Centers for Disease Control and Prevention

Immunization Update

Washington State Immunization Summit 2023

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Disclosures

- JoEllen Wolicki is a federal government employee with no financial interest in or conflict with the manufacturer of any product named in this presentation.
- I will not discuss any off-label uses for vaccines.
- The use of trade names is for identification purposes only and does not imply endorsement by the ACIP or CDC.
- The findings and conclusions in this presentation are those of the author(s) and do not necessarily represent the official position of the Centers for Disease Control and Prevention or ATSDR.

Disclosures

- The recommendations to be discussed are primarily those of the Advisory Committee on Immunization Practices (ACIP).
 - Composed of 15 experts in clinical medicine and public health
 - Provides guidance on use of vaccines and other biologic products to DHHS, CDC, and the U.S. Public Health Service



Next ACIP Meeting June 21–22, 2023

Advisory Committee on Immunization Practices (ACIP) | CDC

Thank You!

Advisory Committee on Immunization Practices 2023 Immunization Schedules

Immunization Schedules: Overview

- Published annually in February
 - Represents current, approved ACIP policy
 - Designed for implementation of ACIP policy
- Two separate schedules
 - Child and adolescent schedule (age birth through 18 years)
 - Adult schedule (age 19 years or older)

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Vaccines in the Child and Add Vaccine Cowd-19	olescent Immunizat	ion Schedule Abbreviation(s) 1vCOV-mRNA 2vCOV-mRNA 1vCOV-aPS	Trade name(s) Comirnaty/Pfize- BioNTech COVID-19 Vaccine SVIICE/WXY-Moderna COVID-19 Vaccine Prizer-BioNTech COVID-19 Vaccine, Bivalent Moderna COVID-19 Vaccine, Bivalent Novaxox COVID-19 Vaccine, Bivalent	How to use the child and adolescent immunization schedule 1 2 2 3 3 4 4 5 Review vaccine trable 1) 9 Vectormineded up vaccination (Table 2) 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
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Pneumococcal conjugate vaccine Pneumococcal polysaccharide vaccine Poliovirus vaccine (inactivated) Rotavirus vaccine		PCV13 PCV15 PPSV23 IPV RV1 RV5	Prevnar 13* Vaxneuvance** Pneumovax 23* IPOL* Rotarix* RotaTeq*	Lowinea the Lux vacane Schedules app for provider at Control of the second se
Tetanus, diphtheria, and acellular pertussis va Tetanus and diphtheria vaccine	ccine	Tdap Td	Adacel Boostrix® Tenivac® Tdvax [™]	• Ceneral Best Practice Guidelines for Immunization (including contraindications and precautions): www.ccd.gov/waceins/hcp/ada/are/general-recs/index.html • Vaccine information statements: • Vaccine information statements
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AppendixRecommended Adult Immunization Schedule, United States, 2023

Guide to Contraindications and Precautions to Commonly Used Vaccines

Adapted from Table 4-1 in Advisory Committee on Immunization Practices (ACIP) General Best Practice Guidelines for Immunization: Contraindication and Precautions available at www.cdc. gov/vaccines/hcp/acip-recs/general-recs/contraindications.html and ACIP's Recommendations for the Prevention and Control of 2022-23 Seasonal Influenza with Vaccines available at www.cdc.gov/mmwr/volumes/71/rr/rr7101a1.htm

For COVID-19 vaccine contraindications and precautions see

www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html#contraindications

Vaccine	Contraindicated or Not Recommended ¹	Precautions ²
Influenza, egg-based, inactivated injectable (IIV4)	 Severe allergic reaction (e.g., anaphylaxis) after previous dose of any influenza vaccine (i.e., any egg-based IIV, ccIIV, RIV, or LAIV of any valency) Severe allergic reaction (e.g., anaphylaxis) to any vaccine component^a (excluding egg) 	 Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine Moderate or severe acute illness with or without fever
Influenza, cell culture-based inactivated injectable [(cclIV4), Flucelvax* Quadrivalent]	Severe allergic reaction (e.g., anaphylaxis) to any ccllV of any valency, or to any component ¹ of ccllV4	 Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine Persons with a history of severe allergic reaction (e.g., anaphylaxis) after a previous dose of any egg-based IIV, RIV, or LAIV of any valency. If using cclV4, administer in medical setting under supervision of health care provider who can recognize and manage severe allergic reactions. May consult an allergist. Moderate or severe acute illness with or without fever
Influenza, recombinant injectable [(RIV4), Flut Quadrivalent]	Severe allergic reaction (e.g., haphylaxis) to any RIV of any valency, or to any component ^a of RIV4	 Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine Persons with a history of severe allergic reaction (e.g., anaphylaxis) after a previous dose of any egg-based IIV, ccIIV, or LAIV of any valency. If using RIV4, administer in medical setting under supervision of health care provider who can recognize and manage severe allergic reactions. May consult an allergist. Moderate or severe acute illness with or without fever
Influenza, live attenu, ed [LAIV4, Flumist* Quadrivalent]	 Severe allergic reaction (e.g. ar phylaxis) after previous dose of any influenza vaccine (i.e., any egg-based IV, ccliv. if, or LAIV of any valency). Severe allergic reaction (e.g. maphylaxis) to any vaccine component³ (excluding egg) Anatomic or functional arcenia for any cause including, but not limited to, medications and HV. Close contacts or caregivers of severely immunosuppressed persons who require a protected environment. Pregnancy Cochlear implant Active communication between the cerebrospinal fluid (CSF) and the oropharynx, nasopharynx, nose, ear, or any other cranial CSF leak. Received influenza antiviral medications oseltamivir or zanamivir within the previous 17 days. 	 Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine Asthma in persons aged 5 years old or older Persons with underlying medical conditions (other than those listed under contraindications) that might predispose to complications after wild-type influenza virus infection [e.g., chronic pulmonary, cardiovascular (except isolated hypertension), renal, hepatic, neurologic, hematologic, or metabolic disorders (including diabetes mellitus)] Moderate or severe acute illness with or without fever
 When a contraindication is contraindications.html When a precaution is prese 	present, a vaccine should NOT be administered. Kroger A, Bahta L, Hunter P. ACIP General Bes	t Practice Guidelines for Immunization. www.cdc.gov/vaccines/hcp/acip-recs/general-recs/ on from the vaccine outweighs the risk for an adverse reaction. Kroger A, Bahta L, Hunter P.

ACIP General Best Practice Guidelines for Immunization. www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html
3. Vaccination providers should check FDA-approved prescribing information for the most complete and updated information, including contraindications, warnings, and precautions. Package inserts for U.S.licensed vaccines are available at www.fda.gov/vaccines-blood-biologics/approved-products/vaccines-licensed-use-united-states.

CDC Interim COVID-19 Immunization Schedule for Persons 6 Months of Age and Older

		COVID-19 Vac Interim COVID-19 Imm for Persons 6 Months of	cine unization So of Age and	chedule Older		L						
	Table 1b. Fo Bivalent P be used. Vaccine ty	or Most People (those who a fizer-BioNTech COVID-19 Vacc pe: mRNA		COVID-19 Vac Interim COVID-19 Immu for Persons 6 Months of	ci uniz f Aj	ne zation Sc ge and (hedule Dider					
	Age	Vaccination History	Table 1c Fo	r Most People (those who ar			COVID-19 Vaccia	ne ation Schedule				
		Unvaccinated: 0 doses	Novavax [®] (Monovalent vaccine) Type: Protein Sub-Unit			for Persons 6 Months of Age and Older						
		1 dose of bivalent vaccine	12 years	1 or more doses of monovalent Novavax vaccine		The followi guidance s Table 1a. F	following tables provide COVID-19 vaccination schedules based on age, health status, and product. For detailed Jance see <u>Interim Clinical Considerations for Use of COVID-19 Vaccines CDC</u> . Ie 1a. For Most People (those who are NOT moderately to severely immunocompromised)					
		2 doses of bivalent varcine	and older			Vaccine ty	pe: mRNA	ent moderna vaccine is no longer recomm	ended and should not be used.			
	6 months			At least 1 dose of bivalent vacci		Age	Vaccination History	Bivalent Vaccine Schedule [†]	Administer			
	4 years ^{‡§}	At least 3 doses of bivalent vac		19 Yaccine remains authoritati to previde a 3 - de 79 Yaccine remains authoritati to previde a 3 - de Madiyuari yaccine, have not received any previou COND 19 -accine, have not received any previou do thermain not receive a doas. This doas is addr to accord 5 465 - Col ⁻ 2 Mericine may consider doap and color. May receive 1 additional biolater of age and color: May receive 1 additional biolater		6 months through 5 years ^{‡9}	Unvaccinated: 0 doses	2 doses. Administer: • Dose 1 now • Dose 2 at least 4–8 weeks [§] after Dose 1	0.25 mL/25 µg from the vial with a			
		Previously vaccinated with mo	50 µg of Matrix- series using any vaccine and wou				1 dose of bivalent vaccine	1 dose. Administer: • Dose 2 at least 4–8 weeks ⁹ after Dose 1	- blue cap and gray label border			
		1 dose of monovalent vaccine	† Persons with a re ‡ Adults 65 years of				At least 2 doses of bivalent vaccine	No dose	No dose			
							Previously vaccinated with monovale	nt mRNA COVID-19 vaccine				
		2 doses of monovalent vaccine At least 1 dose of monovalent va					1 dose of monovalent vaccine	1 dose. Administer: • Dose 2 at least 4–8 weeks ¹ after Dose 1	0.25 mL/25 µg from the vial with a blue cap and gray label border.			
		and 1 dose of bivalent vaccine Unvaccinated: 0 doses					2 doses of monovalent vaccine	1 dose. Administer: • Dose 3 at least 8 weeks (2 months) after Dose 2	0.2 mL/10 µg from the vial with a dark pink cap and yellow label border			
	5 years and older [‡]	1 dose or more doses of monovalent vaccine ⁵					At least 1 dose of monovalent vaccine and 1 dose of bivalent vaccine	No dose	No dose			
		At least 1 dose of bivalent vace					Unvaccinated: 0 doses	1 dose now**	6 through 11 years:			
	 Refer to <u>CDC's in</u> Persons with a m CDC recommendation CDC recommendation waccine from the 	terim Clinical Considerations for specific guida cont SARS-Col-2 Infection may consider delay ds bivalent vaccine doses from the same manu a is recommended. In the following exception same manufacturer be used and a VAERS repc				6 years and older	1 or more doses of monovalent vaccine	1 dose. Administer: • Vaccine at least 8 weeks (2 months) after the previous dose ^{**}	0.25 μg from the vial with a blue cap and gray label border 12 years and older: 0.50 mL/50 μg from the vial with a blue cap and gray label border			
	 Same vaccine Previous dose Percon would 	not available e unknown Lotherwise pet complete the vaccientice review					At least 1 dose of bivalent vaccine	No dose**	No dose**			
2	ables 1 A – C			* Refer to <u>CCC</u> : Interm Circuit Constitutions for specific guidance on interchangeability of vaccine products for all ages. * Refer to <u>CCC</u> : Interm Circuit Constitutions may available all-physicacculation by Innomity from type products and a specific and a specifi								
C	or N	lost Pec	ple			 Person starts Children ages 6 schedule. A thir An 8-week inter 	Dut unable to complete a vaccination series with the san months through 4 years who received bivalent vaccines id dose of either mRNA vaccine (Moderna or Pitzer-BioNTe val between the first and second doses of CDVID-19 vacc	ee CUVUL-19 vacchee due to a contraindication from different manufacturers for the first 2 doses of an mRNA (ich) should be administered at least 8 weeks after the second i ines might be optimal for some people ages 6 months-64 yea	COVID-19 vaccine series should follow a 3-dose lose. rs, especially for males ages 12-39 years, as it may			

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		for Persons 6 Months	of Age and Older	_	8		
	COVID-19 Vac Interim COVID-19 Immu for Persons 6 Months o	cine unization Schedule f Age and Older				ie 1. s) after Dose 2. ⁺ 1. after Dose 2. ⁺	
					Administer	s) after Dose 2.*	0.2 mL/3 p
Table 2a. P	eople Who are Moderately to	Severely Immunocompromised			Moderna: 0.50 mL/50 ug from the	-	from the vial with a maroon c
Bivalent M Vaccine ty	Moderna COVID-19 Vaccine: Mor pe: mRNA	ovalent Moderna vaccine is no longer reco	ommended and should not be used.	ast	vial with a blue cap and gray label border.	1. after Dose 2. ⁴	
Age	Vaccination History	Bivalent Vaccine Schedule	Administer		OR Pfizer-BioNTech: 0.3 mL/30 ug	after Dose 2.4	
	Unvaccinated: 0 doses	3 doses. Administer: • Dose 1 now. • Dose 2 at least 4 weeks after Dose 1. • Dose 3 at least 4 weeks after Dose 2. [‡]	6 months through 11 years:	_	from the vial with a gray cap	i) after Dose 2.*	-
	1 dose of bivalent vaccine only	2 doses. Administer: • Dose 2 at least 4 weeks after Dose 1. • Dose 3 at least 4 weeks after Dose 2. [‡]	cap and gray label border. 12 years and older: 0.50 mL/50 µg from the vial with a blue cap and gray label border.	ages 12 y ages 18 xt availat	ears and older. Administer 0.5 mL/5 µg rS and 50 years and old who completed the primary series le) or unwilling to receive an mRNA vaccine and	e 1.	5 through 1 years:
	2 doses of bivalent vaccine	1 dose. Administer: • Dose 3 at least 4 weeks after Dose 2. [‡]		he clinica DVID-191	at teast 2 months totlowing the tast ij udgement of a healthcare provider and vaccine dose. Refer to CDC's Interim Clinical	e 1. e 2.*	
			6 months through 5 years: 0.2mL/10 μg from the vial with a dark pink cap and yellow label border‡.			2. ⁴ vial v oran	0.2 mL/10 µg from the vial with a orange ca
6 months	3 doses of bivalent vaccine	See tootnote"	6 through 11 years: 0.25 mL/25 µg from the vial with a blue cap and gray label border. 12 years and older: 0.50 mL/50 µg from the vial with a blue cap and gray label border.			e 1. e 2.º	12 years an older: 0.3 mL/30 µg from the via with a gray
older†	Previously vaccinated with mono	valent mRNA COVID-19 vaccine			previous dose.*	cap	
	1 dose of monovalent vaccine	2 doses. Administer: • Dose 2 at least 4 weeks after Dose 1. • Dose 3 at least 4 weeks after Dose 2. [†]	6 months through 11 years: 0.25 mL/25 µg from the vial with a blue cap and orav label border.			s) after Dose 3. ⁺	-
	2 doses of monovalent vaccine	1 dose. Administer: • Dose 3 at least 4 weeks after Dose 2.*	12 years and older: 0.50 mL/50 µg from the vial with a blue cap and gray label border.				
	3 doses of monovalent vaccine	1 dose. Administer: • Dose 4 at least 8 weeks after Dose 3. [‡]	6 months through 5 years: 0.2mL/10 μg from the vial with a dark pink cap and vallow balk border			be administered when FDA authoriza sunknown; or person would otherwis straindication vaccine at least 2 months following to be clinical unknownert of a healthcare	tion requires that e not complete he last
	3 doses of monovalent vaccine and 1 dose of bivalent vaccine	See footnote [‡]	6 through 11 years: 0.25 mL/25 μg from the vial with a blue cap and gray label border.			3VID-19 vaccine dose. Refer to CDC1	Interim Clinical
* Refer to CDCL1 * CDC recommer more than 1 do vaccine from th * Same vaccin * Previous door * Preson woul * Preson wou	and 1 dose of bivalent vaccine	so the control of th	V2 years and older: 0.50 mL/50 ug/from the vial with a blue cap and gray label border.	0	ns with N	/lodera	ite

COVID 10 Vaccino

COVID-19 Vaccine Interim COVID-19 Immunization Schedule for 6 Months of Age and Older (cdc.gov)

* Adults 65 years of age and older: May receive 1 additional bivalent mRNA vaccine dose at least 4 months after

Return of Vaccine-Preventable Diseases

Morbidity and Mortality Weekly Report

Wastewater Testing and Detection of Poliovirus Type 2 Genetically Linked to Virus Isolated from a Paralytic Polio Case — New York, March 9–October 11, 2022

A. Blythe Ryerson, PhD^{1,*}; Daniel Lang, MS^{2,*}; Mohammed A. Alazawi, PhD²; Milagros Neyra, MPH³; Dustin T, Hill, PhD³; Kirsten St. George, PhD^{2,4}; Meghan Fuschino, MS²; Emily Lutterloh, MD²; Bryon Backenson, MS²; Samuel Rulli⁵; Patricia Schnabel Ruppert, DO⁵; Jacqueline Lawler, MPH⁶; Nancy McGraw, MPH⁷; Andrew Knecht, DO⁸; Irina Gelman, DPM⁸; Jane R, Zucker, MD^{1,9}; Enoma Omoregie, PhD⁹; Sarah Kidd, MD¹; David E. Sugerman, MD¹; Jaume Jorba, PhD¹; Nancy Gerloff, PhD¹; Terry Fei Fan Ng, PhD¹; Adriana Lopez, MHS¹; Nina B. Masters, PhD^{1,10}; Jessica Leung, MPH¹; Cara C. Burns, PhD¹; Janell Routh, MD¹; Stephanie R, Bialek, MD¹; M. Steven Oberste, PhD^{1,†}; Eli S. Rosenberg, PhD^{2,11,†}; 2022 U.S. Poliovirus Response Team

> 'Silent' spread of polio in New York drives CDC to consider additional vaccinations for some people



CDC WARNS LARGE OUTBREAK IS POSSIBLE

Measles Exposure at a Large Gathering in Kentucky, February 2023 and Global Measles Outbreaks

Print



Distributed via the CDC Health Alert Network March 3, 2023, 11:15 AM ET CDCHAN-00488

Summary

The Centers for Disease Control and Prevention (CDC) is issuing this Health Alert Network (HAN) Health Advisory to notify clinicians and public health officials about a confirmed measles case at a large gathering. On February 24, 2023, the Kentucky Department for Public Health (KD³H) identified a confirmed case of measles in an unvaccinated individual with a history of recent international travel.

Decline in Vaccination Coverage Among Kindergarteners During the Pandemic

Estimated vaccination coverage among kindergartners by vaccine— United States,2019-20, 2020-21, and 2021–22 school years

Kindergarten Coverage	2019-20 (pre-pandemic)	2020-21 (pandemic)	2021-22 (pandemic)
MMR	95.2%	93.9%	93.5%
DTaP	94.9%	93.6%	93.1%
Polio	95.0%	93.9%	93.5%
Varicella (UTD)	94.8%	93.6%	92.6%

2% drop in kindergarten vaccination coverage since the start of the pandemic

275,000 children, who entered kindergarten during the pandemic, are susceptible to vaccine preventable disease

Seither R, Calhoun K, Yusuf OB, et al. Vaccination Coverage with Selected Vaccines and Exemption Rates Among Children in Kindergarten — United States, 2021–22 School Year. MMWR Morb Mortal Wkly Rep 2023;72:26–32. DOI: <u>http://dx.doi.org/10.15585/mmwr.mm7202a2</u>.



Ensure Everyone is Caught Up! Evidence-Based Strategies

- Use a reminder/recall system to let patients/parents know vaccines are due.
- Give a strong recommendation when talking about vaccines.
- Use provider prompts (computer or handwritten) to remind staff when vaccines are due.
- Assess for needed vaccines at every clinical visit.
- Use standing orders.
- Consider "vaccine-only" visits.

Influenza Update

Time to Prepare for Next Flu Season!

2022–2023 ACIP Recommendations: Influenza

- Annual influenza vaccination is recommended for persons 6 months of age and older without contraindications or precautions
- Note: Influenza vaccine products vary with different age-indications contraindications, and recommendations.



2022–2023 ACIP Recommendations: Influenza



Ages 6 months–64 years No preferential recommendation. Administer any licensed, recommended, and age-appropriate vaccine.



Ages 65 years and older ACIP recommends any high dose or adjuvanted influenza vaccine.

<u>Prevention and Control of Seasonal Influenza with Vaccines: Recommendations of the Advisory Committee on Immunization Practices — United States,</u> 2022–23 Influenza Season | MMWR (cdc.gov)

65 Years Old and Older: Higher Dose and Adjuvanted Vaccines

- ACIP recommends preferential use of higher dose or adjuvanted influenza vaccines for persons 65 years old or older
- Includes these vaccines:
 - Higher dose: High-dose influenza vaccine (Fluzone High-Dose),
 Recombinant Influenza Vaccine (Flublok)
 - -Adjuvanted: Adjuvanted influenza vaccine (Fluad)
 - -No preference between these three
- If none of the three are available, vaccinate with another ageappropriate influenza vaccine

<u>Prevention and Control of Seasonal Influenza with Vaccines: Recommendations of the Advisory Committee on Immunization Practices — United States,</u> <u>2022–23 Influenza Season | MMWR (cdc.gov)</u>

BIVALENT COVID-19 Vaccination Persons Who Are Not Immunocompromised

Bivalent COVID-19 Vaccination Coverage Rates Are Low

16.7% of the total U.S. population have received a bivalent COVID-19 vaccine.

20.2% of U.S. adults ages 18 years or older have received a bivalent COVID-19 vaccine.

New Recommendations for Persons^{*} 6 Years of Age or Older Who HAVE NOT Received Bivalent Vaccine



*Without immunocompromise

New Recommendations for Persons^{*} 6 Years of Age or Older Who HAVE NOT Received Bivalent Vaccine



New Recommendations for Persons^{*} 6 Years of Age and Older Who HAVE Received Bivalent Vaccine



Vaccination is complete. No additional doses are indicated at this time.

*Without immunocompromise

New Recommendations for Those at Higher Risk of Severe Disease: People 65 Years of Age or Older

People 65 years and older who have NOT received a bivalent mRNA dose



People 65 years and older who have already received a bivalent mRNA dose

Children 6 Months Through 5 Years of Age

- Schedule varies based on vaccination status
 - 1. Unvaccinated: No previous doses of any COVID-19 vaccine
 - 2. Vaccinated: Children whose immunization history includes dose(s) of monovalent vaccine
- Vaccine products have different age indications
 - Pfizer-BioNTech: 6 months through 4 years of age
 - Moderna: 6 months through 5 years of age
 - Different presentations based on recipient's vaccination history and health status

Unvaccinated Children^{*} 6 Months Through 5 Years of Age: Moderna COVID-19 Vaccine



Use the blue capped vial with the gray labeled border

*Not immunocompromised.

+An 8-week interval between doses 1 and 2 may be optimal for some people ages 6 months—64 years, especially for males ages 12—39 years, as it may reduce the small risk of myocarditis and pericarditis associated with these vaccines

Unvaccinated Children^{*} 6 Months Through 4 Years of Age: Pfizer-BioNTech COVID-19 Vaccine



*Not immunocompromised.

*An 8-week interval between doses 1 and 2 may be optimal for some people ages 6 months–64 years, especially for males ages 12–39 years, as it may reduce the small risk of myocarditis and pericarditis associated with these vaccines

Children^{*} 6 Months Through 4 Years of Age: Previously Vaccinated with Moderna Monovalent Vaccine

Vaccination	Administer	Schedule	Vial			
1 dose of monovalent Moderna vaccine	Moderna	1 dose 4–8 weeks after the previous dose	Blue cap			
2 doses of monovalent Moderna vaccine	Moderna	1 dose 8 weeks after the previous dose	Pink cap			
1 dose of monovalent AND bivalent Moderna Vaccine	No dose! Pre	No dose! Previously received bivalent vaccine				

Children^{*} 6 Months Through 4 Years of Age: Previously Vaccinated with Pfizer-BioNTech Monovalent Vaccine

Vaccination	Administer	Schedule	Vial
1 dose of monovalent Pfizer-BioNTech vaccine	Pfizer- BioNTech	2 doses. Dose 2: 3–8 weeks after monovalent dose 1. Separate Dose 2 and Dose 3 by at least 8 weeks.	Maroon cap
2 doses of monovalent Pfizer-BioNTech	Pfizer- BioNTech	1 dose at least 8 weeks after monovalent Dose 2.	Maroon cap
3 doses of monovalent Pfizer-BioNTech	Pfizer- BioNTech	1 dose at least 8 weeks after monovalent Dose 3.	Maroon cap
2 doses monovalent Pfizer-BioNTech and 1 dose bivalent Pfizer- BioNTech	No dose! F	Previously received bivalent vaccine	

Children^{*} 5 Years of Age: Previously Vaccinated with Monovalent Vaccine

Vaccination	Administer	Schedule	Vial
Unvaccinated	Moderna	2 doses separated by 4–8 weeks	Blue cap
	Pfizer-BioNTech	1 dose	Orange cap
1 dose of monovalent	Moderna	1 dose 4–8 weeks after the previous dose	Blue cap
Moderna vaccine	OR Pfizer-BioNTech	1 dose at least 8 weeks after the previous dose	Orange cap
2 doses of monovalent	Moderna		Pink cap
Moderna vaccine	OR Pfizer-BioNTech	1 dose 8 weeks after the previous dose	Orange cap
1 or more doses of monovalent Pfizer-BioNTech	Pfizer-BioNTech	1 dose at least 8 weeks after the previous dose	Orange cap
At least 1 dose bivalent of Pfizer-BioNTech (regardless of monovalent vaccine history)	No dose! Previous	sly received bivalent vaccine	

Fewer COVID-19 Vaccine Products in Your Storage Unit



Clinical Considerations for Moderna Bivalent Vaccine Vial with the Blue Cap and Gray-Bordered Label



- Ages: 6 months and older
- Dosage: Varies by age
 - 6 months through 11 years:
 0.25 mL/25 μg
 - 12 years and older: 0.5 mL/50 μg
- Use for persons never vaccinated with bivalent vaccine, including:
 - 6 months and older: Unvaccinated
 - 6 months through 5 years previously vaccinated with only 1 dose of monovalent vaccine
 - 6 years and older vaccinated with 1 or more doses of monovalent vaccine

Clinical Considerations for Moderna Bivalent Vaccine Vial with the Dark Pink Cap and Yellow Box Label

- Ages: 6 months through 5 years
- Dosage: 0.2 mL/10µg
- Route: Intramuscular injection
- Multidose vial = 2 doses
- Use for children who were previously vaccinated with 2 doses of monovalent vaccine



"Booster Doses Only" does not apply

Clinical Considerations for Pfizer-BioNTech Bivalent Vaccine



- Ages 6 months through 4 years
- 0.2 mL/3 μg
- Mix with diluent
- Unvaccinated and previously vaccinated persons

- Ages 5 through 11 years
- 0.2 mL/10 μg

- Mix with diluent
- Unvaccinated and previously vaccinated persons

- Ages 12 years and older
- 0.3 mL/30 μg
- Do NOT mix with diluent
- Unvaccinated and previously vaccinated persons





CDC COVID-19 Vaccine Clinical Resources

COVID-19 Vaccination	Use of $COVID_19$ Vaccines in the	ne United States					
Product Info by U.S. Vaccine +	Print	le officed States					
Interim Clinical Considerations —	Interim Clinical Considerations		U.S. COVID-19 Vaccine Product Information				
Use of COVID-19 Vaccines in the U.S.	Summary of recent changes (last undated March 1)	5 20231	Espanol Print				
Use of COVID-19 Vaccines in the U.S.: Appendices	 New recommendation for children ages 6 months-4 years who BioNTech primary series to receive 1 bivalent Pfizer-BioNTech 	o previously completed a 3-dose monovalent Pfizer- booster dose at least 2 months after completion of the	Find a suite of information storage and handling, safe	n and materials that are needed ety, and reporting.	d for each specific COVID-19 vacci	ne that cover administrati	
FAQs for the Interim Clinical Considerations	 Vaccination providers are now required to report cases of myc COVID-19 Vaccine to the Vaccine Adverse Event Reporting Syst 	carditis and pericarditis after receipt of a Janssen em (VAERS).	Janssen/J&J	Pfizer-	Moderna	Novavax	
Archived COVID-19 Vaccination Schedules				BioNTech			
Myocarditis and Pericarditis	Reference Materials	Get Email Updates			_		
Considerations	Summary Document for Interim Clinical Considerations (Update 13/(12/0922)	ed Receive email updates about this	Interim CC	OVID-19 Immunization	Prevaccina Chasklist	ation Screening	
linical Care +	Interim COVID-19 Immunization Schedule (Updated 12/12/2022)	page.	older	or Ages o montris and	COVID-19 Pre	evaccination Guidelines 🖪	
Provider Requirements and + Support	<u>COVID-19 Vaccination Schedule Infographic</u> <u>COVID-19 Vaccination Schedule Infographic (Immunocompron</u>	nised) Get Email Updates	Find guidance schedules ba:	e for COVID-19 vaccination sed on age and medical	Download a p multiple lang	prevaccination checklist in uages.	
raining and Education	 Special Situations for COVID-19 Vaccination of Children and Ac Age Transitions and Interchangeability. (Updated 12/09/2022) 	iolescents;	condition.		Select Lang	uage ~	
/accine Recipient Education +	<u>FAQs for the Interim Clinical Considerations</u>						
Health Departments +	COVID 10 Varians Decommon lations and Caled	Jac	Requirements, Train	nings, and Resources			
Planning & Partnerships +	COVID-19 Vaccines, Recommendations, and Schedu		Identification, Disposal, ar	nd Reporting of COVID-19	Secretarial Directive on Eligib	pility to Receive Particular	
Vaccine Effectiveness Research	Overview of COVID-19 vaccination Guid	dance for people who are not immunocompromised	vaccine Wastage		COVID-19 Vaccine Boosters	(September 25, 2021)	
COVID-19 Vaccine Data Systems +	Guid	dance for people who are immunocompromised	Vaccine Storage and Hand	dling Toolkit	Secretarial Directive on Eligit	Dility to Receive Particular October 22, 2021)	
			Training and Education				

Interim Clinical Considerations for Use of COVID-19 Vaccines | CDC U.S. COVID-19 Vaccine Product Information | CDC

Pfizer-BioNTech COVID-19 Vaccine At-A-Glance



Guidance below summarizes basic storage, preparation, scheduling, and administration for ALL Pfizer-BioNTech COVID-19 Vaccine products.



Storage and Handling Basics

Find additional guidance on storing the vaccine properly at:

- Vaccine Storage and Handling Toolkit-Updated with COVID-19 Vaccine Storage and Handling Information
- Pfizer-BioNTech COVID-19 Vaccines | FDA
 Pfizer-BioNTech COVID-19 Vaccine | cvdvaccine.com

Vial cap color	BIVALENT Maroon Cap	BIVALENT Gray Cap				
Ages	6 months through 4 years	5 through 11 years	12 years and older			
Supplied in:	MDV: 10 doses per vial Requires diluent	MDV: 10 doses per vial Requires diluent	MDV: 6 doses per vial SDV: 1 dose No diluent			
Storage Temperature: Before Puncture Do NOT store vaccine in a standard freezer	Between: -90°C and -60°C (-130°F and -76°F) until the expiration date* 2°C and 8°C (36°F and 46°F) for up to 10 weeks NOTE: The beyond-use date (10 weeks) replaces the manufacturer's expiration date but NEVER extends it. Always use the earliest date. Do NOT use vaccine if the expiration date or beyond-use date has passed.					
Thawing Frozen Vaccine Do NOT refreeze thawed vaccine	Between: 2°C and 8°C (36°F and 46°F) OR Up to 25°C (77°F) Amount of time needed to thaw vaccine varies based on temperature and number of vials.					
Storage Temperature: After 1st Puncture Do NOT use after 12 hours	age iperature: r Ist Puncture Do NOT use after 12 hours Between: 2°C and 25°C (36°F and 77°F) for up to 12 hours. Discard vial and any unused vaccine after 12 hours.					
nours						

*Vaccine expires 18 months after the manufacture date on the vial. Use Pfizer-BioNTech expiration date tool at <u>lotexpiry.cvd/vaccine.cc</u>

05/16/2023 CS321570-AN

Pfizer-BioNTech COVID-19 Vaccine At A Glance (cdc.gov) Administration of Moderna COVID-19 Vaccines | CDC



Other Vaccine Updates

New Vaccine Products/Indications

Vaccine	Product	What's New
MMR	Priorix (GSK)	Can be used for any dose in MMR series; mixed product series is acceptable
MenACWY	Menveo (GSK)	Single-dose vial Does NOT require reconstitution NOTE : Ages 10 through 55 years differ
PCV20	Prevnar20 (Pfizer)	FDA approved for children 6 weeks of age and older
RV5	Rotarix (GSK)	Does NOT require reconstitution

New Route: MMR II, Varivax, and ProQuad





Subcutaneous injection

OR

Intramuscular injection

Vaccination Resources for Healthcare Providers

Knowledgeable Staff Is Critical

CDC Vaccination Resources for Healthcare Providers

- Schedules App
- Pneumococcal Vaccination App
- Pneumococcal Vaccine Timing for Adults
- Vaccine Catch-up guidance



Immunization Education & Training





Vaccine Storage and Handling Toolkit

Updated with COVID-19 Vaccine Storage and Handling Information Addendum added September 29, 2021



An interactive, web-based immunization training course that include practice.

	YOU CAL SHC Vaccines reduce the • Rout	THE Source the definition of injection	Vacci Need esired tissue tion-site read • Age	to provide an optimal im tions. Needle selection s Gender and weigh for adults (19 years and weigh (19 years and weigh	stration: and Length hould be based on the: t • Injection site	
	when selecting ne	edles to administer inje	ctable vaccines.	- and lengths. In addition, clinic	ai judgment snould be used	
	Route Subcutaneous injection	Age All ages		Needle gauge and length 23–25-gauge 5/8 inch (16 mm)	Injection site Thigh for infants younger than 12 months of age ¹ ; upper outer triceps area for persons 12 months of age and older	
		Neonate, 28 days ar	nd younger	22–25-gauge 5/8 inch (16 mm ²)	Vastus lateralis muscle of anterolateral thigh	
		Infants, 1–12 mon Toddlers, 1–2 year	YOU C		Vaccine Administ Intramuscular (IM) In	ration:
acive training	Intramuscular injection	Children, 3–10 ye: Children, 11–18 yı	Administer t Diphtheria, tr biphtheria, tr hepatitis B (D Diphtheria, tr and Haemop	these vaccines by IM injection etanus, and pertussis (DTaP) - Di etanus, pertussis, polio, and TaP-IPV-Hegs, polio, - Hr hilluis infilierace type b	Infants 11 months of phtheria, tetanus, pertussis, polio, pe b (DTaP-IPV-HepB-Hib) permophilus influenzae type b pentitis 8 (HepB)	age and younger
Videos		Adults, 19 years an = 130 lbs (60 kg) o = 130–152 lbs (60- = Men, 152–260 lb	(DTaP-IPV/Hi Note: Age, recomm schedule for childre "May also be administ	ib) nendations for use, and other indications va en (<u>www.cdc.gov/vaccines/schedules/hcp/i</u> tered by subcutaneous injection.	ry by product. Always review manufacturers' produc imz/child-adolescent.html) before administering vac	t information as well as the current immunization tine.
Title: Comfort and Restraint Techniques		 Women, 152–20 Men, 260 lbs (11 			a late increasion to a surgery and a	desinistes these service the
Short Description: This training demonstrates comfort and restraint techniques. Determine the best position for based on comfort, age, activity level, administration site, and safety. Instruct the parent on how to help the infa stay still so you can administer the vaccine(s) safely.	or the patient ant or child dight umend a nusce i	Women, 200 lbs where the second sec	 Follow asep Use a new, Perform ha 	ptic technique. separate needle and syringe for ea and hygiene before vaccine prepara	re, it's important to prepare and a ich injection. ation, between patients, when changing g	loves (if worn), and any time hands
Title: Assemble a Manufacturer-filled Syringe			[†] Gloves are not requir If worn, perform hand	illed. [†] red unless the person administering the vaccine is d hygiene and change gloves between patients.	likely to come in contact with potentially infectious body fl	uids or has open lesions on the hands.
Short Description: This training addresses how to assemble a manufacturer-filled syringe, available for a variety CDC recommends that providers only prepare vaccines just prior to administration. Always prepare vaccines in area that is not near any area where potentially contaminated items are placed. Title: Single-Dose Vial	y of vaccines. <u>vcdc.</u> a designatec	.ommittee on Immunizat gov/vaccines/hcp/acip-re	 Use the col Administer Use the col 22- to 25 1-inch (2 *** A5/8-inch (16 mm) in 	rrect syringe and needle. r the vaccine using either a 1-mL or rrect gauge and needle length. [‡] - gauge needle 15 mm) needle 15 mm) needle	3-mL syringe.	Vastus Lateralis Muscle 1 in (25 mm)
Short Description: This training addresses how to prepare vaccine from a single-dose vial. A single-dose vial cor dose and should be administered one time to one patient. CDC recommends that providers only prepare and c vaccine just prior to administration.	ntains one draw up any		 Identify th Recommer Use anator the anterio injections- 	ne injection site. nded site: the vastus lateralis muscl mical landmarks to determine the i or lateral aspect of the thigh. The m -above the lateral condule and hele	e in the anterolateral thigh njection site. The muscle is located on iddle third of the muscle is used for we the greater trachanter	Greater trochanter
Title: Expiration Date			3. Administe	er the vaccine correctly.	with greater docharter.	Injection
Short Description: This training addresses how to determine when a vaccine or diluent expires—a critical step i preparation. All vaccines and diluents have an expiration date that indicates the date by which the product must Vaccines and diluents may be used up to and including the expiration date unless the manufacturer indicates or the second secon	n vaccine st be used. otherwise.		 Inject the v a 90-degre Aspiration vaccine. No process that For more in 	vaccine into the middle and thickes te angle and inject all the vaccine in (i.e., pulling back on the plunger) is o large blood vessels are present at at includes aspiration might be mo nformation, see https://www.cdc.uu	It part of the muscle. Insert the needle at the muscle tissue. s not necessary before injecting the the recommended injection site, and a re painful. w/vaccines/hcp/acio-recs/neneral-rocs/	Vastus Lateralis Lateral Patella
Title: <u>Multidose Vial (MDV)</u>			administra If administra	tion.html ering more than one injection in the	same limb, separate the injection sites by 1	First B
Short Description: This training addresses how to prepare vaccine from a multidose vial (MDV), which contains one dose of vaccine. CDC recommends that providers only prepare and draw up any vaccine just prior to admin	more than nistration.		inch, if poss For additional Advisory Comn Vaccine Admi	sible. I information, go to CDC's clinical r nittee on Immunization Practices G inistration section at www.cdc.gov/w	resources on vaccine administration ieneral Best Practice Guidelines for Immun accines/hcp/acip-recs/general-recs/adminis	ization: tration.html

Vaccine administration resource library at www.cdc.gov/vaccines/hcp/admin/resource-library.html

01/19/2022 CS 322033-F

Vaccine Administration Resource Library | CDC

Email Services and Websites

- Questions? Email CDC
- Vaccines and Immunizations website
- HCP education
- COVID-19 vaccine clinical materials
- Vaccinate with Confidence
- Influenza
- Vaccine safety

nipinfo@cdc.gov or CDC INFO | CDC

Vaccines and Immunizations | CDC

Vaccines and Immunizations | CDC

U.S. COVID-19 Vaccine Product Information | CDC

COVID-19 Vaccine Confidence | CDC

Influenza (Flu) | CDC

Vaccine Information and Safety Studies | Vaccine Safety | CDC