

## Influenza A (H1N1) 2009 Monovalent Vaccine

### Catalog No. NR-20347

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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:

Sanofi Pasteur, Inc.

#### Product Description:

Influenza A (H1N1) 2009 (pdm09) Monovalent Vaccine is an inactivated influenza virus vaccine prepared from influenza viruses propagated in embryonated chicken eggs. The virus-containing allantoic fluid was harvested and inactivated with formaldehyde. The 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 non-ionic surfactant, polyethylene glycol p-isooctylphenyl ether (Triton<sup>®</sup> X-100), producing a "split virus". The split virus was further purified and suspended in sodium phosphate-buffered isotonic sodium chloride solution. Gelatin (0.05%) was added as a stabilizer. Each 0.5 mL dose may contain residual amounts of formaldehyde ( $\leq 100$   $\mu$ g), Triton<sup>®</sup> X-100 ( $\leq 0.02\%$ ) and sucrose ( $\leq 2.0\%$ ). Each 0.5 mL dose also contains thimerosal, a mercury derivative, added as a preservative (25  $\mu$ g mercury).

The hemagglutinin content was standardized according to U.S. Public Health Service requirements. Each 0.5 mL dose contains the recommended 15  $\mu$ g of hemagglutinin antigen.

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

#### Material Provided:

Each vial contains 5 mL of split-virus vaccine in phosphate-buffered saline containing 0.05% gelatin and thimerosal (25  $\mu$ g mercury).

#### Packaging/Storage:

NR-20347 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 BEI Resources, NIAID, NIH: Influenza A (H1N1) 2009 Monovalent Vaccine, NR-20347."

#### 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](http://www.cdc.gov/biosafety/publications/bmbl5/index.htm).

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#### References:

- Morse, D., et al. "Use of Influenza A (H1N1) 2009 Monovalent Vaccine: Recommendations of the Advisory Committee on Immunization Practices (ACIP), 2009." MMWR Recomm. Rep. 58 (2009): 1-8. PubMed: [19713882](https://pubmed.ncbi.nlm.nih.gov/19713882/).
- Fiore, A. E., et al. "Prevention and Control of Influenza with Vaccines: Recommendations of the Advisory Committee on Immunization Practices (ACIP), 2010." MMWR Recomm. Rep. 59 (2010): 1-62. PubMed: [20689501](https://pubmed.ncbi.nlm.nih.gov/20689501/).

3. [WHO Recommendations for Influenza Vaccines](#)

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