uniformal staining of the whole nucleus, with several species found. The insert shows H3-di methyl K4 of tw2 cells only. DAPI nuclear staining is shown in blue.



Western blot - Anti-Histone H3 (di methyl K4) antibody - ChIP Grade (ab7766)

All lanes : Anti-Histone H3 (di methyl K4) antibody - ChIP Grade (ab7766) at 1 μ g/ml

Lane 1: HeLa nuclear lysate (triton enriched)

Lane 2: NIH 3T3 nuclear lysate (triton enriched)

Lane 3: PC12 nuclear lysate (triton enriched

Lysates/proteins at 20 µg per lane.

Secondary

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

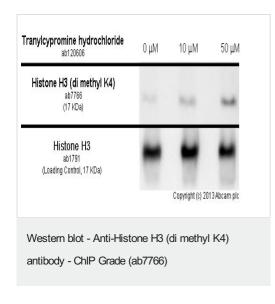
(HRP) at 1/50000 dilution

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

Blocking buffer: 2% BSA

Gel type: MES

Exposure Time: 1 minute



MCF7 cells were incubated at 37°C for 24h with vehicle control (0 μ M) and different concentrations of tranylcypromine hydrochloride (ab120606). Increased expression of Histone 3 K4 di-methyl (ab7766) in MCF7 cells correlates with an increase in tranylcypromine hydrochloride concentration, as described in literature.

Nuclear extracts were prepared with RIPA buffer (containing protease inhibitors and sodium orthovanadate), 10 μ g of each were loaded on the gel and the WB was run under reducing conditions. After transfer the membrane was blocked for an hour using 5% BSA before being incubated with ab7766 at 1 μ g /ml and ab1791 at 1 μ g /ml overnight at 4°C. Antibody binding was detected using an antirabbit HRP secondary antibody (ab97051) at 1/10000 dilution and visualized using ECL development solution.

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