

Zaire Ebolavirus, Mayinga, Infected Cell Lysate, Gamma-Irradiated

Catalog No. NR-49809

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Contributor and Manufacturer:

World Reference Center for Emerging Viruses and Arboviruses, University of Texas Medical Branch, Galveston, Texas, USA, under government contract

Product Description:

A crude preparation of Vero E6 cells infected with Zaire ebolavirus, Mayinga^{1,2} was gamma-irradiated (5 x 10⁶ RADs) on dry ice.

NR-49809 was tested for residual virus following the procedure described by Towner et al.³ No residual virus was recovered.

Material Provided:

Each vial contains approximately 0.5 mL of irradiated infected cell lysate and supernatant from Vero E6 cells infected with Zaire ebolavirus, Mayinga and supplemented with 2% heat-inactivated fetal bovine serum and 0.01 M Tris-HCl (pH 8.5).

Packaging/Storage:

NR-49809 was packaged aseptically, in screw-capped plastic cryovials. The product is provided frozen and should be stored at -70°C or colder immediately upon arrival. Freeze-thaw cycles should be avoided.

Citation:

Acknowledgment for publications should read “The following reagent was obtained through BEI Resources, NIAID, NIH: Zaire Ebolavirus, Mayinga, Infected Cell Lysate, Gamma-Irradiated, NR-49809.”

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. 5th ed. Washington, DC: U.S. Government Printing Office, 2009; see www.cdc.gov/biosafety/publications/bmbl5/index.htm.

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References:

1. McCormick, J. B., et al. “Biologic Differences Between Strains of Ebola Virus from Zaire and Sudan.” J. Infect. Dis. 147 (1983): 264-267. PubMed: 6827142.
2. Sanchez, A., et al. “The Virion Glycoproteins of Ebola Viruses are Encoded in Two Reading Frames and Are Expressed Through Transcriptional Editing.” Proc. Natl. Acad. Sci. USA 93 (1996): 3602-3607. PubMed: 8622982.
3. Towner, J. S., et al. “High-Throughput Molecular Detection of Hemorrhagic Fever Virus Threats with Applications for Outbreak Settings.” J. Infect. Dis. 196 Suppl. 2 (2007) S205-S212. PubMed: 17940951.

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