

# Histone H3K36me2 (H3K36 Dimethyl) Polyclonal Antibody

(Catalog # A-4041)

# **Background**

Histones are basic nuclear proteins that are responsible for the nucleosome structure of the chromosomal fiber in eukaryotes. Nucleosomes consist of approximately 146 bp of DNA wrapped around a histone octamer composed of pairs of each of the four core histones (H2A, H2B, H3, and H4). The chromatin fiber is further compacted through the interaction of a linker histone, H1, with the DNA between the nucleosomes to form higher order chromatin structures. This gene is intronless and encodes a replication-dependent histone that is a member of the histone H3 family. Transcripts from this gene lack polyA tails; instead, they contain a palindromic termination element. This gene is located separately from the other H3 genes that are in the histone gene cluster on chromosome 6p22-p21.3.

## **Description**

Histone H3K36me2 (H3K36 Dimethyl) Polyclonal Antibody. Unconjugated. Raised in: Rabbit.

#### **Formulation**

Buffer: PBS with 0.05% proclin300,50% glycerol, pH7.3.

# **Specificity**

Broad Range, Mouse, Rat, Human

# Isotype

**IgG** 

#### **Uniprot ID**

Q16695/P68431

## **Purification**

Affinity Purified

#### **Immunogen**

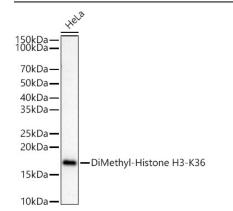
A synthetic dimethylated peptide around K36 of human histone H3 (NP\_003520.1)

## Storage

Shipped at 4°C. Store at -20°C. Avoid multiple freeze/thaw cycles.

#### Application

WB, IHC, IF/ICC, ChIP, ChIPseq, ELISA; Recommended dilution: WB 1:500 - 1:1000, IHC 1:50 - 1:200, IF/ICC 1:50 - 1:200, ChIP 5μg antibody for 5μg-10μg of Chromatin, CHIPseq 1:20 - 1:100, ELISA - recommended starting concentration is 1 μg/mL. Please optimize the concentration based on your specific assay requirements

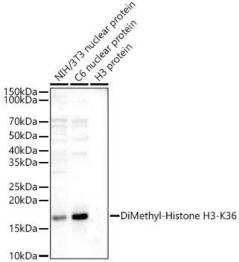


Western blot analysis of lysates from HeLa cells, using Histone H3K36me2 (H3K36 Dimethyl) Polyclonal Antibody at 1:600 dilution. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Exposure time: 180s.



Western blot analysis of various lysates, using Histone H3K36me2

(H3K36 Dimethyl) Polyclonal Antibody at 1:600 dilution.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) at

1:10000 dilution.

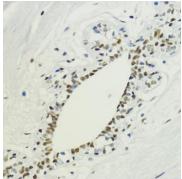
Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

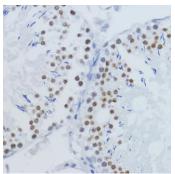
Exposure time: 180s.

	H3R2		H3K4		H3R8		Н3К9		H3R17		H3R26	
	tong	50n9	tong	50n9	tong	50n9	40ng	50ng	40ng	50n9	tong	50ng
me0	0	0	0	0	0	0	0	0	0	0	0	0
me1	0	0	0	0	0	0	0	0	0	0	0	0
me2/ me2a	0	0	0	0	0	0	0	0	0	0	0	0
me3/ me2s	0	0	0	0	0	0	0	0	0	0	0	0
	H3K27		H3K36		H3K56		H3K79		H4R3		H4K20	
me0	0	0	0	0	0	0	0	0	0	0	0	0
me1	0	0	0	0	0	0	0	0	0	0	0	0
me2/ me2a	0	0	0	•	0	0	0	0	0	0	0	0
me3/	0	0	0	0	0	0	0	0	0	0	0	0

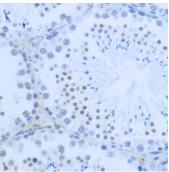
Dot-blot analysis of all sorts of methylation peptides using Histone H3K36me2 (H3K36 Dimethyl) Polyclonal Antibody.



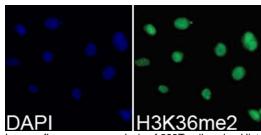
Immunohistochemistry of paraffin-embedded human breast using Histone H3K36me2 (H3K36 Dimethyl) Polyclonal Antibody at dilution of 1:200 (40x lens). Microwave antigen retrieval performed with 0.01M PBS Buffer (pH 7.2) prior to IHC staining.



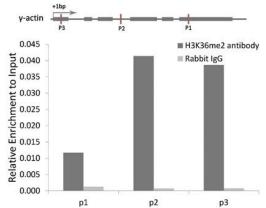
Immunohistochemistry of paraffin-embedded rat testis using Histone H3K36me2 (H3K36 Dimethyl) Polyclonal Antibody at dilution of 1:200 (40x lens). Microwave antigen retrieval performed with 0.01M PBS Buffer (pH 7.2) prior to IHC staining.



Immunohistochemistry of paraffin-embedded mouse testis using Histone H3K36me2 (H3K36 Dimethyl) Polyclonal Antibody at dilution of 1:200 (40x lens). Microwave antigen retrieval performed with 0.01M PBS Buffer (pH 7.2) prior to IHC staining.



Immunofluorescence analysis of 293T cells using Histone H3K36me2 (H3K36 Dimethyl) Polyclonal Antibody. Blue: DAPI for nuclear staining.



Chromatin immunoprecipitation analysis of y-actin gene from 293 cell line, using Histone H3K36me2 (H3K36 Dimethyl) Polyclonal Antibody and rabbit IgG. P1, P2 and P3 were probes located on γ-actin gene as the schematic diagram illustrated. The amount of immunoprecipitated DNA was checked by quantitative PCR. Histogram was constructed by the ratios of the immunoprecipitated DNA to the input.