## FLUCELVAX® QUADRIVALENT



# (Influenza Vaccine) Coding and Billing

As a cell-based influenza vaccine, FLUCELVAX QUADRIVALENT has a unique Current Procedural Terminology (CPT) code.

### Code for the FLUCELVAX QUADRIVALENT vaccine administered

2023-2024 NDC Carton <sup>1</sup>	2023-2024 NDC Unit-of-Use <sup>1</sup>	Presentation <sup>1</sup>	Product Billing CPT Code <sup>2</sup>	Description <sup>1</sup>	CVX Code*2	MVX Code
70461-323-03	70461-323-04	0.5-mL pre-filled syringe	90674	Influenza virus vaccine, quadrivalent (ccIIV4), derived from cell cultures, subunit, contains no preservative or antibiotics, 0.5-mL dosage, for intramuscular use	171	SEQ
70461-423-10	70461-423-11	5-mL multi-dose vial	90756	Influenza virus vaccine, quadrivalent (ccIIV4), derived from cell cultures, subunit, contains no antibiotics, 0.5-mL dosage for intramuscular use	186	SEQ

<sup>\*</sup>CVX=vaccine administered code indicates which product was used and is used in combination with the manufacturer (MVX) code. NDC=National Drug Code

### Code for the administration of FLUCELVAX QUADRIVALENT

Report the appropriate administration code in addition to the CPT code for FLUCELVAX QUADRIVALENT.<sup>2</sup> Note: Medicare (and some other payers) requires use of the Healthcare Common Procedure Coding System (HCPCS) code, G0008, for administration of preventive vaccines, including influenza, regardless of age. Other payers use the appropriate CPT code based on age and counseling provided.<sup>3</sup>

### Include the appropriate International Classification of Diseases, Tenth Revision (ICD-10) diagnosis code

Report the ICD-10 diagnosis code, Z23, indicating an encounter for vaccine administration. The ICD-10 diagnosis code should be linked to both the vaccine and the administration code.<sup>3</sup>

### Determine if modifier 25 is appropriate

When FLUCELVAX QUADRIVALENT is administered on the same date as a significant and separately identifiable Evaluation and Management (E/M) visit, apply modifier 25 to the E/M CPT code, denoting a "significant and separately identifiable" service from the vaccine and vaccine administration service.<sup>4</sup>

CPT Code⁵	Description
90460	Immunization administration through 18 years of age (via any route of administration) with counseling by physician or other qualified healthcare professional; first or only component of each vaccine or toxoid administered
90461 (add-on code)	Immunization administration through 18 years of age via any route of administration, with counseling by physician or other qualified healthcare professional; each additional vaccine or toxoid component administered (list separately in addition to code for primary procedure)
90471	Immunization administration (includes percutaneous, intradermal, subcutaneous, or intramuscular injections); 1 vaccine (single or combination vaccine/toxoid)  NOTE: Report this code for immunization administration of any vaccine that is not accompanied by face-to-face physician or other qualified healthcare professional counseling the patient and/or family, or for patients over 18 years of age.
HCPCS Code <sup>3</sup>	Description
G0008	Seasonal influenza virus vaccine administration
ICD-10 Code <sup>3</sup>	Description
Z23	Encounter for immunization

Visit flu360.com for additional resources and information.

Please see Important Safety Information on next page, and the <u>full US Prescribing Information</u> for FLUCELVAX QUADRIVALENT.

For US Healthcare Professional Use Only This information does not constitute a guarantee or warranty of coverage benefits or reimbursement. Questions?



Call flu360 Customer Service (855) 358-8966, option 2

Note: Some payers may require use of NDCs. If so, determine if the payer requires the carton NDC or the unit-of-use NDC, and then determine if the payer requires the 10-digit or 11-digit format. If 11-digit, add a leading zero to the middle section of numbers.

# FLUCELVAX® QUADRIVALENT (Influenza Vaccine) INDICATION AND IMPORTANT SAFETY INFORMATION



### **INDICATION AND USAGE**

FLUCELVAX QUADRIVALENT is an inactivated vaccine indicated for active immunization for the prevention of influenza disease caused by influenza virus subtypes A and types B contained in the vaccine. FLUCELVAX QUADRIVALENT is approved for use in persons 6 months of age and older.

### **IMPORTANT SAFETY INFORMATION**

#### CONTRAINDICATIONS

Do not administer FLUCELVAX QUADRIVALENT to anyone with a history of severe allergic reactions (e.g. anaphylaxis) to any component of the vaccine.

#### WARNINGS AND PRECAUTIONS

If Guillain-Barré syndrome has occurred within 6 weeks of receipt of a prior influenza vaccine, the decision to give FLUCELVAX QUADRIVALENT should be based on careful consideration of the potential benefits and risks.

Appropriate medical treatment and supervision must be available to manage possible anaphylactic reactions following administration of the vaccine.

Syncope (fainting) can occur in association with administration of injectable vaccines, including FLUCELVAX QUADRIVALENT. Syncope can be accompanied by transient neurological signs such as visual disturbance, paresthesia, and tonic-clonic limb movements. Procedures should be in place to avoid falling injury and to restore cerebral perfusion following syncope by maintaining a supine or Trendelenburg position.

After vaccination with FLUCELVAX QUADRIVALENT, immunocompromised individuals, including those receiving immunosuppressive therapy, may have a reduced immune response

Vaccination with FLUCELVAX QUADRIVALENT may not protect all vaccine recipients against influenza disease.

### **ADVERSE REACTIONS**

In children 6 months through 3 years of age who received FLUCELVAX QUADRIVALENT, the most commonly reported

injection-site adverse reactions were tenderness (27.9%), erythema (25.8%), induration (17.3%) and ecchymosis (10.7%). The most common systemic adverse reactions were irritability (27.9%), sleepiness (26.9%), diarrhea (17.9%) and change of eating habits (17.4%).

In children 2 through 8 years of age who received FLUCELVAX QUADRIVALENT, the most commonly reported injection-site adverse reactions were tenderness (28.7%), pain (27.9%) and erythema (21.3%), induration (14.9%) and ecchymosis (10.0%). The most common systemic adverse reactions were sleepiness (14.9%), headache (13.8%), fatigue (13.8%), irritability (13.8%) and loss of appetite (10.6%).

In children and adolescents 9 through 17 years of age who received FLUCELVAX QUADRIVALENT, the most commonly reported injection-site adverse reactions were injection site pain (21.7%), erythema (17.2%) and induration (10.5%). The most common systemic adverse reactions were headache (18.1%) and fatigue (17.0%).

In adults 18 through 64 years of age who received FLUCELVAX QUADRIVALENT, the most commonly reported injection-site adverse reactions were pain (45.4%), erythema (13.4%) and induration (11.6%). The most common systemic adverse reactions were headache (18.7%), fatigue (17.8%) and myalgia (15.4%).

In adults ≥65 years of age who received FLUCELVAX QUADRIVALENT, the most commonly reported injection-site adverse reactions were pain (21.6%) and erythema (11.9%).

Other adverse events may occur.

To report SUSPECTED ADVERSE REACTIONS, contact CSL Seqirus at 1-855-358-8966 or VAERS at 1-800-822-7967 or www.vaers.hhs.gov.

Before administration, please see the <u>full US Prescribing</u> Information for FLUCELVAX QUADRIVALENT.

FLUCELVAX® QUADRIVALENT is a registered trademark of Seqirus UK Limited or its affiliates.

Information on reimbursement is provided as a courtesy. Due to the rapidly changing nature of the law, Medicare payment policy, and/or reliance on information provided by outside sources, the information provided herein does not constitute a guarantee or warranty that reimbursement will be received or that the codes identified herein are or will remain applicable. This information is provided "as is" and without any other warranty or guarantee, expressed or implied, as to completeness or accuracy, or otherwise.

Providers must confirm or clarify coding and coverage from their respective payers, and are responsible for accurate reporting of products in accordance with particular payer requirements.

References: 1. FLUCELVAX QUADRIVALENT. Package insert. Seqirus Inc; 2023. 2. Centers for Disease Control and Prevention. CPT codes mapped to CVX codes.

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4. American Medical Association. Current Procedural Terminology 2023 (Professional Edition). American Medical Association; 2022. 5. American Academy of Pediatrics.

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