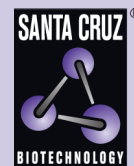


## p27 (F-8): sc-1641



The Power to Question

## BACKGROUND

Cell cycle progression is regulated by a series of cyclin-dependent kinases that consist of catalytic subunits, designated Cdk, and activating subunits, designated cyclins. Orderly progression through the cell cycle requires the activation and inactivation of different cyclin-Cdks at appropriate times. A series of proteins has been recently described that function as "mitotic inhibitors". These include p21, the levels of which are elevated upon DNA damage in G<sub>1</sub> in a p53-dependent manner, p16 and a more recently described p16 related inhibitor designated p15. A p21 related protein, p27, has been described as a negative regulator of G<sub>1</sub> progression and has been speculated to function as a possible mediator of TGF-induced G<sub>1</sub> arrest. p27 interacts strongly with D-type cyclins and Cdk4 *in vitro* and to a lesser extent with cyclin E and Cdk2.

## CHROMOSOMAL LOCATION

Genetic locus: CDKN1B (human) mapping to 12p13.1; Cdkn1b (mouse) mapping to 6 G1.

## SOURCE

p27 (F-8) is a mouse monoclonal antibody raised against amino acids 1-197 representing full length p27 of mouse origin.

## PRODUCT

Each vial contains 200 µg IgG<sub>1</sub> kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

p27 (F-8) is available conjugated to agarose (sc-1641 AC), 500 µg/0.25 ml agarose in 1 ml, for IP; to HRP (sc-1641 HRP), 200 µg/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-1641 PE), fluorescein (sc-1641 FITC), Alexa Fluor® 488 (sc-1641 AF488), Alexa Fluor® 546 (sc-1641 AF546), Alexa Fluor® 594 (sc-1641 AF594) or Alexa Fluor® 647 (sc-1641 AF647), 200 µg/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-1641 AF680) or Alexa Fluor® 790 (sc-1641 AF790), 200 µg/ml, for Near-Infrared (NIR) WB, IF and FCM.

In addition, p27 (F-8) is available conjugated to either TRITC (sc-1641 TRITC, 200 µg/ml) or Alexa Fluor® 405 (sc-1641 AF405), 100 µg/2 ml, for IF, IHC(P) and FCM.

## APPLICATIONS

p27 (F-8) is recommended for detection of p27 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), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500), flow cytometry (1 µg per 1 x 10<sup>6</sup> cells) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

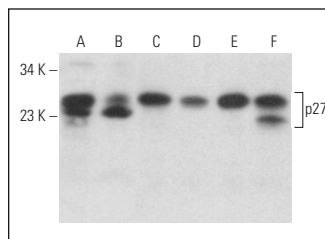
Suitable for use as control antibody for p27 siRNA (h): sc-29429, p27 siRNA (m): sc-29430, p27 shRNA Plasmid (h): sc-29429-SH, p27 shRNA Plasmid (m): sc-29430-SH, p27 shRNA (h) Lentiviral Particles: sc-29429-V and p27 shRNA (m) Lentiviral Particles: sc-29430-V.

Molecular Weight of p27: 27 kDa.

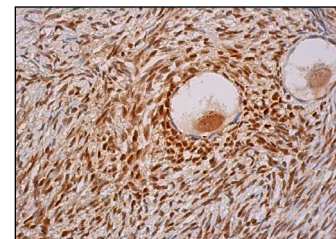
## STORAGE

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

## DATA



p27 (F-8): sc-1641. Western blot analysis of p27 expression in MM-142 (A), RAW 264.7 (B), NAMALWA (C), BJAB (D), Raji (E) and C6 (F) whole cell lysates.



p27 (F-8): sc-1641. Immunoperoxidase staining of formalin fixed, paraffin-embedded human ovary tissue showing nuclear and cytoplasmic staining of ovarian stroma cells and oocytes.

## SELECT PRODUCT CITATIONS

- Kawasaki, H., et al. 1998. Distinct roles of the co-activators p300 and CBP in retinoic-acid-induced F9-cell differentiation. *Nature* 393: 284-289.
- Cheung, B., et al. 1998. Favorable prognostic significance of high-level retinoic acid receptor  $\beta$  expression in neuroblastoma mediated by effects on cell cycle regulation. *Oncogene* 17: 751-759.
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- Paglini, G., et al. 1998. Evidence for the participation of the neuron-specific CDK5 activator P35 during laminin-enhanced axonal growth. *J. Neurosci.* 18: 9858-9869.
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- Nakagawa, H., et al. 2018. Sodium butyrate induces senescence and inhibits the invasiveness of glioblastoma cells. *Oncol. Lett.* 15: 1495-1502.
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- Leem, D.G., et al. 2018. Dammarane-type triterpene ginsenoside-Rg18 inhibits human non-small cell lung cancer A549 cell proliferation via G<sub>1</sub> phase arrest. *Oncol. Lett.* 15: 6043-6049.
- Ma, Y., et al. 2018. Anticancer effect of exogenous hydrogen sulfide in cisplatin-resistant A549/DDP cells. *Oncol. Rep.* 39: 2969-2977.

## RESEARCH USE

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

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