cleaved Rb (172C1094): sc-56217



The Power to Overtion

BACKGROUND

Pediatric cancer retinoblastoma and the formation of other human tumors can be attributed to mutations in the retinoblastoma tumor suppressor gene. The retinoblastoma tumor suppressor gene product, known as Rb or pRb, regulates differentiation, apoptosis and cell cycle control by coordinating the cell cycle, at G_1/S , with transcriptional machinery that includes the heterodimeric E2F family. During G₁, cyclin D (D1, D2, D3)-dependent kinasemediated phosphoryl-ation of Rb at Ser 795 marks the conversion of Rb from a transcriptionally repressive, hypophosphorylated state to an inactive, phosphorylated state, which may be sustained through mitosis by differential phosphorylation of up to 16 putative serine or threonine residues, including Ser 249/Thr 252, Thr 373, Thr 356, Ser 780, Ser 807/Ser 811 and Thr 821/ Thr 826. Hypophosphorylated Rb represses the transcription of genes controlling cell cycle through direct protein-protein interactions, by binding and inactivating the promoters of transcription factors, and through recruitment of histone deacetylase, which deacetylates promoter regions and enhances nucleosome formation, thereby masking transcription enhancing *cis* elements.

REFERENCES

- Bremner, R., Cohen, B.L., Sopta, M., Hamel, P.A., Ingles, C.J., Gallie, B.L. and Phillips, R.A. 1995. Direct transcriptional repression by pRb and its reversal by specific cyclins. Mol. Cell. Biol. 15: 3256-3265.
- Weinberg, R.A. 1995. The retinoblastoma protein and cell cycle control. Cell 81: 323-330.
- 3. Sherr, C.J. 1996. Cancer cell cycles. Science 274: 1672-1677.
- Connell-Crowley, L., Harper, J.W. and Goodrich, D.W. 1997. Cyclin D1/Cdk4 regulates retinoblastoma protein-mediated cell cycle arrest by site-specific phosphorylation. Mol. Biol. Cell 8: 287-301.
- Luo, R.X., Postigo, A.A. and Dean, D.C. 1998. Rb interacts with histone deacetylase to repress transcription. Cell 92: 463-473.
- 6. Driscoll, B., T'Ang, A., Hu, Y.H., Yan, C.L., Fu, Y., Luo, Y., Wu, K.J., Wen, S., Shi, X.H., Barsky, L., Weinberg, K., Murphree, A.L. and Fung, Y.K. 1999. Discovery of a regulatory motif that controls the exposure of specific upstream cyclin-dependent kinase sites that determine both conformation and growth suppressing activity of pRb. J. Biol. Chem. 274: 9463-9471.
- 7. Hu, X., Cress, W. D., Zhong, Q. and Zuckerman, K.S. 2000. Transforming growth factor β inhibits the phosphorylation of pRb at multiple serine/threonine sites and differentially regulates the formation of pRb family-E2F complexes in human myeloid leukemia cells. Biochem. Biophys. Res. Commun. 276: 930-939.
- Barrientes, S., Cooke, C. and Goodrich, D.W. 2000. Glutamic acid mutagenesis of retinoblastoma protein phosphorylation sites has diverse effects on function. Oncogene 19: 562-570.
- 9. LocusLink Report (LocusID: 5925). http://www.ncbi.nlm.nih.gov/LocusLink/

STORAGE

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

CHROMOSOMAL LOCATION

Genetic locus: RB1 (human) mapping to 13q14.2; Rb1 (mouse) mapping to 14 D3.

SOURCE

cleaved Rb (172C1094) is a mouse monoclonal antibody raised against a short amino acid sequence containing the neoepitope at epitope corresponding to the caspase-3 induced cleavage site of Rb of human origin.

PRODUCT

Each vial contains 100 μg lgG_1 in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

cleaved Rb (172C1094) is recommended for detection of the 68 kDa fragment of cleaved retinoblastoma (Rb) of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000) and immuno-precipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)].

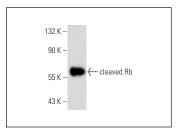
Suitable for use as control antibody for Rb siRNA (h): sc-29468, Rb siRNA (m): sc-29469, Rb shRNA Plasmid (h): sc-29468-SH, Rb shRNA Plasmid (m): sc-29469-SH, Rb shRNA (h) Lentiviral Particles: sc-29468-V and Rb shRNA (m) Lentiviral Particles: sc-29469-V.

Molecular Weight of pro Rb: 106 kDa.

Molecular Weight of cleaved Rb: 68 kDa.

Positive Controls: K-562 whole cell lysate: sc-2203, HeLa whole cell lysate: sc-2200 or A-431 whole cell lysate: sc-2201.

DATA



cleaved Rb (172C1094): sc-56217. Western blot analysis of cleaved Rb expression in K-562 whole cell lysate

RESEARCH USE

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

PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.