10320 Camino Santa Fe, Suite G
San Diego, CA 92121
Tel: 858.875.1900 Fax: 858.875.1999

H3K27me3S28p polyclonal antibody

Rabbit Polyclonal Antibody
Catalog # ABV11345

Specification

H3K27me3S28p polyclonal antibody - Product Information

Application	**CHIP, DB, E, WB**
Primary Accession	P68431
Host	**Rabbit**
Clonality	**Polyclonal**
Isotype	**Rabbit IgG**
Calculated MW	**15404**

H3K27me3S28p polyclonal antibody - Additional Information

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

Positive Control	**Western blot: HeLa cells, IF: HeLa cells, ELISA: Antigen, ChIP: HeLa cells, Dot blot: Histone Peptides.**
Application & Usage	**IF: 1:200, WB: 1:250 – 1:500, ELISA: 1:100 – 1:500, Dot Blot: 1:20,000, ChIP: 1 µl/ChIP, IP: 5 µl/IP.**

Other Names
Histone H3

Target/Specificity
H3K27me3S28p

Antibody Form
Liquid

Appearance
Colorless liquid

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

Handling
The antibody solution should be gently mixed before use.

Reconstitution & Storage
-20 °C

Background Descriptions

Precautions
H3K27me3S28p polyclonal antibody is for research use only and not for use in diagnostic or therapeutic procedures.

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H3K27me3S28p polyclonal antibody - Protein Information

Name H3C1 (HGNC:4766)

Synonyms H3FA, HIST1H3A

Function
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.

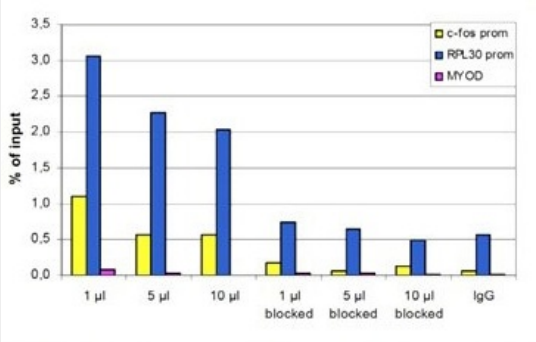
Cellular Location
Nucleus. Chromosome.

H3K27me3S28p polyclonal antibody - Protocols

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

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

H3K27me3S28p polyclonal antibody - Images

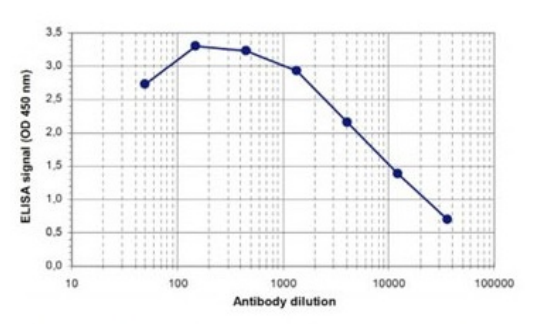


ChIP assays were performed using HeLa cells and the antibody and optimized PCR primer sets for qPCR. A titration of the antibody consisting of 2, 5, 10 and 15 µl per ChIP experiment was analysed. IgG (5 µg/IP) was used as negative control. The Fig shows the recovery, expressed as a % of input (the relative amount of IP DNA compared to input DNA after qPCR analysis).

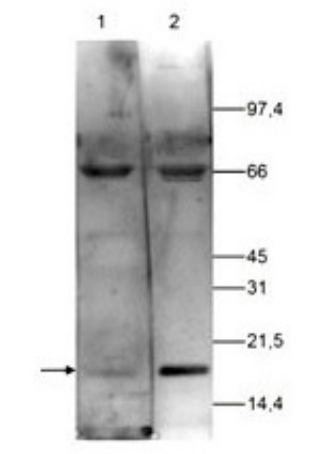
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A Dot Blot analysis was performed to test the cross reactivity of the antibody with peptides containing other modifications of histone H4 and H3 and unmodified H3 sequences. 100 to 0.2 pmol of the peptide containing the respective histone modification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



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 1:8300.



HeLa cells extracts (15 µg) were analysed by WB blot using the antibody.

H3K27me3S28p polyclonal 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. Histone tails undergo numerous post-translational modifications, which either directly or indirectly alter chromatin structure to facilitate transcriptional activation or repression or other nuclear processes. Histone methylation and demethylation is dynamically regulated by respectively histone methyl transferases and histone demethylases. Phosphorylation of H3 on serine 28 is increased during mitosis.