# HSV-1 ICP27 (vP-20): sc-17544



The Power to Question

#### **BACKGROUND**

Herpes simplex virus type 1 (HSV-1) is a neurotropic virus that establishes lifelong latent infections in the sensory neurons of the host. One of the HSV-1 proteins involved in converting the cell into an efficient producer of viral gene products is the infected cell polypeptide 27 or ICP27. HSV-1 immediate-early protein ICP27 is a nuclear phosphoprotein that is required for viral growth during lytic infection. Analysis of viral mutants defective in this function has shown that ICP27 has a number of effects on gene expression including a contribution to the shut off of host protein synthesis, the stimulation of HSV-1 early gene expression and DNA replication, and the induction of late viral gene products. ICP27 performs these functions primarily post-transcriptionally at the level of RNA processing. ICP27 affects three important RNA processing events: polyadenylation, splicing and nuclear RNA export.

# **REFERENCES**

- Hardy, W.R. and Sandri-Goldin, R.M. 1994. Herpes simplex virus inhibits host cell splicing, and regulatory protein ICP27 is required for this effect. J. Virol. 68: 7790-7799.
- Hibbard, M.K. and Sandri-Goldin, R.M. 1995. Arginine-rich regions succeeding the nuclear localization region of the HSV-1 regulatory protein ICP27 are required for efficient nuclear localization and late gene expression.
   J. Virol. 69: 4656-4667.
- Sandri-Goldin, R.M. 1998. ICP27 mediates herpes simplex virus RNA export by shuttling through a leucine-rich nuclear export signal and binding viral intronless RNAs through an RGG motif. Genes Dev. 12: 868-879.
- 4. Zhi, Y., Sciabica, K.S. and Sandri-Goldin., R.M. 1999. Self-interaction of the herpes simplex virus type 1 regulatory protein ICP27. Virology 257: 341-351.
- Stingly, S.W., Garcia Ramirez, J.J., Aguilar, S.A., Simmen, K., Sandri-Goldin, R.M., Ghazal, P.H. and Wagner, E.K. 2000. Global analysis of herpes simplex virus type 1 transcription using an oligonucleotide-based DNA microarray. J. Virol. 74: 9916-9927.

#### **SOURCE**

 $HSV-1\ ICP27\ (vP-20)$  is an affinity purified goat polyclonal antibody raised against a peptide mapping near the N-terminus of  $HSV-1\ ICP27$ .

# **PRODUCT**

Each vial contains 200  $\mu g$  lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-17544 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

## **STORAGE**

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

## **PROTOCOLS**

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

### **APPLICATIONS**

HSV-1 ICP27 (vP-20) is recommended for detection of ICP27 of HSV-1 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).

Molecular Weight of HSV-1 ICP27: 63 kDa.

# **RECOMMENDED SECONDARY REAGENTS**

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048.

#### **SELECT PRODUCT CITATIONS**

- 1. Choudhary, A., Hiscott, P., Hart, C.A., Kaye, S.B., Batterbury, M. and Grierson, I. 2005. Suppression of Thrombospondin 1 and 2 production by herpes simplex virus 1 infection in cultured keratocytes. Mol. Vis. 11: 163-168.
- 2. Yao, F., Murakami, N., Bleiziffer, O., Zhang, P., Akhrameyeva, N.V., Xu, X. and Brans, R. 2010. Development of a regulatable oncolytic herpes simplex virus type 1 recombinant virus for tumor therapy. J. Virol. 84: 8163-8171.

# **RESEARCH USE**

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

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