

## **H4K8ac Antibody**

Purified Rabbit Polyclonal Antibody Catalog # ABV11636

## **Specification**

#### **H4K8ac Antibody - Product Information**

Application
Primary Accession
Host
Clonality
Isotype
Calculated MW

DB, WB, E
P68431
Rabbit
Polyclonal
Rabbit IgG
15404

#### **H4K8ac Antibody - Additional Information**

**Gene ID** 8350;8351;8352;8353;8354;8355; 8356;8357;8358;8968

## Other Names Histone H4

Target/Specificity H4K8ac

### **Formulation**

In PBS with 0.05% (W/V) sodium azide.

## Handling

The antibody solution should be gently mixed before use.

## **Background Descriptions**

#### **Precautions**

H4K8ac Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

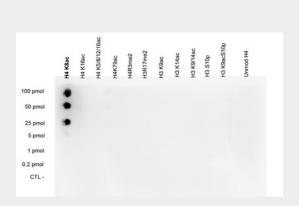
#### **H4K8ac Antibody - Protein Information**

Name H3C1 (<u>HGNC:4766</u>)

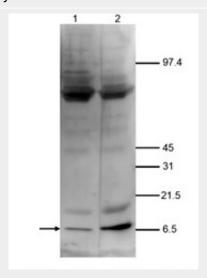
Synonyms H3FA, HIST1H3A

### **Function**

Core component of nucleosome.
Nucleosomes wrap and compact DNA into chromatin, limiting DNA accessibility to the



A Dot Blot analysis was performed to test the cross reactivity of the antibody with peptides containing other modifications of histone H4 and H3 and an unmodified H4 sequence, 100 to 0.2 pmol of the peptide containing the respective histone modification were spotted on a membrane. The Fig shows a high specificity of the antibody for the modification of interest.



Hela cells (15ug) were analysed by WB blot using the H4K8ac antibody.



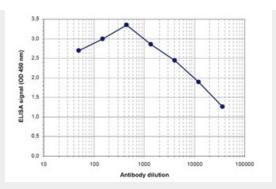
cellular machineries which require DNA as a template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication and chromosomal stability. DNA accessibility is regulated via a complex set of post-translational modifications of histones, also called histone code, and nucleosome remodeling.

**Cellular Location**Nucleus, Chromosome.

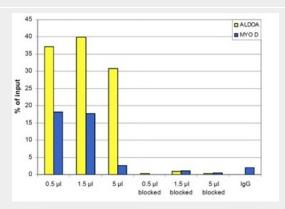
# **H4K8ac Antibody - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- <u>Immunoprecipitation</u>
- Flow Cytomety
- Cell Culture



To determine the titer, an ELISA was performed using a serial dilution of the antibody. The antigen used was a peptide containing the histone modification of interest. By plotting the absorbance against the antibody dilution the titer of the antibody was estimated to be 1:17500.



ChIP assays were performed using human osteosarcoma(U2OS) cells and the antibody and optimized PCR primer sets for qPCR. A titration of the antibody consisting of 2, 5, 10 and 15ul per ChIP experiment was analysed. IgG (5ug/IP) was used as negative control. The Fig shows the recovery, expressed as a %of input (the relative amount of IP DNA compared to input DNA and qPCR analysis). qPCR was performed with primers for the ALDOA promoter (fructose-diphosphate aldolase A) and for the coding region of the myogenic differentiation gene (MYOD), a gene that is inactive at normal conditions.

#### **H4K8ac Antibody - Background**

Histones are the main constituents of the protein part of chromosomes of eukaryotic cells. They are rich in the amino acids arginine and lysine and have been greatly conserved during evolution. Histones pack the DNA into tight masses of chromatin. Histone tails





abcepta

undergo numerous post-translational modifications, which either directly or indirectly alter chromatin structure to facilitate transcriptional activation or repression or other nuclear processes. In addition to the genetic code, combinations of the different histone modifications reveal the so-called "histone code". Histone methylation and demethylation is dynamically regulated by respectively histone methyl transferases and histone demethylases. Acetylation of histone H4 is associated with active gene transcription.