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## Acetyl-Histone H4 (Lys12) Antibody



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Applications:Reactivity:Sensitivity:MW (kDa):Source:UniProt ID:Entrez-Gene Id:WB, IF-ICH M R MkEndogenous11Rabbit#P628058359

Product Usage<br/>InformationApplicationDilutionWestern Blotting1:1000Immunofluorescence (Immunocytochemistry)1:400

Storage Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.

Specificity / Sensitivity

Acetyl-Histone H4 (Lys12) Antibody detects endogenous levels of histone H4 only when acetylated at Lys12. The antibody does not cross-react with other acetylated histones.

Source / Purification Polyclonal antibodies are produced by immunizing animals with a synthetic acetylated peptide

corresponding to residues surrounding Lys12 of human histone H4. Antibodies are purified by protein A

and peptide affinity chromatography.

**Background** Modulation of chromatin structure plays an important role in the regulation of transcription in eukaryotes.

The nucleosome, made up of DNA wound around eight core histone proteins (two each of H2A, H2B, H3, and H4), is the primary building block of chromatin (1). The amino-terminal tails of core histones undergo various posttranslational modifications, including acetylation, phosphorylation, methylation, and ubiquitination (2-5). These modifications occur in response to various stimuli and have a direct effect on the accessibility of chromatin to transcription factors and, therefore, gene expression (6). In most species, histone H2B is primarily acetylated at Lys5, 12, 15, and 20 (4,7). Histone H3 is primarily acetylated at Lys9, 14, 18, 23, 27, and 56. Acetylation of H3 at Lys9 appears to have a dominant role in histone deposition and chromatin assembly in some organisms (2,3). Phosphorylation at Ser10, Ser28, and Thr11 of histone H3 is tightly correlated with chromosome condensation during both mitosis and meiosis (8-10). Phosphorylation at Thr3 of histone H3 is highly conserved among many species and is catalyzed by the kinase haspin. Immunostaining with phospho-specific antibodies in mammalian cells reveals mitotic phosphorylation at

Thr3 of H3 in prophase and its dephosphorylation during anaphase (11).

Background References 1. Workman, J.L. and Kingston, R.E. (1998) Annu Rev Biochem 67, 545-79.

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3. Strahl, B.D. and Allis, C.D. (2000) *Nature* 403, 41-5.

4. Cheung, P. et al. (2000) *Cell* 103, 263-71.

5. Bernstein, B.E. and Schreiber, S.L. (2002) Chem Biol 9, 1167-73.

6. Jaskelioff, M. and Peterson, C.L. (2003) Nat Cell Biol 5, 395-9.

7. Thorne, A.W. et al. (1990) Eur J Biochem 193, 701-13.

8. Hendzel, M.J. et al. (1997) *Chromosoma* 106, 348-60.

9. Goto, H. et al. (1999) J Biol Chem 274, 25543-9.

10. Preuss, U. et al. (2003) Nucleic Acids Res 31, 878-85.

11. Dai, J. et al. (2005) Genes Dev 19, 472-88.

**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 IF-IC: Immunofluorescence (Immunocytochemistry)

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**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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