HP1α/β (C7F11) Rabbit mAb



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Applications: W, IP, IHC-P, IF-IC	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 25	Source/Isotype: Rabbit IgG	UniProt ID: #P45973	Entrez-Gene Id: 23468
Product Usage Information		Application Western Blotting Immunoprecipitation Immunohistochemist	, ,	nistry)		Dilution 1:1000 1:25 1:200 1:100
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.				
Specificity/Sensitivity		HP1 α / β (C7F11) Rabbit mAb detects endogenous levels of total HP1 α and HP1 β protein. The antibody does not cross-react with HP1 γ proteins.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to the carboxy terminus of human HP1 $\alpha\!.$				
Background		Heterochromatin protein 1 (HP1) is a family of heterochromatic adaptor molecules involved in both gene silencing and higher order chromatin structure (1). All three HP1 family members (α , β , and γ) are primarily associated with centromeric heterochromatin; however, HP1 β and γ also localize to euchromatic sites in the genome (2,3). HP1 proteins are approximately 25 kDa in size and contain a conserved amino-terminal chromodomain, followed by a variable hinge region and a conserved carboxy-terminal chromoshadow domain. The chromodomain facilitates binding to histone H3 trimethylated at Lys9, a histone "mark" closely associated with centromeric heterochromatin (4,5). The variable hinge region binds both RNA and DNA in a sequence-independent manner (6). The chromoshadow domain mediates the dimerization of HP1 proteins, in addition to binding multiple proteins implicated in gene silencing and heterochromatin formation, including the SUV39H histone methyltransferase, the DNMT1 and DNMT3a DNA methyltransferases, and the p150 subunit of chromatin assembly factor 1 (CAF-1) (7-9). In addition to contributing to heterochromatin formation and propagation, HP1 and SUV39H1 are also found complexed with retinoblastoma (Rb) and E2F6 proteins, both of which function to repress euchromatic gene transcription in quiescent cells (10,11). HP1 proteins are subject to multiple types of post-translational modifications, including phosphorylation, acetylation, methylation, ubiquitination, and sumoylation, suggesting multiple means of regulation (12-14).				
Background References		 Maison, C. and Almouzni, G. (2004) <i>Nat. Rev. Mol. Cell Biol.</i> 5, 296-304. Minc, E. et al. (2000) <i>Cytogenet. Cell Genet.</i> 90, 279-284. Nielsen, A.L. et al. (2001) <i>Mol. Cell</i> 7, 729-739. Lachner, M. et al. (2001) <i>Nature</i> 410, 116-120. Bannister, A.J. et al. (2001) <i>Nature</i> 410, 120-124. Muchardt, C. et al. (2002) <i>EMBO Rep.</i> 3, 975-981. Yamamoto, K. and Sonoda, M. (2003) <i>Biochem. Biophys. Res. Commun.</i> 301, 287-292. Fuks, F. et al. (2003) <i>Nucleic Acids Res.</i> 31, 2305-2312. Murzina, N. et al. (1999) <i>Mol. Cell</i> 4, 529-540. Nielsen, S.J. et al. (2001) <i>Nature</i> 412, 561-565. Ogawa, H. et al. (2002) <i>Science</i> 296, 1132-1136. Minc, E. et al. (1999) <i>Chromosoma</i> 108, 220-234. Zhao, T. et al. (2001) <i>J. Biol. Chem.</i> 276, 9512-9518. Lomberk, G. et al. (2006) <i>Nat. Cell Biol.</i> 8, 407-415. 				292.

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 IHC-P: Immunohistochemistry (Paraffin) IF-IC:

Immunofluorescence (Immunocytochemistry)

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

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