







Thank you for choosing to immunize with the • Sanofi Influenza Vaccine Portfolio •

It is our ongoing commitment to simplify your vaccine choices for your eligible patients.

2025-2026
Season

	Packaging	Ages ¹⁻³	Presentation ¹⁻³	Dosing Schedule ¹⁻⁴	CPT ^{®a} Codes & Coverage ⁵	NDC ^b Information ⁶		EMR ^c Code
						Package NDC	Unit NDC	
		65 years of age and older	0.5-mL single-dose pre-filled syringe	1 dose	90662 Covered by most health plans and Medicare Part B	49281-125-65	49281-125-88	
		9 years of age and older	0.5-mL single-dose pre-filled syringe	1 dose	90673 Covered by most health plans and Medicare Part B	49281-725-10	49281-725-88	
		6 months of age and older	0.5-mL single-dose pre-filled syringe	1 or 2 doses Please refer to the Prescribing Information for further information on dosing schedule.	90656 Covered by most health plans and Medicare Part B	49281-425-50	49281-425-88	
			5-mL multi-dose vial Maximum of 10 doses can be withdrawn					90657 (0.25-mL dose) 90658 (0.5-mL dose) Covered by most health plans and Medicare Part B

^aCPT (Current Procedural Terminology) is a registered trademark of the American Medical Association. ^bNDC=National Drug Code. ^cEMR=electronic medical record. ^dACIP=Advisory Committee on Immunization Practices.

Notice: This coding guide is provided for informational purposes only and does not constitute legal or reimbursement advice. It is not intended to substitute for the physician's independent diagnosis or treatment of each patient. The information contained herein is gathered from various resources and is subject to change. Providers are solely responsible for the accuracy of all coding and claims submitted for reimbursement to any third-party payer.

Professional Indication:

Fluzone, Flublok, and Fluzone High-Dose are vaccines indicated for active immunization for the prevention of disease caused by influenza A virus subtypes and type B virus contained in (or in the case of Flublok, represented by antigens contained in) the vaccine. Fluzone is approved for use in persons 6 months of age and older. Flublok is approved for use in individuals 9 years of age and older. Fluzone High-Dose is approved for use in persons 65 years of age and older.

IMPORTANT SAFETY INFORMATION FOR FLUZONE® (INFLUENZA VACCINE), FLUBLOK® (INFLUENZA VACCINE), AND FLUZONE® HIGH-DOSE (INFLUENZA VACCINE)

Do not administer Fluzone, Flublok, or Fluzone High-Dose to anyone with a history of a severe allergic reaction (e.g., anaphylaxis) to any component of the vaccine (including egg protein for Fluzone and Fluzone High-Dose). Fluzone and Fluzone High-Dose should not be administered to anyone who has had a severe allergic reaction after previous dose of any influenza vaccine.

Appropriate medical treatment must be immediately available to manage potential anaphylactic reactions following administration of the vaccine.

If Guillain-Barré syndrome has occurred within 6 weeks following previous influenza vaccination, the decision to give Fluzone, Flublok, or Fluzone High-Dose should be based on careful consideration of the potential benefits and risks.

If Fluzone, Flublok, or Fluzone High-Dose are administered to immunocompromised persons, including those receiving immunosuppressive therapy, the expected immune response may not be attained.

Vaccination with Fluzone, Flublok, or Fluzone High-Dose may not protect all recipients.

Syncope (fainting) has been reported following vaccination with Fluzone, Flublok and Fluzone High-Dose. Procedures should be in place to avoid injury from fainting.

For Fluzone, in children 6 months through 8 years of age, the most common injection-site adverse reactions were pain or tenderness and redness; the most common solicited systemic adverse reactions were irritability, drowsiness (6 months through 35 months), and myalgia (3 years through 8 years). In adults 18 through 64 years of age, the most common injection-site adverse reaction was pain; the most common solicited systemic adverse reactions were headache and myalgia. In adults over 65 years of age, the most common injection-site adverse reaction was pain; the most common solicited systemic adverse reactions were headache, myalgia, and malaise.

For Flublok, in children 9 through 17 years of age who received Flublok Quadrivalent, the most common solicited injection-site adverse reaction was pain; the most common solicited systemic adverse reactions were myalgia, headache, and malaise. In adults 18 through 64 years of age who received Flublok, the most common injection site adverse reaction was pain; the most common solicited systemic adverse reactions were headache, fatigue, and myalgia. In adults 65 years of age and older who received Flublok, the most common injection-site adverse reaction was pain; the most common solicited systemic adverse reactions were fatigue and headache.

For Fluzone High-Dose, in adults 65 years of age and older, the most common injection-site reaction was pain; the most common solicited systemic adverse reactions were myalgia, malaise, and headache.

For Fluzone, Flublok, and Fluzone High-Dose, other adverse reactions may occur.

Please click to access or refer to the accompanying Prescribing Information for Fluzone, Flublok, or Fluzone High-Dose.

References: 1. Fluzone High-Dose. Prescribing Information. Sanofi Pasteur Inc. 2. Flublok. Prescribing Information. Protein Sciences Corporation. 3. Fluzone. Prescribing Information. Sanofi Pasteur Inc. 4. Grohskopf LA, Blanton LH, Ferdinands JM, et al. Prevention and control of seasonal influenza with vaccines: recommendations of the Advisory Committee on Immunization Practices — United States, 2024-2025 influenza season. *MMWR Recomm Rep.* 2022;71(1):1-28. 5. Seasonal Influenza Vaccines Pricing. Centers for Medicare & Medicaid Services. January 04, 2023. Accessed June 9, 2025. <https://www.cms.gov/medicare/payment/part-b-drugs/vaccine-pricing>. 6. Data on file. Sanofi 2023.