

A27L Protein from Vaccinia Virus (WR) with C-terminal Histidine Tag, Recombinant from Baculovirus

Catalog No. NR-22133

For research use only. Not for use in humans.

Contributor:

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Manufacturer:

BEI Resources

Product Description:

A full-length recombinant form of the A27L membrane glycoprotein of the Western Reserve (WR) strain of vaccinia virus containing a C-terminal histidine-tag was produced in Sf9 insect cells using a baculovirus expression system and purified using nickel affinity chromatography. The predicted protein sequence is shown in Figure 1. The full length A27L protein is 110 residues (GenPept: [P11258](#)).^{1,2,3}

Material Provided:

Each vial contains 50 to 200 µg of NR-22133 in PBS (pH 7.4) with 0.05% polysorbate (v/v). The protein concentration, expressed as mg/mL, is shown on the Certificate of Analysis.

Packaging/Storage:

NR-22133 was packaged aseptically in cryovials. The product is provided on dry ice and should be stored at -20°C or colder immediately upon arrival. Repeated freeze-thaw cycles of this product should be avoided.

Functional Activity:

NR-22133 was demonstrated to be functionally active based on its reactivity with a mouse monoclonal antibody to A27L (BEI Resources NR-569).

Citation:

Acknowledgment for publications should read “The following reagent was obtained through BEI Resources, NIAID, NIH: A27L Protein from Vaccinia Virus (WR) with C-terminal Histidine Tag, Recombinant from Baculovirus, NR-22133.”

Biosafety Level: 1

Appropriate safety procedures should always be used with this material. Laboratory safety is discussed in the following publication: U.S. Department of Health and Human Services, Public Health Service, Centers for Disease Control and

Prevention, and National Institutes of Health. [Biosafety in Microbiological and Biomedical Laboratories](#). 6th ed. Washington, DC: U.S. Government Printing Office, 2020.

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References:

- Rodriguez, J. F. and M. Esteban. “Mapping and Nucleotide Sequence of the Vaccinia Virus Gene that Encodes a 14-Kilodalton Fusion Protein.” *J. Virol.* 61 (1987): 3550–3554. PubMed: 2822962.
- Amegadzie, B. Y., B. Y. Ahn, and B. Moss. “Identification, Sequence, and Expression of the Gene Encoding a Mr 35,000 Subunit of the Vaccinia Virus DNA-Dependent RNA Polymerase.” *J. Biol. Chem.* 266 (1991): 13712–13718. PubMed: 1856205.
- Chung, C. S. et al. “A27L Protein Mediates Vaccinia Virus Interaction with Cell Surface Heparan Sulfate.” *J. Virol.* 72 (1998): 1577-1585. PubMed: 9445060.
- Ward, B. M. “Visualization and Characterization of the Intracellular Movement of Vaccinia Virus Intracellular Mature Virions.” *J. Virol.* 79 (2005): 4755–4763. PubMed: 15795261.

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Figure 1: Predicted Protein Sequence

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1  DPMDGTLFPG  DDDLAI PATE  FFSTKAAK KP  EAKREAI VKA  DEDDNEETLK
51  QRLTNLEK KI  TNVTTKFE QI  EKCKRNDE V  LFRLENHA ET  LRAAMISLAK
101 KIDVQTGR RP  YEHHHHHH
    
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VAC-WR A27L protein – Residues 3 to 112 [represents all 110 amino acid residues of the A27L protein from Vaccinia Virus (WR)
 (GenPept: [P11258](#))
 Plasmid-derived amino acids- Residues 1 and 2
 Hexa-histidine tag – Residues 113 to 118