SANTA CRUZ BIOTECHNOLOGY, INC.

HSV-1 (202): sc-57864



BACKGROUND

The herpes simplex virus (HSV) (also known as cold sore, night fever or fever blister) is a virus that causes a contagious disease. The HSV-1 strain generally appears in the orafacial organs. All herpes viruses are morphologically identical: they have a large double-stranded DNA genome and the virion consists of an icosahedral nucleocapsid which is surrounded by a lipid bilayer envelope. Following primary infection, the virus establishes a latent infection in the host and may reactivate at any stage. Reactivation is frequently, but not always, associated with further disease.

REFERENCES

- 1. McCormick, F.P. and Newton, A.A. 1975. Polyamine metabolism in cells infected with herpes simplex virus. J. Gen. Virol. 127: 25-33.
- 2. Trusal, L.R., Anthony, A. and Docherty, J.J. 1975. Differential feulgendeoxyribonucleic acid hydrolysis patterns of herpes simplex virus type 1 and type 2 infected cells. J. Histochem. Cytochem. 23: 283-238.
- Rapp, F. and Buss, E.R. 1976. Comparison of herpes simplex virus isolates using a quantitative selection assay for transformation. Intervirology 6: 72-82.
- Purifoy, D.J. 1976. Comparison of DNA polymerase activities induced by herpes simplex virus types 1 and 2. Intervirology 6: 356-366.
- Slomka, M.J. 1996. Seroepidemiology and control of genital herpes: the value of type specific antibodies to herpes simplex virus. Commun. Dis. Rep. CDR Rev. 3: R41-45.
- Mador, N., Panet, A. and Steiner, I. 2002. The latency-associated gene of herpes simplex virus type 1 (HSV-1) interferes with superinfection by HSV-1. J. Neurovirol. 8: 97-102.
- Burton, E.A., Huang, S., Goins, W.F. and Glorioso, J.C. 2003. Use of the herpes simplex viral genome to construct gene therapy vectors. Methods Mol. Med. 76: 1-31.
- Ni, Y., Goldman, D., Hoffman, B. and Brooks, P.J. 2003. Overexpression of an epitope-tagged serotonin transporter in serotonin neurons of the dorsal raphe nucleus using a defective HSV-1 vector. Behav. Brain Res. 138: 133-143.
- Nozawa, N., Daikoku, T., Koshizuka, T., Yamauchi, Y., Yoshikawa, T. and Nishiyama, Y. 2003. Subcellular localization of herpes simplex virus type 1 UL51 protein and role of palmitoylation in Golgi apparatus targeting. J Virol. 77: 3204-3216.

SOURCE

HSV-1 (202) is a mouse monoclonal antibody raised against HSV-1 and -2 infected cells.

PRODUCT

Each vial contains 100 μ l ascites containing lgG_{2a} with < 0.1% sodium azide.

RESEARCH USE

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

APPLICATIONS

HSV-1 (202) is recommended for detection of herpes simplex virus 1 by immunofluorescence (starting dilution to be determined by researcher, dilution range 1:50-1:2500).

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Immunofluorescence: use goat anti-mouse IgG-FITC: sc-2010 (dilution range: 1:100-1:400) or goat anti-mouse IgG-TR: sc-2781 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

STORAGE

For immediate and continuous use, store at 4° C for up to one month. For sporadic use, freeze in working aliquots in order to avoid repeated freeze/ thaw cycles. If turbidity is evident upon prolonged storage, clarify solution by centrifugation.

PROTOCOLS

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