

p21 (H-164): sc-756



The Power to Question

BACKGROUND

It is now well established that cyclins play a positive role in promoting cell cycle transitions via their ability to associate with and activate their cognate cyclin-dependent kinases (Cdks). Cdk2 associates with cyclins A, D and E and has been implicated in the control of the G₁ to S phase transition in mammals. A novel Cdk-interacting protein, p21 (also designated WAF1/CIP1), has been identified in cyclin A, cyclin D1, cyclin E and Cdk2 immunoprecipitates. p21 is a potent, tight-binding inhibitor of Cdks and can inhibit the phosphorylation of Rb by cyclin A-Cdk 2, cyclin E-Cdk2, cyclin D1-Cdk4 and cyclin D2-Cdk4 complexes. Expression of p21 is inducible by wildtype, but not mutant, p53. The mouse homolog of p21 is designated CAP20.

REFERENCES

- Sherr, C.J. 1993. Mammalian G₁ cyclins. *Cell* 73: 1059-1065.
- Harper, J.W., et al. 1993. The p21 Cdk-interacting protein Cip1 is a potent inhibitor of G₁ cyclin-dependent kinases. *Cell* 75: 805-816.

CHROMOSOMAL LOCATION

Genetic locus: CDKN1A (human) mapping to 6p21.2; Cdkn1a (mouse) mapping to 17 A3.3.

SOURCE

p21 (H-164) is a rabbit polyclonal antibody raised against amino acids 1-164 representing full length p21 of human origin.

PRODUCT

Each vial contains 200 µg IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

p21 (H-164) is recommended for detection of p21 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000); partially cross-reactive with the related mitotic inhibitory protein, p27.

Suitable for use as control antibody for p21 siRNA (h): sc-29427, p21 siRNA (m): sc-29428, p21 shRNA Plasmid (h): sc-29427-SH, p21 shRNA Plasmid (m): sc-29428-SH, p21 shRNA (h) Lentiviral Particles: sc-29427-V and p21 shRNA (m) Lentiviral Particles: sc-29428-V.

Molecular Weight of p21: 21 kDa.

Positive Controls: C32 nuclear extract: sc-2136, C32 + PMA nuclear extract: sc-2137 or HeLa nuclear extract: sc-2120.

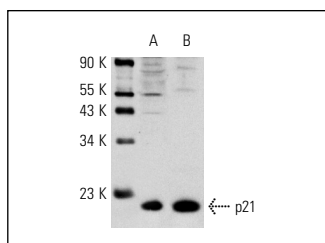
STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

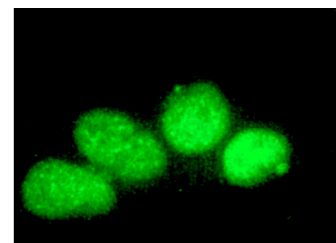
RESEARCH USE

For research use only, not for use in diagnostic procedures.

DATA



Western blot analysis of p21 expression in C32 nuclear extracts (A,B). Antibodies tested include p21 (H-164): sc-756 (A) and p21 (C-19)-G: sc-397-G (B).



p21 (H-164): sc-756. Immunofluorescence staining of methanol-fixed C32 cells showing nuclear staining.

SELECT PRODUCT CITATIONS

- Russell, A., et al. 1999. Inhibitory effect of p21 in MDF-7 cells is overcome by its coordinated stabilization with D-type cyclins. *Oncogene* 18: 6454-6459.
- Su, D., et al. 2010. Role of p38 MAPK pathway in BMP4-mediated Smad-dependent premature senescence in lung cancer cells. *Biochem. J.* 433: 333-343.
- Marcar, L., et al. 2010. Mage-A cancer/testis antigens inhibit p53 function by blocking its interaction with chromatin. *Cancer Res.* 70: 10362-10370.
- Cho, S.J., et al. 2010. RNPC1 modulates the RNA-binding activity of, and cooperates with, HuR to regulate p21 mRNA stability. *Nucleic Acids Res.* 38: 2256-2267.
- Sistrunk, C., et al. 2011. Skp2 is necessary for Myc-induced keratinocyte proliferation but dispensable for Myc oncogenic activity in the oral epithelium. *Am. J. Pathol.* 178: 2470-2477.
- Sze, S.C., et al. 2011. Regulation of p21, MMP-1, and MDR-1 expression in human colon carcinoma HT29 cells by Tian Xian liquid, a chinese medicinal formula, *in vitro* and *in vivo*. *Integr. Cancer Ther.* 10: 58-69.
- Knobel, P.A., et al. 2011. Inhibition of REV3 expression induces persistent DNA damage and growth arrest in cancer cells. *Neoplasia* 13: 961-970.
- O'Dell, M.R., et al. 2012. Kras(G12D) and p53 mutation cause primary intrahepatic cholangiocarcinoma. *Cancer Res.* 72: 1557-1567.
- Capparelli, C., et al. 2012. CTGF drives autophagy, glycolysis and senescence in cancer-associated fibroblasts via HIF1 activation, metabolically promoting tumor growth. *Cell Cycle* 11: 2272-2284.

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Try **p21 (F-5): sc-6246** or **p21 (F-8): sc-271610**, our highly recommended monoclonal alternatives to p21 (H-164). Also, for AC, HRP, FITC, PE, Alexa Fluor® 488 and Alexa Fluor® 647 conjugates, see **p21 (F-5): sc-6246**.