# DP-1/3 (L7-7): sc-101031



The Power to Question

#### **BACKGROUND**

The human retinoblastoma gene product appears to play an important role in the negative regulation of cell proliferation. Functional inactivation of Rb can be mediated either through mutation or as a consequence of interaction with DNA tumor virus-encoded proteins. Of all the Rb associations described to date, the identification of a complex between Rb and the transcription factor E2F most directly implicates Rb in regulation of cell proliferation. E2F was originally identified through its role in transcriptional activation of the adeno-virus E2 promoter. Sequences homologous to the E2F binding site have been found upstream of a number of genes that encode proteins with putative functions in the  $G_1$  and S phases of the cell cycle. E2F-1 forms heterodimers with a second protein, designated DP-1, forming an "active" E2F transcriptional regulatory complex. Additional members of the E2F family include E2F-2, E2F-3, E2F-4, E2F-5 and DP-2.

#### **REFERENCES**

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- 2. Nevins, J.R. 1992. E2F: a link between the Rb tumor suppressor protein and viral oncoproteins. Science 258: 424-429.
- 3. Helin, K., et al. 1993. Heterodimerization of the transcription factors E2F-1 and DP-1 leads to cooperative *trans*-activation. Genes Dev. 7: 1850-1861.
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- 5. Ginsberg, D., et al. 1994. E2F-4, a new member of the E2F transcription factor family, interacts with p107. Genes Dev. 8: 2665-2679.
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- 7. Zhang, Y., et al. 1995. Cloning and characterization of human DP2, a novel dimerization partner of E2F. Oncogene 10: 2085-2093.
- 8. Ishida, H., et al. 2005. Identification and characterization of novel isoforms of human DP-1: DP-1 $\alpha$  regulates the transcriptional activity of E2F1 as well as cell cycle progression in a dominant-negative manner. J. Biol. Chem. 280: 24642-24648.
- 9. Reimer, D., et al. 2006. Heterogeneous cross-talk of E2F family members is crucially involved in growth modulatory effects of interferon- $\gamma$  and EGF. Cancer Biol. Ther. 5: 771-776.

## **CHROMOSOMAL LOCATION**

Genetic locus: TFDP1 (human) mapping to 13q34, TFDP3 (human) mapping to Xq26.2.

### **SOURCE**

DP-1/3 (L7-7) is a mouse monoclonal antibody raised against recombinant DP-1/3 of human origin.

## **PRODUCT**

Each vial contains 100  $\mu g$   $lgG_{2a}$  kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

## **APPLICATIONS**

DP-1/3 (L7-7) is recommended for detection of DP-1/3 of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g 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).

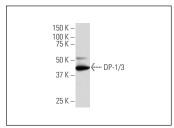
Molecular Weight of DP-1/3: 46 kDa.

Positive Controls: IMR-32 cell lysate: sc-2409.

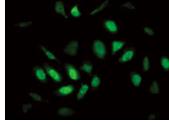
## **RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG $\kappa$  BP-HRP: sc-516102 or m-lgG $\kappa$  BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker<sup>TM</sup> Molecular Weight Standards: sc-2035, UltraCruz\* Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-lgG $\kappa$  BP-FITC: sc-516140 or m-lgG $\kappa$  BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz\* Mounting Medium: sc-24941 or UltraCruz\* Hard-set Mounting Medium: sc-359850.

## DATA



DP-1/3 (L7-7): sc-101031. Western blot analysis of DP-1/3 expression in IMR-32 whole cell lysate.



DP-1/3 (L7-7): sc-101031. Immunofluorescence staining of paraformaldehyde-fixed HeLa cells showing nuclear localization.

#### **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.

### **PROTOCOLS**

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