

p-p27 (Thr 198): sc-130603

BACKGROUND

p27 associates with cyclin and cyclin-dependent kinase complexes to inhibit their kinase activity and contribute to the control of cell proliferation. p27 is phosphorylated on many sites, including threonine 187, *in vivo*, with the predominant phosphorylation site being serine 10. The extent of serine 10 phosphorylation by proline-directed kinase is markedly increased in cells in the G₀-G₁ phase of the cell cycle compared to cells in the S or M phase. p27 concentration is regulated predominantly by posttranslational mechanisms. p27 is degraded by both the ubiquitin-proteasome pathway and ubiquitin-independent proteolysis. Regulation of ubiquitin-mediated proteolysis is often achieved through ubiquitination of the targeted phosphorylated protein, which renders it more susceptible to degradation.

REFERENCES

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- Hannon, G.J., et al. 1994. p15INK4B is a potential effector of TGFβ-induced cell cycle arrest. *Nature* 371: 257-260.
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- Polyak, K., et al. 1994. Cloning of p27KIP1, a cyclin-dependent kinase inhibitor and a potential mediator of extracellular antimitogenic signals. *Cell* 78: 59-66.
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CHROMOSOMAL LOCATION

Genetic locus: CDKN1B (human) mapping to 12p13.1.

SOURCE

p-p27 (Thr 198) is a rabbit polyclonal antibody raised against a short amino acid sequence containing Thr 198 phosphorylated p27 of human origin.

PRODUCT

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

STORAGE

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

APPLICATIONS

p-p27 (Thr 198) is recommended for detection of Thr 198 phosphorylated p27 of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for p27 siRNA (h): sc-29429, p27 shRNA Plasmid (h): sc-29429-SH and p27 shRNA (h) Lentiviral Particles: sc-29429-V.

Molecular Weight of p-p27: 27 kDa.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western blotting: use goat anti-rabbit IgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit IgG-HRP: sc-2030 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto B Blocking Reagent: sc-2335 (use 50 mM NaF, sc-24988, as diluent), Western Blotting Luminol Reagent: sc-2048 and Lambda Phosphatase: sc-200312A.

SELECT PRODUCT CITATIONS

- Coëffier, M., et al. 2013. Enteral delivery of proteins stimulates protein synthesis in human duodenal mucosa in the fed state through a mammalian target of rapamycin-independent pathway. *Am. J. Clin. Nutr.* 97: 286-294.

RESEARCH USE

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

PROTOCOLS

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