Cell Signaling Store at -200 Mono-Methyl-Histone H3 (Lys36) (D9J1D) Rabbit mAb TECHNOLOGY® Orders: 877-616-CELL (2355) orders@cellsignal.com Support: 877-678-TECH (8324) Web: info@cellsignal.com cellsignal.com 3 Trask Lane | Danvers | Massachusetts | 01923 | USA For Research Use Only. Not for Use in Diagnostic Procedures. Applications: Reactivity: Sensitivity: MW (kDa): Source/Isotype: UniProt ID: Entrez-Gene Id: WB, IP, IF-IC, FC-FP, H M R Mk Endogenous 17 Rabbit IgG #P68431 8350 ChIP, ChIP-seq, C&R, C&T For optimal ChIP and ChIP-seq results, use 10 µl of antibody and 10 µg of chromatin (approximately 4 x Product Usage 10<sup>6</sup> cells) per IP. This antibody has been validated using SimpleChIP<sup>®</sup> Enzymatic Chromatin IP Kits. Information The CUT&RUN dilution was determined using CUT&RUN Assay Kit #86652. The CUT&Tag dilution was determined using CUT&Tag Assay Kit #77552. Application Dilution Western Blotting 1:1000 Immunoprecipitation 1:50 Immunofluorescence (Immunocytochemistry) 1:1600 Flow Cytometry (Fixed/Permeabilized) 1:400 Chromatin IP 1:50 Chromatin IP-seq 1:50 CUT&RUN 1.50CUT&Tag 1:50 Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than Storage 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody. For a carrier free (BSA and azide free) version of this product see product #76466. Specificity / Sensitivity Mono-Methyl-Histone H3 (Lys36) (D9J1D) Rabbit mAb recognizes endogenous levels of histone H3 only when mono-methylated at Lys36. The antibody does not cross-react with non-methylated, di-methylated, or tri-methylated Lys36. In addition, the antibody does not cross-react with mono-methylated histone H3 Lys4, Lys9, Lys27, or Lys79. Species predicted to Hamster, Bovine react based on 100% sequence homology: Source / Purification Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding mono-methylated Lys36 of human histone H3 protein. Background The nucleosome, made up of four core histone proteins (H2A, H2B, H3, and H4), is the primary building block of chromatin. Originally thought to function as a static scaffold for DNA packaging, histones have now been shown to be dynamic proteins, undergoing multiple types of post-translational modifications, including acetylation, phosphorylation, methylation, and ubiquitination (1). Histone methylation is a major determinant for the formation of active and inactive regions of the genome and is crucial for the proper programming of the genome during development (2,3). Arginine methylation of histones H3 (Arg2, 17, 26) and H4 (Arg3) promotes transcriptional activation and is mediated by a family of protein arginine methyltransferases (PRMTs), including the co-activators PRMT1 and CARM1 (PRMT4) (4). In contrast, a more diverse set of histone lysine methyltransferases has been identified, all but one of which contain a conserved catalytic SET domain originally identified in the Drosophila Su(var)3-9, Enhancer of zeste, and Trithorax proteins. Lysine methylation occurs primarily on histones H3 (Lys4, 9, 27, 36, 79) and H4 (Lys20) and has been implicated in both transcriptional activation and silencing (4). Methylation of these lysine

residues coordinates the recruitment of chromatin modifying enzymes containing methyl-lysine binding

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	modules such as chromodomains (HP1, PRC1), PHD fingers (BPTF, ING2), tudor domains (53BP1), and WD-40 domains (WDR5) (5-8). The discovery of histone demethylases, such as PADI4, LSD1, JMJD1, JMJD2, and JHDM1, has shown that methylation is a reversible epigenetic marker (9).
Background References	<ol> <li>Peterson, C.L. and Laniel, M.A. (2004) <i>Curr Biol</i> 14, R546-51.</li> <li>Kubicek, S. et al. (2006) <i>Ernst Schering Res Found Workshop</i>, 1-27.</li> <li>Lin, W. and Dent, S.Y. (2006) <i>Curr Opin Genet Dev</i> 16, 137-42.</li> <li>Lee, D.Y. et al. (2005) <i>Endocr Rev</i> 26, 147-70.</li> <li>Daniel, J.A. et al. (2005) <i>Cell Cycle</i> 4, 919-26.</li> <li>Shi, X. et al. (2006) <i>Nature</i> 442, 96-9.</li> <li>Wysocka, J. et al. (2006) <i>Nature</i> 442, 86-90.</li> <li>Wysocka, J. et al. (2005) <i>Cell</i> 121, 859-72.</li> <li>Trojer, P. and Reinberg, D. (2006) <i>Cell</i> 125, 213-7.</li> </ol>
Species Reactivity	Species reactivity is determined by testing in at least one approved application (e.g., western blot).
Western Blot Buffer	IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.
Applications Key	WB: Western Blotting IP: Immunoprecipitation IF-IC: Immunofluorescence (Immunocytochemistry) FC-FP: Flow Cytometry (Fixed/Permeabilized) ChIP: Chromatin IP ChIP-seq: Chromatin IP-seq C&R: CUT&RUN C&T: CUT&Tag
Cross-Reactivity Key	H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected
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