p53 (1C12) Mouse mAb



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WB, IP, IF-IC, FC-FP, H M R Hm Mk Endogenous 53 Mouse IgG1 #P04637 7157 ChIP	, , -, - ,	Reactivity: H M R Hm Mk	Sensitivity: Endogenous	MW (kDa): 53	Source/Isotype: Mouse IgG1	UniProt ID: #P04637	Entrez-Gene Id: 7157
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Product Usage Information

For optimal ChIP results, use 2.5 μ I of antibody and 10 μ g of chromatin (approximately 4 x 10⁶ cells) per IP. This antibody has been validated using SimpleChIP® Enzymatic Chromatin IP Kits.

Application	Dilution
Western Blotting	1:1000
Immunoprecipitation	1:500
Immunofluorescence (Immunocytochemistry)	1:2000
Flow Cytometry (Fixed/Permeabilized)	1:1600
Chromatin IP	1:200

Storage

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20° C. Do not aliquot the antibody.

For a carrier free (BSA and azide free) version of this product see product #15755.

Specificity / Sensitivity

p53 (1C12) Mouse mAb detects endogenous levels of total p53 protein.

Species predicted to react based on 100% sequence homology:

Rabbit

Source / Purification

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser20 of human p53.

Background

The p53 tumor suppressor protein plays a major role in cellular response to DNA damage and other genomic aberrations. Activation of p53 can lead to either cell cycle arrest and DNA repair or apoptosis (1). p53 is phosphorylated at multiple sites in vivo and by several different protein kinases in vitro (2,3). DNA damage induces phosphorylation of p53 at Ser15 and Ser20 and leads to a reduced interaction between p53 and its negative regulator, the oncoprotein MDM2 (4). MDM2 inhibits p53 accumulation by targeting it for ubiquitination and proteasomal degradation (5,6), p53 can be phosphorylated by ATM, ATR, and DNA-PK at Ser15 and Ser37. Phosphorylation impairs the ability of MDM2 to bind p53, promoting both the accumulation and activation of p53 in response to DNA damage (4,7). Chk2 and Chk1 can phosphorylate p53 at Ser20, enhancing its tetramerization, stability, and activity (8,9). p53 is phosphorylated at Ser392 in vivo (10,11) and by CAK in vitro (11). Phosphorylation of p53 at Ser392 is increased in human tumors (12) and has been reported to influence the growth suppressor function. DNA binding, and transcriptional activation of p53 (10,13,14). p53 is phosphorylated at Ser6 and Ser9 by CK1 δ and CK1 ϵ both in vitro and in vivo (13,15). Phosphorylation of p53 at Ser46 regulates the ability of p53 to induce apoptosis (16). Acetylation of p53 is mediated by p300 and CBP acetyltransferases. Inhibition of deacetylation suppressing MDM2 from recruiting HDAC1 complex by p19 (ARF) stabilizes p53. Acetylation appears to play a positive role in the accumulation of p53 protein in stress response (17). Following DNA damage, human p53 becomes acetylated at Lys382 (Lys379 in mouse) in vivo to enhance p53-DNA binding (18). Deacetylation of p53 occurs through interaction with the SIRT1 protein, a deacetylase that may be involved in cellular aging and the DNA damage response (19).

Background References

- 1. Levine, A.J. (1997) Cell 88, 323-31.
- 2. Meek, D.W. (1994) Semin Cancer Biol 5, 203-10.
- 3. Milczarek, G.J. et al. (1997) Life Sci 60, 1-11.
- 4. Shieh, S.Y. et al. (1997) Cell 91, 325-34.
- 5. Chehab, N.H. et al. (1999) Proc Natl Acad Sci U S A 96, 13777-82.

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- 6. Honda, R. et al. (1997) FEBS Lett 420, 25-7.
- 7. Tibbetts, R.S. et al. (1999) Genes Dev 13, 152-7.
- 8. Shieh, S.Y. et al. (1999) EMBO J 18, 1815-23.
- 9. Hirao, A. et al. (2000) Science 287, 1824-7.
- 10. Hao, M. et al. (1996) J Biol Chem 271, 29380-5.
- 11. Lu, H. et al. (1997) $\textit{Mol Cell Biol}\ 17,\,5923-34.$
- 12. Ullrich, S.J. et al. (1993) Proc Natl Acad Sci U S A 90, 5954-8.
- 13. Kohn, K.W. (1999) Mol Biol Cell 10, 2703-34.
- 14. Lohrum, M. and Scheidtmann, K.H. (1996) Oncogene 13, 2527-39.
- 15. Knippschild, U. et al. (1997) Oncogene 15, 1727-36.
- 16. Oda, K. et al. (2000) Cell 102, 849-62.
- 17. Ito, A. et al. (2001) EMBO J 20, 1331-40.
- 18. Sakaguchi, K. et al. (1998) Genes Dev 12, 2831-41.
- 19. Solomon, J.M. et al. (2006) Mol Cell Biol 26, 28-38.

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

WB: Western Blotting **IP**: Immunoprecipitation **IF-IC**: Immunofluorescence (Immunocytochemistry) **FC-FP**: Flow Cytometry (Fixed/Permeabilized) **ChIP**: Chromatin IP

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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