RPA70/RPA1 (4D9) Rat mAb



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Applications: W, IP, IF-IC	Reactivity: H Mk	Sensitivity: Endogenous	MW (kDa): 70	Source/Isotype: Rat IgG1	UniProt ID: #P27694	Entrez-Gene Id: 6117
Product Usage Information		Application Western Blotting Immunoprecipitation Immunofluorescence	(Immunocytochem	istry)		Dilution 1:1000 1:50 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		RPA70 (4D9) Rat mAb detects endogenous levels of total RPA70 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a recombinant MBP-fusion protein corresponding to the carboxy-terminal sequence of human RPA70.				
Background		RPA70 (HSSB, REPA1, RF-A, RP-A, p70) is a component of a heterotrimeric complex, composed of 70, 32/30, and 14 kDa subunits, collectively known as RPA. RPA is a single-stranded DNA binding protein, whose DNA binding activity is believed to reside entirely in the 70 kDa subunit. The complex is required for almost all aspects of cellular DNA metabolism such as DNA replication (1-3), recombination, cell cycle and DNA damage checkpoints, and all major types of DNA repair including nucleotide excision, base excision, mismatch, and double-strand break repairs (4-7). In response to genotoxic stress in eukaryotic cells, RPA has been shown to associate with the Rad9/Rad1/Hus1 (9-1-1) checkpoint complex (8). RPA is hyperphosphorylated upon DNA damage or replication stress by checkpoint kinases including ataxia telangiectasia mutated (ATM), ATM and Rad3-related (ATR), and DNA-dependent protein kinase (DNA-PK) (9-11). Phosphorylation of RPA32 occurs at serines 4, 8, and 33 (11). Hyperphosphorylation may alter RPA-DNA and RPA-protein interactions. In addition to the checkpoint partners, RPA interacts with a wide variety of protein partners, including proteins required for normal replication such as RCF, PCNA, and Pol α, and also proteins involved in SV40 replication, such as DNA polymerase I and SV40 large T antigen (10,12).				
Background References		1. Liu, V.F. and Weaver, D.T. (1993) <i>Mol. Cell Biol.</i> 13, 7222-31. 2. Wobbe, C.R. et al. (1987) <i>Proc. Natl. Acad. Sci. USA</i> 84, 1834-8. 3. Fairman, M.P. and Stillman, B. (1988) <i>EMBO J.</i> 7, 1211-8. 4. Wold, M.S. and Kelly, T. (1988) <i>Proc. Natl. Acad. Sci. USA</i> 85, 2523-7. 5. Zhou, B.B. and Elledge, S.J. (2000) <i>Nature</i> 408, 433-9. 6. Kastan, M.B. and Bartek, J. (2004) <i>Nature</i> 432, 316-23. 7. Sancar, A. et al. (2004) <i>Annu. Rev. Biochem.</i> 73, 39-85. 8. Guo, S. et al. (2006) <i>J Biol Chem</i> 281, 21607-16. 9. Wu, X. et al. (2005) <i>Oncogene</i> 24, 4728-35. 10. Binz, S.K. et al. <i>DNA Repair (Amst)</i> 3, 1015-24. 11. Nuss, J.E. et al. (2005) <i>Biochemistry</i> 44, 8428-37. 12. Yuzhakov, A. et al. (1999) <i>EMBO J.</i> 18, 6189-99.				
Species Reacti	vity	Species reactivity is de	etermined by testin	g in at least one approve	ed 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)

Cross-Reactivity Key

H: Human Mk: Monkey

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