biei resources

SUPPORTING INFECTIOUS DISEASE RESEARCH

Fluzone[®] Influenza Virus Vaccine, 2009-2010 Formula

Catalog No. NR-19878

This reagent is the property of the U.S. Government.

For research use only. Not for human use.

Contributor:

National Institutes of Allergy and Infectious Diseases (NIAID), National Institutes of Health (NIH)

Manufacturer:

Sanofi Pasteur, Inc., Swiftwater, Pennsylvania, U.S.A.

Product Description:

NR-19878 is a split-virus vaccine prepared from influenza A/Brisbane/59/2007 IVR-148 viruses (H1N1), A/Uruguay/716/2007 NYMC X-175C (H3N2) [an A/Brisbane/10/2007 (H3N2)-like virus] and B/Brisbane/60/2008 that were predicted to circulate in the United States during the 2009 to 2010 influenza season.¹

NR-19878 was prepared from the extra-embryonic fluid of embryonated chicken eggs. The fluid containing the virus was harvested and inactivated with formaldehyde. The inactivated virus was concentrated and purified in a linear sucrose density gradient solution using a continuous flow centrifuge. The virus was then chemically disrupted using a nonionic surfactant (Triton[®] X-100) producing a split-virus. The split-virus was further purified by chemical means and suspended in sodium phosphate-buffered isotonic sodium chloride solution. Gelatin 0.05% was added as a stabilizer.²

The hemagglutinin content was standardized according to U.S. Public Health Service requirements. Each 0.25 mL syringe contains the recommended ratio of 7.5 μ g each of the hemagglutinin antigens from influenza viruses A/Brisbane/59/2007 IVR-148 (H1N1), A/Uruguay/716/2007 NYMC X-175C (H3N2) and B/Brisbane/60/2008.²

Please note that this vaccine preparation is being released <u>for</u> research use only and not for human use. Vaccines produced for the 2009-2010 season are now past their expiration dates.

Material Provided:

Each syringe contains 0.25 mL of split-virus vaccine in phosphate-buffered saline containing 0.05% gelatin. The product was formulated without preservatives.

Packaging/Storage:

NR-19878 contains a pre-filled 0.25 mL syringe. The product is provided on refrigerated bricks and should be stored at 2°C to 8°C immediately upon arrival. Do not freeze.

Citation:

Acknowledgment for publications should read "The following reagent was obtained through the NIH Biodefense and Emerging Infections Research Resources Repository, NIAID, NIH: Fluzone[®] Influenza Virus Vaccine, 2009-2010 Formula, NR-19878."

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. <u>Biosafety</u> in <u>Microbiological and Biomedical Laboratories</u>. 5th ed. Washington, DC: U.S. Government Printing Office, 2007; see www.cdc.gov/od/ohs/biosfty/bmbl5/bmbl5toc.htm.

Disclaimers:

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References:

- Fiore, A. E., et al. "Prevention and Control of Seasonal Influenza with Vaccines: Recommendations of the Advisory Committee on Immunization Practices (ACIP), 2009." <u>MMWR Recomm. Rep.</u> 58 (2009): 1-52. PubMed: 19644442.
- 2. <u>http://www.fda.gov/BiologicsBloodVaccines/Vaccines/ApprovedProducts/ucm112854.htm</u>

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