

# Resource Summary Report

Generated by [RRID](#) on Aug 4, 2026

## Rabbit Anti-Histone H3, dimethyl (Lys36) Monoclonal Antibody, Unconjugated, Clone C75H12

RRID:AB\_1030983

Type: Antibody

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### Proper Citation

Cell Signaling Technology Cat# 2901, RRID:AB\_1030983

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### Antibody Information

**URL:** [http://antibodyregistry.org/AB\\_1030983](http://antibodyregistry.org/AB_1030983)

**Proper Citation:** Cell Signaling Technology Cat# 2901, RRID:AB\_1030983

**Target Antigen:** Histone H3, dimethyl (Lys36)

**Host Organism:** rabbit

**Clonality:** monoclonal

**Comments:** Applications: W, IHC-P, IF-IC, F. Consolidation on 10/2018: AB\_10287336, AB\_1030983, AB\_10310188.

Info: Independent validation by the NYU Lagone was performed for: IHC. This antibody was found to have the following characteristics: Functional in human:TRUE, NonFunctional in human:FALSE, Functional in animal:FALSE, NonFunctional in animal:FALSE

**Antibody Name:** Rabbit Anti-Histone H3, dimethyl (Lys36) Monoclonal Antibody, Unconjugated, Clone C75H12

**Description:** This monoclonal targets Histone H3, dimethyl (Lys36)

**Target Organism:** monkey, rat, simian, mouse, human

**Clone ID:** Clone C75H12

**Antibody ID:** AB\_1030983

**Vendor:** Cell Signaling Technology

**Catalog Number:** 2901

**Record Creation Time:** 20250820T065454+0000

**Record Last Update:** 20250821T011157+0000

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## Ratings and Alerts

- Independent validation by the NYU Lagone was performed for: IHC. This antibody was found to have the following characteristics: Functional in human:TRUE, NonFunctional in human:FALSE, Functional in animal:FALSE, NonFunctional in animal:FALSE - NYU Langone's Center for Biospecimen Research and Development <https://med.nyu.edu/research/scientific-cores-shared-resources/center-biospecimen-research-development>

No alerts have been found for Rabbit Anti-Histone H3, dimethyl (Lys36) Monoclonal Antibody, Unconjugated, Clone C75H12.

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## Data and Source Information

**Source:** [Antibody Registry](#)

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## Usage and Citation Metrics

We found 18 mentions in open access literature.

**Listed below are recent publications.** The full list is available at [RRID](#) .

Gökbuget D, et al. (2025) KMT2C/KMT2D-dependent H3K4me1 mediates changes in DNA replication timing and origin activity during a cell fate transition. Cell reports, 44(2), 115272.

Moreno Rueda LY, et al. (2025) Single-cell analysis of neoplastic plasma cells identifies myeloma pathobiology mediators and potential targets. Cell reports. Medicine, 6(2), 101925.

Zhang Q, et al. (2024) EZH2/G9a interact to mediate drug resistance in non-small-cell lung cancer by regulating the SMAD4/ERK/c-Myc signaling axis. Cell reports, 43(2), 113714.

Liu H, et al. (2024) Enforced activation of the CREB/KDM2B axis prevents alcohol-induced embryonic developmental delay. Cell reports, 43(12), 115075.

Hamagami N, et al. (2023) NSD1 deposits histone H3 lysine 36 dimethylation to pattern non-CG DNA methylation in neurons. Molecular cell, 83(9), 1412.

Gong M, et al. (2023) Transcriptional and metabolic programs promote the expansion of follicular helper T cells in lupus-prone mice. iScience, 26(5), 106774.

Hamagami N, et al. (2023) NSD1 deposits histone H3 lysine 36 dimethylation to pattern non-CG DNA methylation in neurons. bioRxiv : the preprint server for biology.

Fang L, et al. (2023) Methionine restriction promotes cGAS activation and chromatin untethering through demethylation to enhance antitumor immunity. *Cancer cell*, 41(6), 1118.

Zheng Y, et al. (2023) Histone methylation mediated by NSD1 is required for the establishment and maintenance of neuronal identities. *Cell reports*, 42(12), 113496.

Xu Y, et al. (2022) The KRAS-G12D mutation induces metabolic vulnerability in B-cell acute lymphoblastic leukemia. *iScience*, 25(3), 103881.

Li Y, et al. (2022) Histone methylation antagonism drives tumor immune evasion in squamous cell carcinomas. *Molecular cell*, 82(20), 3901.

Harpaz N, et al. (2022) Single-cell epigenetic analysis reveals principles of chromatin states in H3.3-K27M gliomas. *Molecular cell*, 82(14), 2696.

Farhangdoost N, et al. (2021) Chromatin dysregulation associated with NSD1 mutation in head and neck squamous cell carcinoma. *Cell reports*, 34(8), 108769.

Yan R, et al. (2021) Decoding dynamic epigenetic landscapes in human oocytes using single-cell multi-omics sequencing. *Cell stem cell*, 28(9), 1641.

Chaouch A, et al. (2021) Histone H3.3 K27M and K36M mutations de-repress transposable elements through perturbation of antagonistic chromatin marks. *Molecular cell*, 81(23), 4876.

Harutyunyan AS, et al. (2020) H3K27M in Gliomas Causes a One-Step Decrease in H3K27 Methylation and Reduced Spreading within the Constraints of H3K36 Methylation. *Cell reports*, 33(7), 108390.

Streubel G, et al. (2018) The H3K36me2 Methyltransferase Nsd1 Demarcates PRC2-Mediated H3K27me2 and H3K27me3 Domains in Embryonic Stem Cells. *Molecular cell*, 70(2), 371.

Li Q, et al. (2018) Bisphenol A and Phthalates Modulate Peritoneal Macrophage Function in Female Mice Involving SYMD2-H3K36 Dimethylation. *Endocrinology*, 159(5), 2216.