

Flulaval[®] Influenza Virus Vaccine, Suspension for Intramuscular Injection, 2008-2009 Formula

Catalog No. NR-17422

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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 and Distributor:

ID Biomedical Corporation of Quebec and GlaxoSmithKline

Product Description:

NR-17422 is a trivalent, split-virion, inactivated influenza virus vaccine that contains the hemagglutinin and neuraminidase surface antigens from an influenza A/Brisbane/59/07 (H1N1)-like virus (A/Brisbane/59/07 IVR-148), an A/Brisbane/10/07 (H3N2)-like virus (A/Uruguay/716/07 NYMC X-175C) and the B/Florida/4/06 virus that were predicted to circulate in the United States during the 2008 to 2009 influenza season.¹

NR-17422 was prepared from virus propagated in the allantoic cavity of embryonated hens' eggs. Each of the influenza virus strains was produced and purified separately. The virus was inactivated with ultraviolet light treatment followed by formaldehyde treatment and then purified by centrifugation and disrupted with sodium deoxycholate. Thimerosal, a mercury derivative, was added as a preservative. Each 0.5 mL dose (10 doses per vial) contains 25 µg of mercury and may also contain residual amounts of eggs proteins, formaldehyde and sodium deoxycholate. Antibiotics were not used in the manufacture of this vaccine.

The hemagglutinin content was standardized according to U.S. Public Health Service requirements. Each 0.5 mL dose contains the recommended ratio of 15 µg each of the hemagglutinin antigens from influenza viruses A/Brisbane/59/07 IVR-148 (H1N1), A/Uruguay/716/07 NYMC X-175C (H3N2) and B/Florida/4/06.

Please note that this vaccine preparation is being released for research use only and not for human use. Vaccines produced for the 2008-2009 season are now past their expiration dates.

Material Provided:

Each vial contains 5 mL (10 doses) of purified surface antigen vaccine in phosphate-buffered saline containing 25 µg of mercury.

Packaging/Storage:

NR-17422 is packaged in a 5 mL multi-dose vial. 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: Flulaval[®] Influenza Virus Vaccine, Suspension for Intramuscular Injection, 2008-2009 Formula, NR-17422."

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, 2007; see www.cdc.gov/od/ohs/biosfty/bmb15/bmb15toc.htm.

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References:

1. Fiore, A. E., et al. "Prevention and Control of Influenza: Recommendations of the Advisory Committee on Immunization Practices (ACIP), 2008." MMWR Recomm. Rep. 57 (2008): 1-60. PubMed: 18685555.

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