

Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade ab45166

リコンビナント **RabMAb**

★★★★★ **1 Abreviews** **21 References** 画像数 **13**

製品の概要

製品名	Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade
製品の詳細	Rabbit monoclonal [EP1002Y] to Histone H4 (acetyl K8) - ChIP Grade
由来種	Rabbit
アプリケーション	適用あり: Flow Cyt (Intra), ChIP, ChIP-sequencing, WB, IHC-P, ICC/IF, IP
種交差性	交差種: Mouse, Rat, Human
免疫原	Synthetic peptide within Human Histone H4 aa 1-100 (N terminal) (acetyl K8). The exact sequence is proprietary. Database link: <u>P62805</u>
ポジティブ・コントロール	WB: HeLa whole cell lysate +TSA, C6 cell lysate, C6 cell + TSA lysate, NIH/3T3 +TSA whole cell lysate. IHC-P: Human normal colon FFPE tissue sections, mouse kidney paraffin-embedded tissue sections, rat kidney paraffin-embedded tissue sections. ICC/IF: C6 + TSA lysates. ChIP: Chromatin prepared from HeLa cells ChIP-Seq: HeLa Cells
特記事項	This product is a recombinant monoclonal antibody, which offers several advantages including: <ul style="list-style-type: none"> - High batch-to-batch consistency and reproducibility - Improved sensitivity and specificity - Long-term security of supply - Animal-free production For more information <u>see here</u> . Our RabMAb [®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <u>RabMAb[®] patents</u> .

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. Avoid freeze / thaw cycle.
バッファー	pH: 7.20 Preservative: 0.01% Sodium azide Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.17% BSA

精製度	Protein A purified
ポリ/モノ	モノクローナル
クローン名	EP1002Y
アイソタイプ	IgG

アプリケーション

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ab45166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確認ください。

アプリケーション	Abreviews	特記事項
Flow Cyt (Intra)		Use at an assay dependent concentration.
ChIP	★★★★★ (1)	Use 2 µg for 25 µg of chromatin.
ChIP-sequencing		Use 4µg for 10 ⁷ cells.
WB		1/5000 - 1/10000. Predicted molecular weight: 11 kDa.
IHC-P		1/250 - 1/2500. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. See <u>IHC antigen retrieval protocols</u> .
ICC/IF		1/150 - 1/500.
IP		1/20 - 1/50.

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 H4 family.
翻訳後修飾	<p>Acetylation at Lys-6 (H4K5ac), Lys-9 (H4K8ac), Lys-13 (H4K12ac) and Lys-17 (H4K16ac) occurs in coding regions of the genome but not in heterochromatin.</p> <p>Citrullination at Arg-4 (H4R3ci) by PAD4 impairs methylation.</p> <p>Monomethylation and asymmetric dimethylation at Arg-4 (H4R3me1 and H4R3me2a, respectively) by PRMT1 favors acetylation at Lys-9 (H4K8ac) and Lys-13 (H4K12ac).</p> <p>Demethylation is performed by JMJD6. Symmetric dimethylation on Arg-4 (H4R3me2s) by the PRDM1/PRMT5 complex may play a crucial role in the germ-cell lineage.</p> <p>Monomethylated, dimethylated or trimethylated at Lys-21 (H4K20me1, H4K20me2, H4K20me3). Monomethylation is performed by SET8. Trimethylation is performed by SUV420H1 and SUV420H2 and induces gene silencing.</p> <p>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</p>

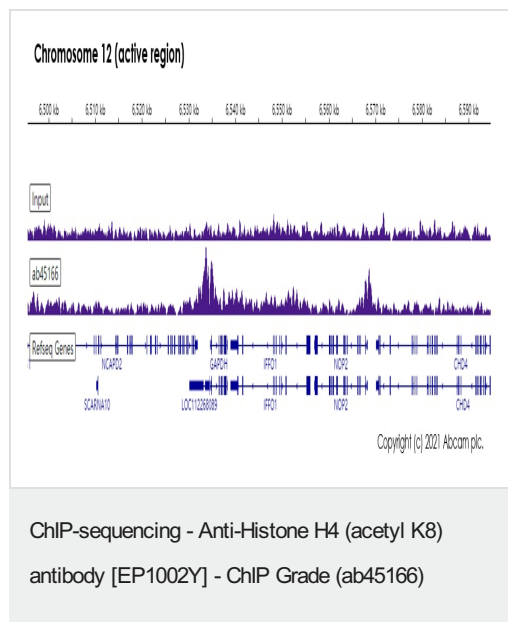
proteins. Monoubiquitinated at Lys-92 of histone H4 (H4K91ub1) in response to DNA damage. The exact role of H4K91ub1 in DNA damage response is still unclear but it may function as a licensing signal for additional histone H4 post-translational modifications such as H4 Lys-21 methylation (H4K20me).

Sumoylated, which is associated with transcriptional repression.

細胞内局在

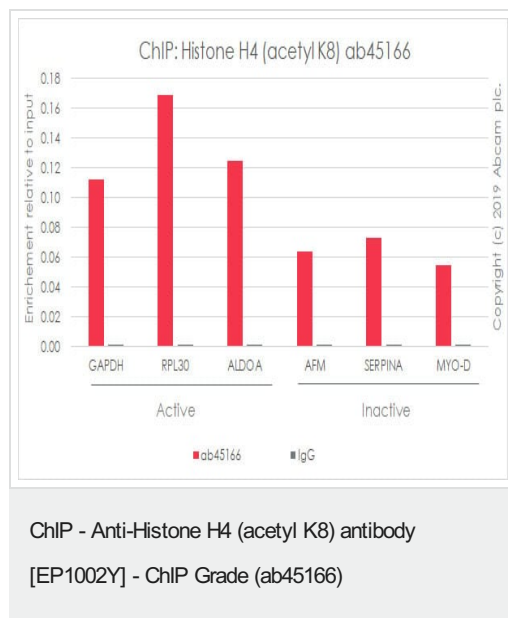
Nucleus. Chromosome.

画像

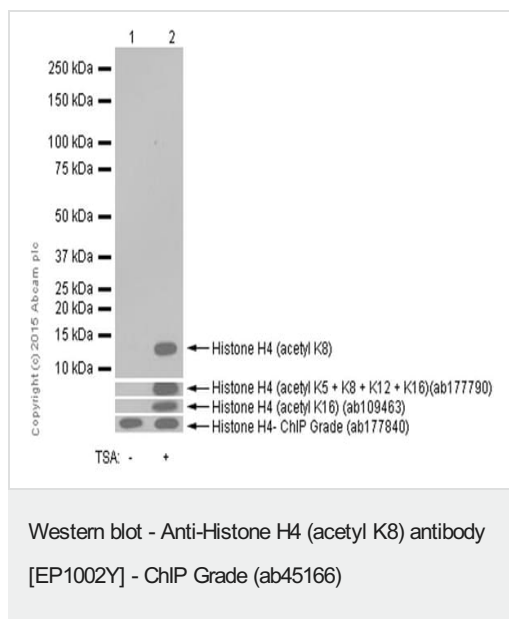


Chromatin was prepared from HeLa cells. Cells were fixed with 1% formaldehyde for 10 minutes. ChIP was performed with 10^7 HeLa cells and 4 μ g of ab45166 [EP1002Y]. ChIP DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 30 million reads.

Additional screenshots of mapped reads can be downloaded [here](#).



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 25 μ g of chromatin, 2 μ g of ab45166 (red), and 20 μ l of Protein A/G sepharose beads. No antibody was added to the beads control (grey). 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 H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166) at 1/5000 dilution

Lane 1 : Untreated HeLa (human cervix adenocarcinoma) whole cell lysate

Lane 2 : HeLa (human cervix adenocarcinoma) treated with Trichostatin A whole cell lysate

Lysates/proteins at 10 µg per lane.

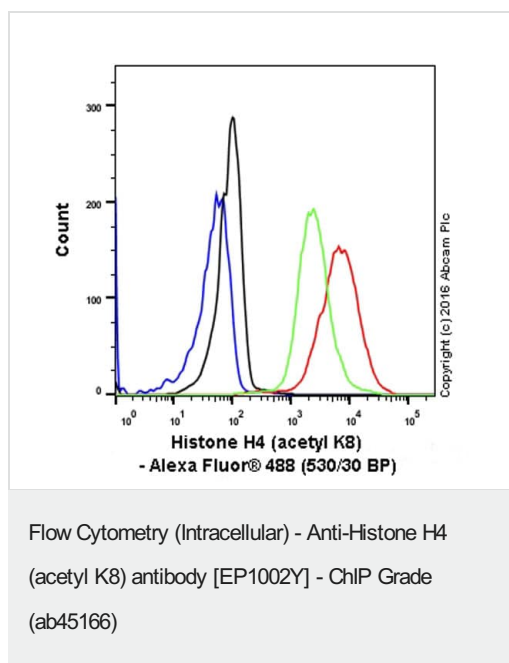
Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/20000 dilution

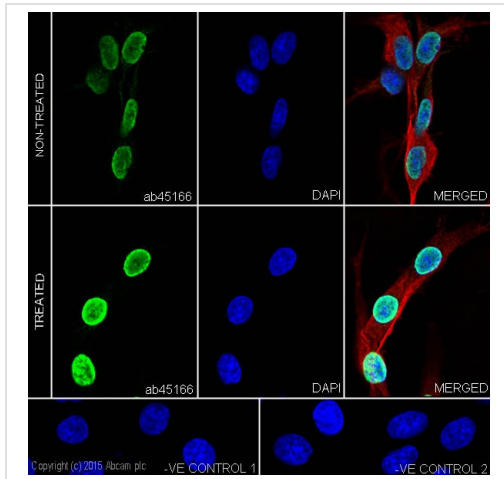
Predicted band size: 11 kDa

Observed band size: 11 kDa

Blocking and diluting buffer 5% NFDM/TBST

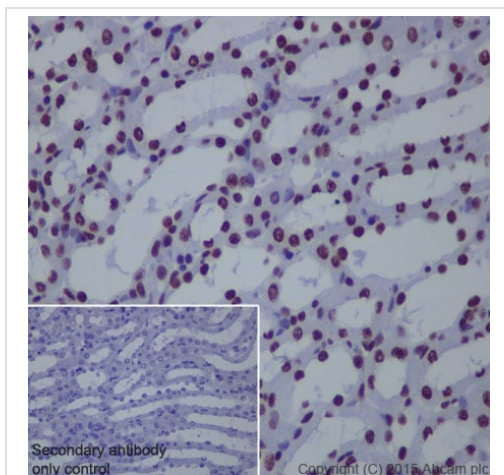


Intracellular Flow Cytometry analysis of HeLa (human cervix adenocarcinoma) treated (Red)/untreated (Green) with 500ng/ml Trichostatin A for 4 hours with purified ab45166 at 1/20 dilution. The secondary antibody was Goat anti rabbit IgG (Alexa Fluor® 488) at 1/2000 dilution. A Rabbit monoclonal IgG (Black) was used as the isotype control and cells without incubation with primary antibody and secondary antibody (Blue) were used as unlabeled control.



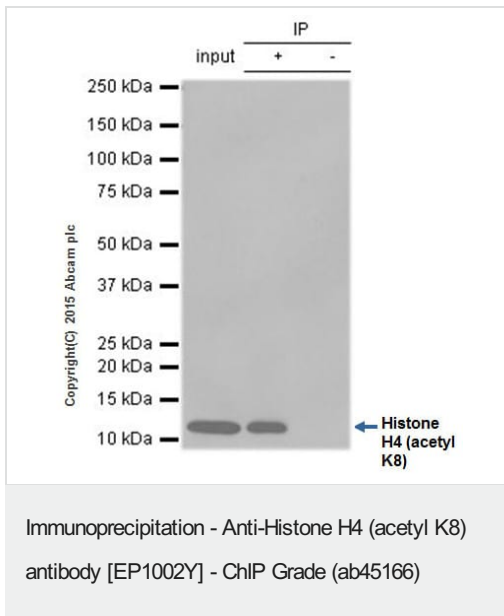
Immunocytochemistry/ Immunofluorescence - Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166)

Immunocytochemistry/immunofluorescence staining of 4% paraformaldehyde fixed; 0.1% triton X 100 permeabilized C6 (rat glioma) cells (non-treated-top panels) and (C6 + TSA(500ng/ml, 4hr)-middle panels) with purified ab45166 at dilution of 1/150. The secondary antibody used was Alexa Fluor® 488; goat anti-rabbit IgG (**ab150077**) at a dilution of 1/1000. Nucleus was counter-stained with DAPI (blue). **ab7291**, a mouse anti-tubulin antibody (1/1000) was used to stain tubulin along with **ab150120** (AlexaFluor®594 goat anti-mouse secondary, 1/1000) shown in the top right and middle right hand panels. The negative controls are shown in the bottom two panels- for negative control 1 rabbit primary antibody and anti-mouse secondary antibody (**ab150120**) was used. For negative control 2 mouse primary antibody (**ab7291**) and anti-rabbit secondary antibody (**ab150077**) was used.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166)

Immunohistochemical staining of paraffin-embedded mouse kidney sections labelling Histone H4 (acetyl K8) with purified ab45166 at dilution of 1:2500. The secondary antibody used was **ab97051**; a goat anti-rabbit IgG H&L (HRP) at dilution of 1/500. The sample was counter-stained with hematoxylin. Antigen retrieval was performed using EDTA Buffer; pH 9.0. PBS was used instead of the primary antibody as the negative control and is shown in the inset.



ab45166 (purified) at 1/20 immunoprecipitating Histone H4 (acetyl K8) in HeLa treated with Trichostatin A whole cell lysate.

Lane 1 (input): HeLa treated with Trichostatin A whole cell lysate (10 µg)

Lane 2 (+): ab45166 + HeLa treated with Trichostatin A whole cell lysate.

Lane 3 (-): Rabbit monoclonal IgG (**ab172730**) instead of ab45166 in HeLa treated with Trichostatin A whole cell lysate.

For western blotting, **ab131366** VeriBlot for IP Detection Reagent (HRP) was used for detection (1/10000).

Blocking buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM /TBST.

Lanes 1-2 : Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166) at 1/20 dilution

Lane 3 : Rabbit IgG, monoclonal [EPR25A] - Isotype Control (**ab172730**) at 1/20 dilution

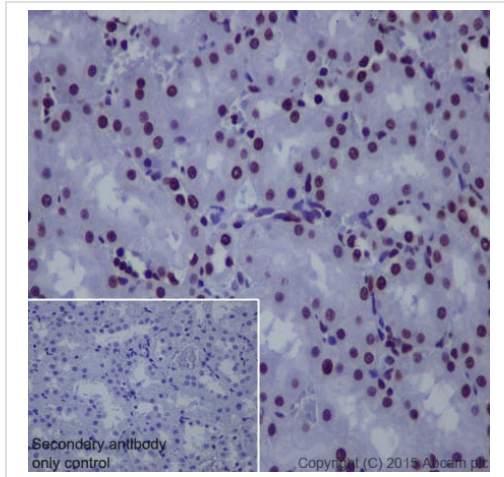
Lane 1 : HeLa (human cervix adenocarcinoma) treated with Trichostatin A whole cell lysate at 10 µg

Lanes 2-3 : HeLa (human cervix adenocarcinoma) treated with Trichostatin A whole cell lysate

Secondary

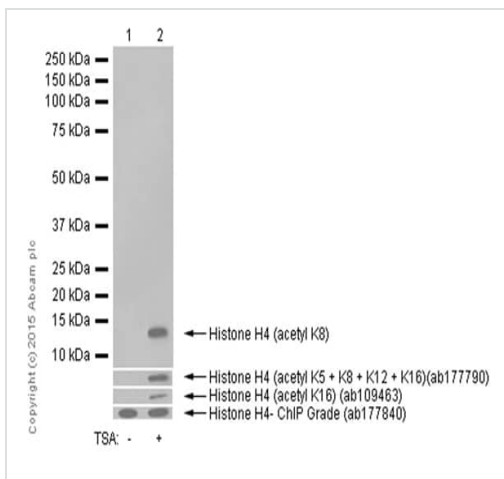
All lanes : VeriBlot for IP Detection Reagent (HRP) (**ab131366**) at 1/10000 dilution

Observed band size: 11 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166)

Immunohistochemical staining of paraffin-embedded rat kidney sections labelling Histone H4 (acetyl K8) with purified ab45166 at dilution of 1:2500. The secondary antibody used was **ab97051**; a goat anti-rabbit IgG H&L (HRP) at dilution of 1/500. The sample was counter-stained with hematoxylin. Antigen retrieval was performed using EDTA Buffer; pH 9.0. PBS was used instead of the primary antibody as the negative control and is shown in the inset.



Western blot - Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166)

All lanes : Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166) at 1/5000 dilution

Lane 1 : Untreated C6 (rat glioma) whole cell lysate

Lane 2 : C6 (rat glioma) treated with Trichostatin A whole cell lysate

Lysates/proteins at 10 µg per lane.

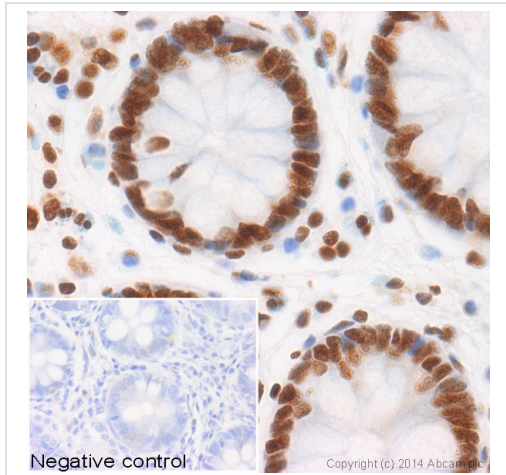
Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) (**ab97051**) at 1/20000 dilution

Predicted band size: 11 kDa

Observed band size: 11 kDa

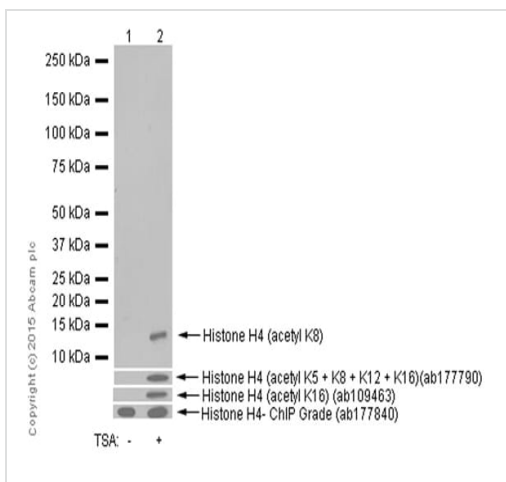
Blocking and diluting buffer 5% NFDm/TBST



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166)

Immunohistochemical analysis of formalin fixed paraffin embedded human colon tissue sections labelling Histone H4 (acetyl K8) with unpurified ab45166 at dilution of 1/200.

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.



Western blot - Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166)

All lanes : Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166) at 1/5000 dilution

Lane 1 : Untreated NIH/3T3 (mouse embryo) whole cell lysate

Lane 2 : NIH/3T3 (mouse embryo) treated with Trichostatin A whole cell lysate

Lysates/proteins at 10 µg per lane.

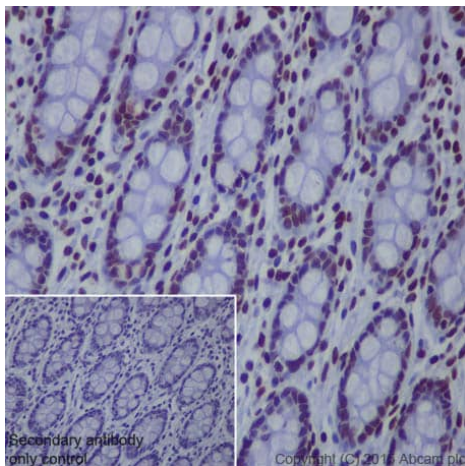
Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/20000 dilution

Predicted band size: 11 kDa

Observed band size: 11 kDa

Blocking and diluting buffer 5% NFDm/TBST



Immunohistochemical staining of paraffin-embedded human colon sections labelling Histone H4 (acetyl K8) with purified ab45166 at dilution of 1:2500. The secondary antibody used was **ab97051**; a goat anti-rabbit IgG H&L (HRP) at dilution of 1/500. The sample was counter-stained with hematoxylin. Antigen retrieval was performed using EDTA Buffer; pH 9.0. PBS was used instead of the primary antibody as the negative control and is shown in the inset.

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166)

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Anti-Histone H4 (acetyl K8) antibody [EP1002Y] - ChIP Grade (ab45166)

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