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Mono-Methyl-Histone H3 (Lys36) (D9J1D) Rabbit mAb



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Applications: W, IP, IF-IC, FC-FP, ChIP, ChIP-seq, C&R, C&T	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 17	Source/Isotype: Rabbit IgG	UniProt ID: #P68431	Entrez-Gene Id: 8350	
Product Usage Information		For optimal ChIP and ChIP-seq 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.					
		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	
		Immunoprecipitatior	า			1:50	
		Immunofluorescence	e (Immunocytochem	nistry)		1:1600	
		Flow Cytometry (Fixe	-	<u>,</u>		1:400	
		Chromatin IP				1:50	
		Chromatin IP-seq				1:50	
		CUT&RUN				1:50	
		CUT&Tag				1:50	
		condiag				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. Do not aliquot the antibody.					
		For a carrier free (BS/	A and azide free) vei	rsion of this product see	product #76466.		
Specificity/Sensiti	vity	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 based on 100% sec homology		Hamster, Bovine					
Source / Purificati	on	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 <i>Drosophila</i> 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 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	 Peterson, C.L. and Laniel, M.A. (2004) <i>Curr Biol</i> 14, R546-51. Kubicek, S. et al. (2006) <i>Ernst Schering Res Found Workshop</i>, 1-27. Lin, W. and Dent, S.Y. (2006) <i>Curr Opin Genet Dev</i> 16, 137-42. Lee, D.Y. et al. (2005) <i>Endocr Rev</i> 26, 147-70. Daniel, J.A. et al. (2005) <i>Cell Cycle</i> 4, 919-26. Shi, X. et al. (2006) <i>Nature</i> 442, 96-9. Wysocka, J. et al. (2006) <i>Nature</i> 442, 86-90. Wysocka, J. et al. (2005) <i>Cell</i> 121, 859-72. 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 BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.		
Applications Key	W: 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 Mk: Monkey		
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