

Applications: Reactivity: Sensitivity: MW (kDa): Source/Isotype: UniProt ID: Entrez-Gene Id: HMRMk Rabbit IgG #Q08945 W, ChIP Endogenous 81 6749 For optimal ChIP results, use 10 μ l of antibody and 10 μ g of chromatin (approximately 4 x 10⁶ cells) per Product Usage IP. This antibody has been validated using SimpleChIP[®] Enzymatic Chromatin IP Kits. Information Application Dilution Western Blotting 1:1000 Chromatin IP 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. Specificity/Sensitivity SSRP1 (E1Y8D) Rabbit mAb recognizes endogenous levels of total SSRP1 protein. This antibody also cross-reacts with protein of unknown origin at 140 kDa in some cell lines. Species predicted to react Chicken based on 100% sequence homology Source / Purification Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val145 of human SSRP1 protein. Suppressor of Ty-16 (SPT16) and structure-specific recognition protein-1 (SSRP1) are subunits of the Background facilitates chromatin transcription (FACT) complex that is essential for transcription elongation (1,2). FACT facilitates RNA polymerase-dependent transcription of chromatin templates by destabilizing the nucleosomes within the open reading frames of active genes (3-5). FACT destabilizes the nucleosomes, which would otherwise act as barriers to RNA polymerase transcription activity, by disrupting histonehistone and histone-DNA contacts that lead to the eviction of the histone H2A-H2B dimer (2,3,6). FACT may also function as a histone chaperone to reassemble nucleosomes after RNA polymerase passage (7). In addition to transcription, FACT activity has been shown to have a role in DNA replication in yeast and in DNA repair by contributing to the activation of p53 by CK2 and by facilitating histone H2AX-H2B exchange upon DNA damage (8,9). **Background References** 1. Winkler, D.D. and Luger, K. (2011) / Biol Chem 286, 18369-74. 2. Orphanides, G. et al. (1999) Nature 400, 284-8. 3. Orphanides, G. et al. (1998) Cell 92, 105-16. 4. Birch, J.L. et al. (2009) EMBO J 28, 854-65. 5. Orphanides, G. and Reinberg, D. (2000) Nature 407, 471-5. 6. Keller, D.M. and Lu, H. (2002) J Biol Chem 277, 50206-13. 7. Belotserkovskaya, R. et al. (2003) Science 301, 1090-3. 8. Schlesinger, M.B. and Formosa, T. (2000) Genetics 155, 1593-606. 9. Heo, K. et al. (2008) Mol Cell 30, 86-97. **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 ChIP: Chromatin IP **Cross-Reactivity Key** H: Human M: Mouse R: Rat Mk: Monkey Trademarks and Patents Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc.

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