

Mono-Methyl-Histone H3 (Lys9) (D1P5R) Rabbit mAb



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Applications: W, IP, IF-IC, ChIP	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 17	Source/Isotype: Rabbit IqG	UniProt ID: #P68431	Entrez-Gene Id: 8350	
VV, 1P, 1P-1C, CHIP	H IVI K IVIK	Endogenous	17	Kabbit IgG	#100431	6550	
Product Usage Information		For optimal ChIP results, use 10 µl of antibody and 10 µg of chromatin (approximately 4 x 10 ⁶ cells) per IP. This antibody has been validated using SimpleChIP [®] Enzymatic Chromatin IP Kits.					
		Application				Dilution	
		Western Blotting				1:1000	
		Immunoprecipitation				1:100	
		Immunofluorescence Chromatin IP	e (Immunocytochem	iistry)		1:3200 1:50	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. <i>Do not aliquot the antibody.</i>					
		For a carrier free (BSA and azide free) version of this product see product #68406.					
Specificity/Sensitivity		Mono-Methyl-Histone H3 (Lys9) (D1P5R) Rabbit mAb recognizes endogenous levels of histone H3 protein only when mono-methylated at Lys9. This antibody does not cross-react with non-methylated, di-methylated, or tri-methylated Lys9. In addition, the antibody does not cross-react with other known mono-methylated, di-methylated, or tri-methylated lysines on histones H3, H4, H2A, or H2B.					
Species predicted to react based on 100% sequence homology		Xenopus, Zebrafish, F	Pig, S. cerevisiae, Ho	rse			
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding mono-methyl Lys9 of human histone H3 protein.					
block of chromatin. Origin now been shown to be dy modifications, including a methylation is a major der is crucial for the proper pr of histones H3 (Arg2, 17, 2 family of protein arginine (PRMT4) (4). In contrast, a but one of which contain a Su(var)3-9, Enhancer of ze H3 (Lys4, 9, 27, 36, 79) and silencing (4). Methylation enzymes containing meth (BPTF, ING2), tudor domai			originally thought to be dynamic proteins, ing acetylation, pho- or determinant for the er programming of 17, 26) and H4 (Arg nine methyltransfer ist, a more diverse s tain a conserved cat of zeste, and Trithou of and H4 (Lys20) and tion of these lysine methyl-lysine bindin omains (53BP1), and is PADI4, LSD1, JMJD	up of four core histone proteins (H2A, H2B, H3, and H4), is the primary building ginally thought to function as a static scaffold for DNA packaging, histones have dynamic proteins, undergoing multiple types of post-translational gracetylation, phosphorylation, methylation, and ubiquitination (1). Histone determinant for the formation of active and inactive regions of the genome and programming of the genome during development (2,3). Arginine methylation (7, 26) and H4 (Arg3) promotes transcriptional activation and is mediated by a ne methyltransferases (PRMTs), including the co-activators PRMT1 and CARM1 and conserved catalytic SET domain originally identified in the <i>Drosophila</i> zeste, and Trithorax proteins. Lysine methylation occurs primarily on histones and H4 (Lys20) and has been implicated in both transcriptional activation and of these lysine residues coordinates the recruitment of chromatin modifying ethyl-lysine binding modules such as chromodomains (HP1, PRC1), PHD fingers mains (53BP1), and WD-40 domains (WDR5) (5-8). The discovery of histone PADI4, LSD1, JMJD1, JMJD2, and JHDM1, has shown that methylation is a			
Background References		1. Peterson, C.L. and Laniel, M.A. (2004) <i>Curr Biol</i> 14, R546-51. 2. Kubicek, S. et al. (2006) <i>Ernst Schering Res Found Workshop</i> , 1-27. 3. Lin, W. and Dent, S.Y. (2006) <i>Curr Opin Genet Dev</i> 16, 137-42. 4. Lee, D.Y. et al. (2005) <i>Endocr Rev</i> 26, 147-70. 5. Daniel, J.A. et al. (2005) <i>Cell Cycle</i> 4, 919-26. 6. Shi, X. et al. (2006) <i>Nature</i> 442, 96-9. 7. Wysocka, J. et al. (2006) <i>Nature</i> 442, 86-90. 8. Wysocka, J. et al. (2005) <i>Cell</i> 121, 859-72. 9. Trojer, P. and Reinberg, D. (2006) <i>Cell</i> 125, 213-7.					

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 nonfat

dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key W: Western Blotting IP: Immunoprecipitation IF-IC: Immunofluorescence (Immunocytochemistry)

ChIP: Chromatin IP

Cross-Reactivity Key H: Human M: Mouse R: Rat Mk: Monkey

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