

## PCAF (C14G9) Rabbit mAb



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<b>Applications:</b> V, IP, ChIP, ChIP-seq	<b>Reactivity:</b> H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 93	<b>Source/Isotype:</b> Rabbit IgG	
Product Usage Information		For optimal ChIP and ChIP-seq results, use 20 $\mu$ l of antibody and 10 $\mu$ g of chromatin (approximately 4 10 <sup>6</sup> cells) per IP. This antibody has been validated using SimpleChIP® Enzymatic Chromatin IP Kits.			
		Application			Dilution
		Western Blotting			1:1000
		Immunoprecipitation			1:100
		Chromatin IP			1:25
		Chromatin IP-seq			1:25
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.			
Specificity/Sensitivity		PCAF (C14G9) Rabbit mAb detects endogenous levels of total PCAF protein. The antibody does not cross-react with the related GCN5L2 protein.			
Species predicted to react based on 100% sequence homology		Bovine, Horse			
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to the amino terminus of human PCAF protein.			
Background		p300/CBP-associated factor (PCAF), also known as lysine acetyl-transferase 2B (KAT2B), is a transcriptional adaptor protein and histone acetyl-transferase (HAT) that functions as the catalytic subunit of the PCAF transcriptional co-activator complex (1). PCAF is 73% homologous to GCN5L2, another HAT protein found in similar complexes (1,2). Like GCN5L2, PCAF acetylates histone H3 on Lys14 and histone H4 on Lys8, both of which contribute to gene activation by modulating chromatin structure and recruiting additional co-activator proteins that contain acetyl-lysine binding bromodomains (3). PCAF also acetylates non-histone proteins including transcriptional activators (p53, E2F1, MyoD), general transcription factors (TFIIE $\beta$ and TFIIF) and architectural DNA binding proteins (HMGA1 and HMG17) (4-10). Acetylation of these proteins regulates their nuclear localization, protein stability, DNA binding, and co-activator association.			
Background References		<ol> <li>Nagy, Z. and Tora, L. (2007) Oncogene 26, 5341-57.</li> <li>Yang, X.J. et al. (1996) Nature 382, 319-24.</li> <li>Schiltz, R.L. et al. (1999) J Biol Chem 274, 1189-92.</li> <li>Bannister, A.J. and Miska, E.A. (2000) Cell Mol Life Sci 57, 1184-92.</li> <li>Liu, L. et al. (1999) Mol Cell Biol 19, 1202-9.</li> <li>Martínez-Balbás, M.A. et al. (2000) EMBO J 19, 662-71.</li> <li>Sartorelli, V. et al. (1999) Mol Cell 4, 725-34.</li> <li>Imhof, A. et al. (1997) Curr Biol 7, 689-92.</li> <li>Munshi, N. et al. (1998) Mol Cell 2, 457-67.</li> <li>Herrera, J.E. et al. (1999) Mol Cell Biol 19, 3466-73.</li> </ol>			
Species Reactivit	v		· · · · ·		lication (e.g., western blot).

Species Reactivity

**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 ChIP: Chromatin IP ChIP-seq: Chromatin IP-seq

**Cross-Reactivity Key** 

H: Human M: Mouse R: Rat Mk: Monkey

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