

PRACTICE ALERT

Influenza vaccine update, 2021-22

Give the vaccine by the end of October. The CDC states that COVID-19 vaccines can be co-administered with influenza vaccines, but reactogenicity is possible.

During the 2020-2021 influenza season, fewer cases of influenza were reported than in any previous year since 1997, when data were first recorded.¹ FIGURE 1² shows the dramatic decline in the number of influenza-positive clinical samples reported to the Centers for Disease Control and Prevention (CDC) during the 2020-2021 influenza season compared with the 2019-2020 season. There was only one pediatric death attributed to influenza in 2020-2021, compared with a mean of 177 per year in the previous 3 seasons.

Deaths attributed to pneumonia and influenza were recorded over a recent 5-year period, with COVID-19 added in early mid-2020 (FIGURE 2).¹ Total cumulative deaths for 2020-2021 were extremely high, mostly due to COVID-19. Of the relatively few influenza cases last season, 37.5% were caused by influenza A and 62.5% by influenza B. The extremely low incidence of influenza precludes determining influenza vaccine effectiveness for last season.¹

In addition, other common respiratory pathogens—parainfluenza, adenoviruses, rhinoviruses, enteroviruses, and common coronaviruses—circulated last winter at historic lows.³ All of these historic lows can be attributed to the measures taken to mitigate the effect of the COVID-19 pandemic, including masks, social distancing, closure of certain venues that normally attract large crowds, and the closure of schools with a resulting increase in schooling at home. With the anticipated relaxation of these measures in 2021-2022, we can expect more influenza and other respiratory ailments due to common pathogens.

Updates to influenza vaccine recommendations

At its June 2021 meeting, the Advisory Committee on Immunization Practices (ACIP) approved the influenza vaccine recommendations for the 2021-2022 season.⁴ The central recommendation is unchanged: Everyone \geq 6 months of age should receive a vaccine unless they have a contraindication. Updates to the previous recommendations include the content of the 2021 vaccines, the specific vaccines that will be available for different age groups, the timing of vaccine administration, advice on co-administration with COVID-19 vaccines, and the list of contraindications and precautions based on vaccine type.⁴

Viral composition of US vaccines for the 2021-22 season

The antigens that will be included in the 2021-2022 influenza vaccines are listed in **TABLE 1.**⁴ The B strains are the same as last year; the A strains have been updated. The H3N2 strain is the same in all vaccines, but the H1N1 strain differs based on whether the vaccine is egg based or non-egg based. The advantage of non-egg-based vaccines is that the production process does not take as long and can be delayed in an attempt to better reflect the influenza stains in worldwide circulation.

The influenza vaccines expected to be available for the 2021-22 season

TABLE 2⁴ lists the influenza vaccines approved for use in the United States and the ages for which they are approved.⁴ All products for 2021-2022 will be quadrivalent, containing 2 type-A and 2 type-B antigens. The only

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FIGURE 1

Far fewer influenza-positive tests were reported in the 2020-21 season than during the 2019-20 season²



FIGURE 2 US mortality rates for COVID-19 have far exceeded those for pneumonia and influenza¹

Five-year data reported through week ending July 24, 2021



PIC, pneumonia, influenza, and/or COVID-19.

change in age indications is that cell culture-based inactivated influenza vaccine (ccIIV4) (Flucelvax Quadrivalent) is now approved for use starting at age 2 years; previously it was approved starting at age 4 years.⁴

Timing of vaccination

The onsets and peaks of influenza disease occur at different times each year and can also vary by geographic location. An analysis of 36 influenza seasons starting in 1982 showed that peak activity occurred most frequently in February (15 seasons), followed by December (7 seasons), and January and March (6 seasons each).⁵ Only once did peak activity occur in October and once in November. This information, along with observational studies showing the waning of influenza vaccine effectiveness after 5 to 6 months, especially in the elderly, informed the ACIP decision to modify their recommendation on the timing of vaccination. The recommendation now states that vaccine should be administered by the end of October and that July and August would have been too early, especially for older adults.

Children ages 6 months through 8 years who have not been vaccinated previously require 2 doses separated by at least 4 weeks, and the first dose should be administered early enough to allow for the second by the end of October.⁴ Children who require only 1 dose can also receive the vaccine as soon as it is available, as there is less evidence that vaccine effectiveness wanes in children.

Earlier administration is also recommended for pregnant women in their third trimester. Delaying vaccination in this group could result in postpartum administration of the vaccine, thereby depriving infants of protection against influenza illness during their first 6 months after birth.⁴

Co-administration of influenza and COVID-19 vaccines

Current guidance from the CDC states that COVID-19 vaccines can be co-administered with other vaccines including influenza vaccines.⁶ However, there are no data by which to judge the efficacy of each vaccine in coadministration or the potential for increased adverse reactions. ACIP advises caution on 2 points: (1) physicians should watch for updated guidance as more information becomes available, and (2) there is the potential for increased reactogenicity after co-administration, especially with the more reactogenic influenza vaccines: adjuvanted inactivated influenza vaccine (aIIV4) and high-dose inactivated influenza

TABLE 1

2021-2022 influenza vaccine composition⁴

Egg-based	IIV4s and	LAIV4
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A/Victoria/2570/2019 (H1N1)pdm09-like virus ^a				
A/Cambodia/e0826360/2020 (H3N2)-like virusª				
B/Washington/02/2019 (Victoria lineage)-like virus				
B/Phuket/3073/2013 (Yamagata lineage)-like virus				
Cell-culture-based IIV4 and RIV4				
A/Wisconsin/588/2019 (H1N1)pdm09-like virusª				
A/Cambodia/e0826360/2020 (H3N2)-like virusª				
B/Washington/02/2019 (Victoria lineage)-like virus				
B/Phuket/3073/2013 (Yamagata lineage)-like virus				

IIV4, quadrivalent inactivated influenza vaccine; LAIV4, quadrivalent live-attenuated influenza vaccine; RIV4, quadrivalent recombinant influenza vaccine. ^a Updated from 2020-2021 vaccine.

vaccine (HD-IIV4). Moreover, these vaccines and the co-administered COVID-19 vaccine should be injected into different limbs.

Contraindications and precautions

Serious allergic reactions to influenza vaccines are rare—about 1.3 incidents per million doses administered.⁷ However, a previous severe allergic reaction to a particular vaccine or to any component of the vaccine is a con-

TABLE 2

Influenza vaccines expected to be available in the United States for the 2021-2022 flu season⁴

All vaccines listed are quadrivalent.

Vaccine type	0-6 mo	6-23 mo	2-17 у	18-49 y	50-64 y	≥ 65 y	
Standard-dose, unadjuvanted inactivated (IIV4)	N/A	Afluria Quadrivalent					
		Fluarix Quadrivalent					
		FluLaval Quadrivalent					
		Fluzone Quadrivalent					
Cell culture-based inactivated (IIV4)	N/A			Flucelvax Quadrivalent			
Adjuvanted inactivated (allV4)	N/A					Fluad Quadrivalent	
High-dose inactivated (HD-IIV4)	N/A					Fluzone High-dose Quadrivalent	
Recombinant (RIV4)		N/A		Flublok Quadriv		alent	
Live attenuated (LAIV4)	Ν	I/A	FluMist Quadrivalent		N/A		

Egg based Non-egg based

IIV4, quadrivalent inactivated influenza vaccine; LAIV4, quadrivalent live attenuated influenza vaccine; N/A, not approved for age group; RIV4, quadrivalent recombinant influenza vaccine.

traindication for use of that vaccine. In addition, a history of severe allergic reaction to any influenza vaccine is a contraindication for all egg-based vaccines.

There are 2 precautions for all influenza vaccines: a concurrent moderate or severe acute illness (with or without fever), and a history of Guillain-Barré syndrome within 6 weeks of receiving any influenza vaccine. An additional precaution for ccIIV4 and recombinant influenza vaccine (RIV4) is a history of severe allergic reaction after administration of any other influenza vaccine. Administration of RIV4 or ccIIV4 to someone with such a history should occur in a medical setting and be supervised by someone who can recognize and treat a severe reaction.

Live attenuated influenza vaccine (LAIV) continues to have a considerably longer list of contraindications, which can be found in the published recommendations for 2021-2022.⁸

Final advice

The upcoming influenza season has the potential to be clinically challenging with the possibility of co-existing high rates of both COVID-19 and influenza. Recommend both influenza and COVID-19 vaccination to patients. Also, be sure to encourage and practice other preventive measures such as masking in crowds, frequent hand washing, isolation when sick, respiratory hygiene, and (for physicians) selected prescribing of influenza antiviral medications and meticulous office-based infection control practices.⁹ JFP

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SHOULDER DISLOCATION

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