

FLUARIX® Influenza Virus Vaccine, 2010-2011 Formula

Catalog No. NR-31799

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For research use only. Not for human use.

Contributor:

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

Manufacturer:

GlaxoSmithKline Biologicals, Dresden, Germany

Product Description:

NR-31799 is a split virus vaccine prepared from influenza viruses A/California/07/2009 NYMC X-181 (H1N1)pdm09, A/Victoria/210/2009 NYMC X-187 (H3N2) (an A/Perth/16/2009-like virus) and B/Brisbane/60/2008 that were predicted to circulate in the United States during the 2010 to 2011 influenza season.

NR-31799 was prepared from influenza viruses propagated in embryonated chicken eggs. Each of the influenza viruses was produced and purified separately. After harvesting the virus-containing fluids, each influenza virus was concentrated and purified by zonal centrifugation using a linear sucrose density gradient solution containing detergent to disrupt the viruses. Following dilution, the vaccine was further purified by diafiltration. Each influenza virus solution was inactivated by the consecutive effects of sodium deoxycholate and formaldehyde leading to the production of a "split virus." Each inactivated split virus was then suspended in sodium phosphate-buffered isotonic sodium chloride solution. The vaccine was formulated from the three split virus solutions.

The hemagglutinin content was standardized according to U.S. Public Health Service requirements. Each 0.5 mL of vaccine contains the recommended ratio of 15 µg of the hemagglutinin antigen for each of the three influenza strains recommended for the 2010-2011 Northern Hemisphere influenza season: A/California/07/2009 NYMC X-181 (H1N1)pdm09, A/Victoria/210/2009 NYMC X-187 (H3N2) (an A/Perth/16/2009-like virus) and B/Brisbane/60/2008.^{1,2}

Please note that this vaccine preparation is being released for research use only and not for human use. Vaccines produced for the 2010-2011 season are now past their expiration dates.

Material Provided:

Each syringe contains 0.5 mL of split virus vaccine in sodium phosphate-buffered isotonic sodium chloride solution. The product is preservative-free and does not contain thimerosal.

Packaging/Storage:

NR-31799 is packaged in a 0.5 mL prefilled 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 BEI Resources, NIAID, NIH: FLUARIX® Influenza Virus Vaccine, 2010-2011 Formula, NR-31799."

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/bmb15/index.htm.

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References:

1. Fiore, A. E., et al. "Prevention and Control of Seasonal Influenza with Vaccines: Recommendations of the Advisory Committee on Immunization Practices (ACIP), 2010." MMWR Recomm. Rep. 59 (2010): 1-62. PubMed: 20689501.
2. <http://www.fda.gov/BiologicsBloodVaccines/Vaccines/ApprovedProducts/ucm112850.htm>

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