

cyclin D1 (M-20): sc-718



The Power to Question

BACKGROUND

The proliferation of eukaryotic cells is controlled at specific points in the cell cycle, particularly at the G₁ to S and the G₂ to M transitions. It is well established that the Cdc2 p34-cyclin B protein kinase plays a critical role in the G₂ to M transition while cyclin A associates with Cdk2 p33 and functions in S phase. Considerable effort directed towards the identification of G₁ cyclins has led to the isolation of cyclin D, cyclin C and cyclin E. Of these, cyclin D corresponds to a putative human oncogene, designated PRAD1, which maps at the site of the Bcl1 rearrangement in certain lymphomas and leukemias. Two additional human type D cyclins, as well as their mouse homologs, have been identified. Evidence has established that members of the cyclin D family function to regulate phosphorylation of the retinoblastoma gene product, thereby activating E2F transcription factors.

REFERENCES

1. Draetta, G. 1990. Cell cycle control in eukaryotes: molecular mechanisms of Cdc2 activation. *Trends Biol. Sci.* 15: 378-383.
2. Xiong, Y., et al. 1991. Human D-type cyclin. *Cell* 65: 691-699.
3. Xiong, Y., et al. 1992. Molecular cloning and chromosomal mapping of CCND genes encoding human D-type cyclins. *Genomics* 13: 575-584.

CHROMOSOMAL LOCATION

Genetic locus: CCND1 (human) mapping to 11q13.3, CCND2 (human) mapping to 12p13.32; Ccnd1 (mouse) mapping to 7 F5, Ccnd2 (mouse) mapping to 6 F3.

SOURCE

cyclin D1 (M-20) is an affinity purified rabbit polyclonal antibody raised against a peptide mapping at the C-terminus of cyclin D1 of human origin.

PRODUCT

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

Blocking peptide available for competition studies, sc-718 P, (100 µg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

cyclin D1 (M-20) is recommended for detection of cyclin D1 and, to a lesser extent, cyclin D2 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) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Molecular Weight of cyclin D1: 37 kDa.

Positive Controls: cyclin D1 (h4): 293T Lysate: sc-171254, C32 nuclear extract: sc-2136 or KNRK nuclear extract: sc-2141.

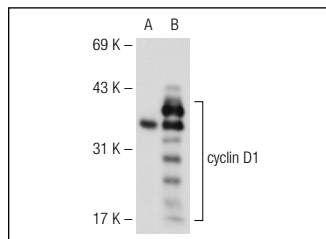
RESEARCH USE

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

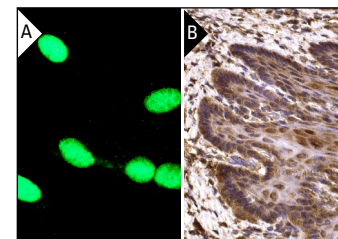
STORAGE

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

DATA



cyclin D1 (M-20): sc-718. Western blot analysis of cyclin D1 expression in non-transfected: sc-117752 (A) and human cyclin D1 transfected: sc-171254 (B) 293T whole cell lysates.



cyclin D1 (M-20): sc-718. Immunofluorescence staining of methanol-fixed C32 cells showing nuclear localization (A). Immunoperoxidase staining of formalin fixed, paraffin-embedded human esophagus tissue showing nuclear and cytoplasmic staining of squamous epithelial cells (B).

SELECT PRODUCT CITATIONS

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3. Lappano, R., et al. 2012. MIBE acts as antagonist ligand of both estrogen receptor α and GPER in breast cancer cells. *Breast Cancer Res.* 14: R12.
4. Lappano, R., et al. 2012. Two novel GPER agonists induce gene expression changes and growth effects in cancer cells. *Curr. Cancer Drug Targets* 12: 531-542.
5. De Marco, P., et al. 2012. Insulin-like growth factor-I regulates GPER expression and function in cancer cells. *Oncogene* 32: 678-688.
6. Catalano, M.G., et al. 2012. Cytotoxic activity of the histone deacetylase inhibitor panobinostat (LBH589) in anaplastic thyroid cancer *in vitro* and *in vivo*. *Int. J. Cancer* 130: 694-704.
7. Ling, C., et al. 2012. A deuterated analog of dasatinib disrupts cell cycle progression and displays anti-non-small cell lung cancer activity *in vitro* and *in vivo*. *Int. J. Cancer* 131: 2411-2419.
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9. Lee, S.H., et al. 2012. Capsaicin represses transcriptional activity of β -catenin in human colorectal cancer cells. *J. Nutr. Biochem.* 23: 646-655.
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