

Immunization Updates from the CDC and DHA: Catching Kids Up and Keeping Them Healthy in the Post Covid Era

A. Patricia Wodi, M.D., F.A.A.P. Immunization Services Division National Center for Immunization and Respiratory Diseases Centers for Disease Control and Prevention Atlanta Elisha Hall, Ph.D., R.D. Health Education Specialist National Center for Immunization and Respiratory Diseases Centers for Disease Control and Prevention Atlanta

> 28 April 2022 1120 - 1220 (ET)

Navy CAPT Gregory H Gorman, M.D., M.H.S. Executive Director, Defense Health Board Chair, DHA Complex Pediatrics Clinical Community, Falls Church, Va.

1

UNCLASSIFIED

Presenter(s)

A. Patricia Wodi, M.D., F.A.A.P. Immunization Services Division National Center for Immunization and Respiratory Diseases Centers for Disease Control and Prevention Atlanta Elisha Hall, Ph.D., R.D. Health Education Specialist National Center for Immunization and Respiratory Diseases at the Centers for Disease Control and Prevention Atlanta

Navy CAPT Gregory H Gorman, M.D., MHS Executive Director, Defense Health Board Chair, DHA Complex Pediatrics Clinical Community, Falls Church, VA



UNCLASSIFIED Medically Ready Force... Ready Medical Force



A. Patricia Wodi, M.D., F.A.A.P.



- A. Patricia Wodi, M.D., is a public health physician with the Centers for Disease Control and Prevention's Immunization Services Division, located in the National Center for Immunization and Respiratory Diseases. Prior to joining the CDC, Dr. Wodi worked in clinical practice and clinical drug development for over 15 years.
- At the CDC, she is the co-lead for the Advisory Committee on Immunization Practice Combined Immunization Schedule Work Group, editor for the Epidemiology and Prevention of Vaccine-Preventable Disease textbook (Pink Book), provides immunization educational resources and training to healthcare providers, and has worked on several vaccine safety research studies.
- Dr. Wodi holds a Doctor of Medicine degree from the College of Medicine, University of Port-Harcourt in Nigeria, and is board certified in general pediatrics and pediatric infectious diseases.





Elisha Hall, Ph.D., R.D.



- Elisha Hall, Ph.D., R.D., is a Health Education Specialist with the National Center for Immunization and Respiratory Diseases at the Centers for Disease Control and Prevention. As an educator with 11 years of experience, she creates, delivers, and evaluates immunizationrelated educational material and training.
- Dr. Hall first joined the CDC in 2017, transitioning to immunizations in 2020. In addition to being the lead editor on CDC's Epidemiology and Prevention of Vaccine-Preventable Diseases, otherwise known as the "Pink Book", she has spent the majority of the past two years in CDC's COVID-19 emergency response, holding various education and leadership roles within its Vaccine Task Force, most recently as the Clinical Guidelines Lead.
- She holds a BS and an MS in Nutrition and Health Sciences and a PhD in Human Sciences from the University of Nebraska-Lincoln.





Navy CAPT Gregory H. Gorman, M.D., M.H.S.



- CAPT Gregory Gorman is the Executive Director of the Defense Health Board and Chair of the Defense Health Agency Complex Pediatrics Clinical Community. He is a Professor of Pediatrics at the F. Edward Hébert School of Medicine at the Uniformed Services University and a practicing pediatric nephrologist for the Military Health System and the National Institutes of Health.
- A graduate of Georgetown University and the Washington University School of Medicine in St. Louis, he is board certified in Pediatrics, Pediatric Nephrology, and Clinical Informatics. He earned a Masters in Epidemiology at the Johns Hopkins Bloomberg School of Public Health.
- CAPT Gorman has authored over 30 peer-reviewed journal articles focusing on the health of military-connected children, childhood disease epidemiology, and military medicine. He is a member of the American Academy of Pediatrics, the American Society of Pediatric Nephrology, the American Medical Informatics Association, and AMSUS, the Society of Federal Health Professionals







Disclosures

- Dr. A. Patricia Wodi, Dr. Elisha Hall and CAPT Gregory Gorman have no relevant financial or non-financial relationships to disclose relating to the content of this activity; or presenter(s) must disclose the type of affiliation/financial interest (e.g., employee, speaker, consultant, principal investigator, grant recipient) with company name(s) included.
- The views expressed in this presentation are those of the authors and do not necessarily reflect the official policy or position of the Department of Defense, Centers for Disease Control and Prevention, nor the U.S. Government.
- Use of trade names of vaccine products is for identification purposes and does not imply endorsement by the Centers for Disease Control and Prevention (CDC).





Disclosures

- Use of vaccines in a manner not approved by the U.S. Food and Drug Administration will be discussed.
 - But in accordance with recommendations by the Advisory Committee on Immunization Practices (ACIP) This continuing education activity is managed and accredited by the Defense Health Agency, J-7, Continuing Education Program Office (DHA, J-7, CEPO). DHA, J-7, CEPO and all accrediting organizations do not support or endorse any product or service mentioned in this activity.
- DHA, J-7, CEPO staff, as well as activity planners and reviewers have no relevant financial or nonfinancial interest to disclose.
- Commercial support was not received for this activity.





Centers for Disease Control and Prevention National Center for Immunization and Respiratory Diseases



Child and Adolescent Immunization Schedule Catch-up Vaccination

A. Patricia Wodi, MD, FAAP Public Health Physician Communication and Education Branch Immunization Services Division National Center for Immunization and Respiratory Diseases

Learning Objectives

At the conclusion of this session participants will be able to:

- 1. Identify the various sections of CDC's childhood and adolescent immunization schedule.
- 2. Describe how to use the immunization schedule for children and adolescents who are behind on the recommended vaccination schedule.
- 3. Learn which vaccines can be co-administered.
- 4. Locate current CDC vaccination resources.



Overview of child and adolescent immunization schedule

Clinical scenario on catch-up vaccination

Co-administration of vaccines

Vaccination resources for healthcare providers

Child and Adolescent Immunization Schedule

Child and Adolescent Immunization Schedule

- Covers birth through 18 years
- Updated each year
 - Represents current, approved Advisory **Committee on Immunization Practices** (ACIP) policy
 - Designed for implementation of ACIP policy

Published in February

 MMWR Notice to Readers – announcement of availability of schedules on CDC website

Vaccines in the Child and Adolescent Immuniz	ation Schedule	•	Houston	an the shild	and adala	continue.	uninatio		
Vaccine	Abbreviation(s)	Trade name(s)	How to u	se the child	and adole	scent mini	unizatio		
Dengue vaccine	DEN4CYD	Dengvaxia*	schedule		1				
Diphtheria, tetanus, and acellular pertussis vaccine	DTaP	Daptacel* Infanrix*	1	2	3	4	5		
Diphtheria, tetanus vaccine	DT	No trade name	Determine recommended	Determine	for additional	types, frequencies,	contraindicat		
Haemophilus influenzae type b vaccine	Hib (PRP-T) Hib (PRP-OMP)	ActHIB* Hiberix* PedvaxHIB*	vaccine by age (Table 1)	interval for catch- up vaccination (Table 2)	recommended vaccines by medical condition or other indication (Table 3)	intervals, and considerations for special situations	and precauti for vaccine ty (Appendix)		
Hepatitis A vaccine	НерА	Havrix* Vaqta*				(Notes)			
Hepatitis B vaccine	НерВ	Engerix-B* Recombivax HB*	Recommended b	withe Achikopy Comm	ttee on Immunization	Diractices funnue of	r nau kranninas la		
Human papillomavirus vaccine	HPV	Gardasil 9*	and approved by	the Centers for Diseas	e Control and Prevent	tion (www.cdc.gov),	American Acade		
Influenza vaccine (inactivated)	IIV4	Multiple	of Pediatrics (www	w.aap.org), American /	cademy of Family Ph	ysicians (www.aafp.o	arg), American		
Influenza vaccine (live, attenuated)	LAIV4	FluMist* Quadrivalent	(www.midwife.or	 and Gynecolog and Gynecolog and Gynecolog 	of Physician Associat	American College of tes (www.aapa.org), i	and National		
Measles, mumps, and rubella vaccine	MMR	M-M-BII*	Association of Pee	diatric Nurse Practition	ers (www.napnap.org	g).			
Meningococcal serogroups A, C, W, Y vaccine	MenACWY-D	Menactra*	Dement						
	MenACWY-CRM	Menveo*	 Suspected cases of reportable vaccine-preventable diseases or outbreaks to your state or local heal 						
	MenACWY-TT	MenQuadfi*	department			,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,			
Meningococcal serogroup B vaccine	MenB-4C	Bexsero*	 Clinically significant adverse events to the Vaccine Adverse Event Reporting System (VAERS) 						
	MenB-FHbp	Trumenba*	www.vacis.mis.go	SV 01 800-822-7907					
Pneumococcal 13-valent conjugate vaccine	PCV13	Prevnar 13*	Questions or	comments					
Pneumococcal 23-valent polysaccharide vaccine	PPSV23	Pneumovax 23*	Contact www.cdc.g	ov/cdc-info or 800-CD	C-INFO (800-232-4636)), in English or Spanis	h, 8 a.m8 p.m. l		
Poliovirus vaccine (inactivated)	IPV	IPOL*	Monday anough H	iday, excloding noiday	9				
Rotavirus vaccine	RV1 RV5	Rotarix* RotaTeq*	Download www.cdc/	I the CDC Vaccine Sche gow/vaccines/schedule	dules app for provider s/hcp/schedule-app.h	s at tml			
Tetanus, diphtheria, and acellular pertussis vaccine	Tdap	Adacel* Boostrix*	Helpful infor	mation					
Tetanus and diphtheria vaccine	Td	Tenivac" Tdvax"	Complete Advisor www.cdc.gov/vac General Rest Practic	ry Committee on Immu cines/hcp/acip-recs/in ice Guidelines for Immu	dex.html vization (including con	(P) recommendations	2 weautions):		
Varicella vaccine	VAR	Varivax*	www.cdc.gov/vac	cines/hcp/acip-recs/ge	neral-recs/index.html		condensition.		
Combination vaccines (use combination vaccines instead of sepa	rate injections when a	ppropriate)	 Vaccine information 	on statements:	International				
DTaP, hepatitis B, and inactivated poliovirus vaccine	DTaP-HepB-IPV	Pediarix*	Manual for the Su	cones/ncp/vis/index.nc	mi reventable Diseases				
DTaP, inactivated poliovirus, and Haemophilus influenzae type b vaccine	DTaP-IPV/Hib	Pentacel*	(including case id	entification and outbre	ak response):				
DTaP and inactivated poliovirus vaccine	DTaP-IPV	Kinrix* Quadracel*	ACIP Shared Clinic	cines/pubs/surv-manu cal Decision-Making Re cines/acin/acin-scrim-	commendations		Scan QR for acce online sci		
DTaP, inactivated poliovirus, Haemophilus influenzae type b, and hepatitis B vaccine	DTaP-IPV-Hib- HepB	Vaxelis*	and a sumary	conceptacip score	and a set of				
Measles mumos nihella and varicella vaccine	MMRV	ProOuad*	1 II m	U.S. De	epartment of		1000		

Recommended Child and Adolescent Immunization Schedule



UNITED STATES

an OR co

Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger

Vaccines in the Child and Adolescent Immunizat	ion Schedule*	
Vaccine	Abbreviation(s)	Trade name(s)
Dengue vaccine	DEN4CYD	Dengvaxia*
Diphtheria, tetanus, and acellular pertussis vaccine	DTaP	Daptacel* Infanrix*
Diphtheria, tetanus vaccine	DT	No trade name
Haemophilus influenzae type b vaccine	Hib (PRP-T) Hib (PRP-OMP)	ActHIB* Hiberix* PedvaxHIB*
Hepatitis A vaccine	НерА	Havrix® Vaqta®
Hepatitis B vaccine	НерВ	Engerix-B* Recombivax HB*
Human papillomavirus vaccine	HPV	Gardasil 9*
Influenza vaccine (inactivated)	IIV4	Multiple
Influenza vaccine (live, attenuated)	LAIV4	FluMist [®] Quadrivalent
Measles, mumps, and rubella vaccine	MMR	M-M-R II*
Meningococcal serogroups A, C, W, Y vaccine	MenACWY-D	Menactra*
	MenACWY-CRM	Menveo*
	MenACWY-TT	MenQuadfi*
Meningococcal serogroup B vaccine	MenB-4C	Bexsero*
	MenB-FHbp	Trumenba*
Pneumococcal 13-valent conjugate vaccine	PCV13	Prevnar 13*
Pneumococcal 23-valent polysaccharide vaccine	PPSV23	Pneumovax 23*
Poliovirus vaccine (inactivated)	IPV	IPOL*
Rotavirus vaccine	RV1 RV5	Rotarix* RotaTeq*
Tetanus, diphtheria, and acellular pertussis vaccine	Tdap	Adacel* Boostrix*
Tetanus and diphtheria vaccine	Td	Tenivac® Tdvax™
Varicella vaccine	VAR	Varivax*
Combination vaccines (use combination vaccines instead of separate	e injections when app	propriate)

	,,	
DTaP, hepatitis B, and inactivated poliovirus vaccine	DTaP-HepB-IPV	Pediarix*
DTaP, inactivated poliovirus, and Haemophilus influenzae type b vaccine	DTaP-IPV/Hib	Pentacel®
DTaP and inactivated poliovirus vaccine	DTaP-IPV	Kinrix* Quadracel*
DTaP, inactivated poliovirus, Haemophilus influenzae type b, and hepatitis B vaccine	DTaP-IPV-Hib- HepB	Vaxelis*
Measles, mumps, rubella, and varicella vaccine	MMRV	ProQuad*

*Administer recommended vaccines if immunization history is incomplete or unknown. Do not restart or add doses to vaccine series for extended intervals between doses. When a vaccine is not administered at the recommended age, administer at a subsequent visit. The use of trade names is for identification purposes only and does not imply endorsement by the ACIP or CDC.



Recommended by the Advisory Committee on Immunization Practices (www.cdc.gov/vaccines/acip) and approved by the Centers for Disease Control and Prevention (www.cdc.gov), American Academy of Pediatrics (www.aap.org), American Academy of Family Physicians (www.aafp.org), American College of Obstetricians and Gynecologists (www.acog.org), American College of Nurse-Midwives (www.midwife.org), American Academy of Physician Associates (www.aap.org), and National Association of Pediatric Nurse Practitioners (www.napnap.org).

Report

 Suspected cases of reportable vaccine-preventable diseases or outbreaks to your state or local health department

 Clinically significant adverse events to the Vaccine Adverse Event Reporting System (VAERS) at www.vaers.hhs.gov or 800-822-7967

Questions or comments

Contact www.cdc.gov/cdc-info or 800-CDC-INFO (800-232-4636), in English or Spanish, 8 a.m.–8 p.m. ET, Monday through Friday, excluding holidays

Download the CDC Vaccine Schedules app for providers at www.cdc.gov/vaccines/schedules/hcp/schedule-app.html

Helpful information

 Complete Advisory Committee on Immunization Practices (ACIP) recommendations: www.cdc.gov/vaccines/hcp/acip-recs/index.html

 General Best Practice Guidelines for Immunization (including contraindications and precautions): www.cdc.gov/vaccines/hcp/acip-recs/general-recs/index.html

- Vaccine information statements:
- www.cdc.gov/vaccines/hcp/vis/index.html
- Manual for the Surveillance of Vaccine-Preventable Diseases (including case identification and outbreak response): www.cdc.gov/vaccines/pubs/surv-manual
- ACIP Shared Clinical Decision-Making Recommendations www.cdc.gov/vaccines/acip/acip-scdm-faqs.html



U.S. Department of Health and Human Services Centers for Disease Control and Prevention



UNITED STATES



Table 1 Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2022

These recommendations must be read with the notes that follow. For those who fall behind or start late, provide catch-up vaccination at the earliest opportunity as indicated by the green bars. To determine minimum intervals between doses, see the catch-up schedule (Table 2).

Vaccine	Birth	1 mo	2 mos	4 mos	6 mos	9 mos	12 mos	15 mos	18 mos	19–23 mos	2–3 yrs	4–6 yrs	7–10 yrs	11–12 yrs	13–15 yrs	16 yrs	17–18 yrs
Hepatitis B (HepB)	1ª dose	< 2 nd	dose>		4		3 rd dose		>								
Rotavirus (RV): RV1 (2-dose series), RV5 (3-dose series)			1 st dose	2 nd dose	See Notes												
Diphtheria, tetanus, acellular pertussis (DTaP <7 yrs)			1ª dose	2 nd dose	3 rd dose			∢ 4 th d	oseÞ			5 th dose					
Haemophilus influenzae type b (Hib)			1 st dose	2 nd dose	See Notes		4 ^{3rd or 4 See 1}	th dose, Notes									
Pneumococcal conjugate (PCV13)			1 st dose	2 nd dose	3 rd dose		∢ 4 th c	lose —>									
Inactivated poliovirus (IPV <18 yrs)			1ª dose	2 nd dose	•		3 rd dose					4 th dose					
Influenza (IIV4)							A	nnual vacci	nation 1 or	2 doses				Annua	lvaccinatior	1 dose on	у
Influenza (LAIV4)											Annua 1 o	l vaccinatio r 2 doses	n	Annua	lvaccinatior	1 dose on	у
Measles, mumps, rubella (MMR)					See N	Notes	4 1** c	lose>				2 nd dose					
Varicella (VAR)							∢ 1* c	lose>				2 nd dose					
Hepatitis A (HepA)					See N	Notes		2-dose serie	s, See Note	s							
Tetanus, diphtheria, acellular pertussis (Tdap ≥7 yrs)														1 dose			
Human papillomavirus (HPV)														See Notes			
Meningococcal (MenACWY-D ≥9 mos, MenACWY-CRM ≥2 mos, MenACWY-TT ≥2years)								See Notes						1ª dose		2 nd dose	
Meningococcal B (MenB-4C, MenB- FHbp)															See No	tes	
Pneumococcal polysaccharide (PPSV23)														See Notes			
Dengue (DEN4CYD; 9-16 yrs)													Se	eropositive i (S	n endemic a ee Notes)	reas only	
Range of recommended ages for all children	Range of r for catch-u	ecommend up vaccinati	led ages ion	Rar for	nge of recon certain high	nmended a n-risk group	ges s	Recomr can beg	mended vac jin in this ag	cination le group	Re	ecommende n shared clin	ed vaccination	on based n-making	No	recommer t applicabl	ndation/ e

Table 2 Recommended Catch-up Immunization Schedule for Children and Adolescents Who Start Late or Who Are More than 1 Month Behind, United States, 2022

The table below provides catch-up schedules and minimum intervals between doses for children whose vaccinations have been delayed. A vaccine series does not need to be restarted, regardless of the time that has elapsed between doses. Use the section appropriate for the child's age. Always use this table in conjunction with Table 1 and the Notes that follow.

			Children age 4 months through 6 years		
Vaccine	Minimum Age for		Minimum Interval Between Doses		
	Dose 1	Dose 1 to Dose 2	Dose 2 to Dose 3	Dose 3 to Dose 4	Dose 4 to Dose 5
Hepatitis B	Birth	4 weeks	8 weeks and at least 16 weeks after first dose minimum age for the final dose is 24 weeks		
Rotavirus	6 weeks Maximum age for first dose is 14 weeks, 6 days.	4 weeks	4 weeks maximum age for final dose is 8 months, 0 days		
Diphtheria, tetanus, and acellular pertussis	6 weeks	4 weeks	4 weeks	6 months	6 months
Haemophilus influenzae type b	6 weeks	No further doses needed if first dose was administered at age 15 months or older. 4 weeks if first dose was administered before the 1 ^s birthday. 8 weeks (as final dose) if first dose was administered at age 12 through 14 months.	No further doses needed 4 weeks if current age is younger than 12 months and first dose was administered at younger than age 7 months and at least 1 previous dose was PRP-T (ActHib [*] , Pentacel [*] , Hiberk [*]), Vaxelis [*] or unknown 8 weeks and age 12 through 39 months and first dose was administered at age 7 through 11 months; OR If current age is 12 through 59 months and first dose was administered before the 1 st birthday and second dose was administered at younger than 15 months; OR If both doses were PedvaxHiB [*] and were administered before the 1st birthday	8 weeks (as final dose) This dose only necessary for children age 12 through 59 months who received 3 doses before the 1 st birthday.	
Pneumococcal conjugate	6 weeks	No further doses needed for healthy children if first dose was administered at age 24 months or older 4 weeks if first dose was administered before the 1 st birthday 8 weeks (as final dose for healthy children) If first dose was administered at the 1 st birthday or after	No further doses needed for healthy children if previous dose was administered at age 24 months or older 4 weeks if current age is younger than 12 months and previous dose was administered at <7 months old 8 weeks (as final dose for healthy children) if previous dose was administered between 7–11 months (wait until at least 12 months old); OR if current age is 12 months or older and at least 1 dose was administered before age 12 months	8 weeks (as final dose) This dose only necessary for children age 12 through 59 months who received 3 doses before age 12 months or for children at high risk who received 3 doses at any age.	
Inactivated poliovirus	6 weeks	4 weeks	4 weeks if current age is <4 years 6 months (as final dose) if current age is 4 years or older	6 months (minimum age 4 years for final dose)	
Measles, mumps, rubella	12 months	4 weeks			
Varicella	12 months	3 months			
Hepatitis A	12 months	6 months			
Meningococcal ACWY	2 months MenACWY-CRM 9 months MenACWY-D 2 years MenACWY-TT	8 weeks	See Notes	See Notes	
			Children and adolescents age 7 through 18 years		
Meningococcal ACWY	Not applicable (N/A)	8 weeks			
Tetanus, diphtheria; tetanus, diphtheria, and acellular pertussis	7 years	4 weeks	4 weeks if first dose of DTaP/DT was administered before the 1 st birthday 6 months (as final dose) if first dose of DTaP/DT or Tdap/Td was administered at or after the 1 st birthday	6 months if first dose of DTaP/DT was administered before the 1 st birthday	
Human papillomavirus	9 years	Routine dosing intervals are recommended.			
Hepatitis A	N/A	6 months			
Hepatitis B	N/A	4 weeks	8 weeks and at least 16 weeks after first dose		
Inactivated poliovirus	N/A	4 weeks	6 months A fourth dose is not necessary if the third dose was administered at age 4 years or older and at least 6 months after the previous dose.	A fourth dose of IPV is indicated if all previous doses were administered at <4 years or if the third dose was administered <6 months after the second dose.	
Measles, mumps, rubella	N/A	4 weeks			
Varicella	N/A	3 months if younger than age 13 years.			
Dengue	9 years	6 months	6 months		



Recommended Child and Adolescent Immunization Schedule by Medical Indication, United States, 2022

Always use this table in conjunction with Table 1 and the Notes that follow.

					11	DICATION				
			HIV infectio	n CD4+ count ¹						
VACCINE	Pregnancy	Immunocom- promised status (excluding HIV infection)	<15% or total CD4 cell count of <200/mm ³	≥15% and total CD4 cell count of ≥200/mm ³	Kidney failure, end-stage renal disease, or on hemodialysis	Heart disease or chronic lung disease	CSF leak or cochlear implant	Asplenia or persistent complement component deficiencies	Chronic liver disease	Diabetes
Hepatitis B										
Rotavirus		SCID ²								
Diphtheria, tetanus, and acellular pertussis (DTaP)										
Haemophilus influenzae type b										
Pneumococcal conjugate										
Inactivated poliovirus										
Influenza (IIV4)										
Influenza (LAIV4)						Asthma, wheezing: 2–4yrs ³				
Measles, mumps, rubella	*									
Varicella	*									
Hepatitis A										
Tetanus, diphtheria, and acellular pertussis (Tdap)										
Human papillomavirus	*									
Meningococcal ACWY										
Meningococcal B										
Pneumococcal polysaccharide										
Dengue										
Vaccination according to routine schedule recommended	o the	Recommended for persons with an additic factor for which the vac would be indicated	onal risk ccine	Vaccination is recom and additional doses necessary based on r condition or vaccine.	mended, may be medical See Notes.	Precaution—vaccine might be indicated if benefit of protection outweighs risk of adverse reaction	Contraindio recommen not be adm *Vaccinate	ated or not ded—vaccine should inistered after pregnancy	No recomme applicable	endation/not

1 For additional information regarding HIV laboratory parameters and use of live vaccines, see the General Best Practice Guidelines for Immunization, "Altered Immunocompetence," at

www.cdc.gov/vaccines/hcp/acip-recs/general-recs/immunocompetence.html and Table 4-1 (footnote J) at www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html. 2 Severe Combined Immunodeficiency 3 LAIV4 contraindicated for children 2–4 years of age with asthma or wheezing during the preceding 12 months 16

Notes Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2022

Adolescent vaccination of children who received MenACWY prior to

ongoing increased risk of meningococcal disease (e.g., those with complement deficiency, HIV, or asplenia): Follow the booster schedule

Children for whom boosters are not recommended (e.g., a healthy

meningococcal disease is endemic): Administer MenACWY according

Children for whom boosters are recommended because of an

child who received a single dose for travel to a country where

Meningococcal serogroup A,C,W,Y vaccination (minimum age: 2 months [MenACWY-CRM, Menveo], 9 months [MenACWY-D, Menactra], 2 years [MenACWY-TT, MenQuadfi])

Routine vaccination

2-dose series at age 11–12 years; 16 years

Catch-up vaccination Age 13–15 years: 1 dose now and I

(minimum interval: 8 weeks) Age 16–18 years: 1 dose

Special situations Anatomic or functional asplenia **HIV infection, persistent complet** complement inhibitor (e.g., eculi:

 Menveo - Dose 1 at age 2 months: 4-dose

and 12 months) Dose 1 at age 3-6 months: 3- or 4

after dose 1 and after age 12 mor

- Dose 1 at age 24 months or older

Persistent complement compo

• Menactra

inhibitor use:

2-dose series (minimum interval: 6 months) at age 12–23 months 3 if applicable] at least 8 weeks af

Catch-up vaccination received at age 7 months or olde Unvaccinated persons through age 18 years should complete a least 12 weeks later and after age Dose 1 at age 7–23 months: 2-do 2-dose series (minimum interval: 6 months)

Notes

Hepatitis A vaccination

Routine vaccination

months.

Persons who previously received 1 dose at age 12 months or older

(minimum age: 12 months for routine vaccination)

age 10 years:

for persons at increased risk.

should receive dose 2 at least 6 months after dose 1. Adolescents age 18 years or older may receive the combined HepA and HepB vaccine, Twinrix®, as a 3-dose series (0, 1, and 6 months) or

4-dose series (3 doses at 0, 7, and 21-30 days, followed by a booster Age 9-23 months: 2-dose series Age 24 months or older: 2-dose dose at 12 months).

Anatomic or functional aspleni International travel infection:

 Persons traveling to or working in countries with high or intermediate Age 9-23 months: Not recomm endemic hepatitis A (www.cdc.gov/travel/): Age 24 months or older: 2-dose Infants age 6-11 months: 1 dose before departure; revaccinate Menactra® must be administer with 2 doses, separated by at least 6 months, between age 12-23

of PCV13 series. MenQuadfi[®] Dose 1 at age 24 months or older

Unvaccinated age 12 months or older: Administer dose 1 as soon Travel in countries with hyperend as travel is considered disease, including countries in the the Hajj (www.cdc.gov/travel/): Hepatitis B vaccination Children less than age 24 months: Menveo[®] (age 2-23 months) (minimum age: birth)

· Dose 1 at age 2 months: 4-dose and 12 months) Dose 1 at age 3-6 months: 3- or 3 if applicable] at least 8 weeks a received at age 7 months or old least 12 weeks later and after ad Dose 1 at age 7–23 months: 2-d after dose 1 and after age 12 mc Menactra® (age 9-23 months) · 2-dose series (dose 2 at least 12

administered as early as 8 week Mother is HBsAg-positive: Children age 2 years or older: 1 do: MenQuadfi

First-year college students who liv previously vaccinated at age 16 ye * 1 dose Menveo[®], Menactra[®], or N

vaccine (total of 4 doses) beginning at age 1 month. - Test for HBsAg and anti-HBs at age 9–12 months. If HepB series is delayed, test 1-2 months after final dose.

Birth dose (monovalent HepB vaccine only)

- All medically stable infants ≥2.000 grams: 1 dose within 24 hours of

- Infants <2,000 grams: Administer 1 dose at chronological age

1 month or hospital discharge (whichever is earlier and even if

(in separate limbs) within 12 hours of birth, regardless of birth

weight. For infants <2,000 grams, administer 3 additional doses of

• Mother's HBsAg status is unknown:

Mother is HBsAg-negative:

weight is still <2,000 grams).

- Administer HepB vaccine within 12 hours of birth, regardless of birth weight.
- For infants <2,000 grams, administer HBIG in addition to HepB vaccine (in separate limbs) within 12 hours of birth. Administer 3 additional doses of vaccine (total of 4 doses) beginning at age 1 month.
- Determine mother's HBsAg status as soon as possible. If mother is HBsAg-positive, administer HBIG to infants ≥2,000 grams as soon as possible, but no later than 7 days of age.

Routine series

- 3-dose series at age 0, 1–2, 6–18 months (use monovalent HepB) vaccine for doses administered before age 6 weeks)
- Infants who did not receive a birth dose should begin the series as soon as feasible (see Table 2).

containing HepB is used after the birth d Minimum age for the final (3rd or 4th) dc Notes Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2022 Minimum intervals: dose 1 to dose 2:4

Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2022

mellitus:

Special situations

be administered during same visit.

Underlying conditions below: When both PCV13 and PPSV23 are

indicated, administer PCV13 first. PCV13 and PPSV23 should not

Chronic heart disease (particularly cyanotic congenital heart disease and cardiac failure); chronic lung disease (including

asthma treated with high-dose, oral corticosteroids); diabetes

8 weeks / dose 1 to dose 3: 16 weeks (wh substitute "dose 4" for "dose 3" in these c Catch-up vaccination

For vaccination recommendations for persons ages 19 years or older, see the Recommended Adult Immunization Schedule, 2022.

Administration of 4 doses is permitted when a combination vaccine
 Pregnancy: Pregnancy: testing not needed before vaccination: HPV

Additional information

COVID-19 Vaccination COVID-19 vaccines are recommended for use within the scope of the Emergency Use Authorization or Biologics License Application for the particular vaccine. ACIP recommendations for the use of COVID-19 vaccines can be found at www.cdc.gov

vaccines/hcp/acip-recs/vacc-specific/covid-19.html CDC's interim clinical considerations for use of COVID-19 vaccines can be found at www.cdc.gov/vaccines/covid-19/clinica considerations/covid-19-vaccines-us.html.

Consult relevant ACIP statements for detailed recommendations at

Within a number range (e.g., 12–18), a dash (–) should be read as

Vaccine doses administered ≤4 days before the minimum age or

interval are considered valid. Doses of any vaccine administered

≥5 days earlier than the minimum age or minimum interval should

not be counted as valid and should be repeated as age appropriate.

The repeat dose should be spaced after the invalid dose by the

Recommended and minimum ages and intervals between vaccine

· Information on travel vaccination requirements and recommendations

Immunization at www.cdc.gov/vaccines/hcp/acip-recs/general-recs/

Circumstances (In: Kimberlin DW, Brady MT, Jackson MA, Long SS, eds.

The National Vaccine Injury Compensation Program (VICP) is a no-fault

alternative to the traditional legal system for resolving vaccine injury

claims. All routine child and adolescent vaccines are covered by VICP

except for pneumococcal polysaccharide vaccine (PPSV23). For more

information, see www.hrsa.gov/vaccinecompensation/index.html.

Red Book: 2018 Report of the Committee on Infectious Diseases. 31st ed.

mmunocompetence.html, and Immunization in Special Clinical

www.cdc.gov/vaccines/hcp/acip-recs/general-recs/timing.html.

doses, in General Best Practice Guidelines for Immunization at

For vaccination of persons with immunodeficiencies, see

Table 8-1, Vaccination of persons with primary and secondary

immunodeficiencies, in General Best Practice Guidelines for

Itasca, IL: American Academy of Pediatrics; 2018:67-111).

· For information about vaccination in the setting of a vaccine-

preventable disease outbreak, contact your state or local health

recommended minimum interval. For further details, see Table 3-1.

For calculating intervals between doses, 4 weeks = 28 days. Intervals of

www.cdc.gov/vaccines/hcp/acip-recs/index.html.

≥4 months are determined by calendar months.

is available at www.cdc.gov/travel/.

"through"

department.

For other catch-up guidance, see Table 2 **Special situations**

Unvaccinated persons should complete

Adolescents age 11–15 years may use ar

Adolescents age 18 years or older may r

(Heplisav-B*) at least 4 weeks apart.

Adolescents age 18 years or older may re

and HepB vaccine, Twinrix®, as a 3-dose

4-dose series (3 doses at 0, 7, and 21-30

schedule with at least 4 months betwee

months.

 Revaccination is not generally recommendation normal immune status who were vaccin adolescents, or adults.

Recombivax HB^e only)

dose at 12 months)

 Post-vaccination serology testing and 10mlU/mL) is recommended for certain Infants born to HBsAg-positive motified

Hemodialysis patients Other immunocompromised person For detailed revaccination recommendation

vaccines/hcn/acin-recs/vacc-specific/hepl Human papillomavirus vacci (minimum age: 9 years)

Routine and catch-up vaccinat Administer HepB vaccine and hepatitis B immune globulin (HBIG) HPV vaccination routinely recommende start at age 9 years) and catch-up HPV all persons through age 18 years if not a 2- or 3-dose series depending on age at

> Age 9–14 years at initial vaccination: months (minimum interval: 5 months; too soon) Age 15 years or older at initial vaccir

months, 6 months (minimum intervals dose 2 to dose 3: 12 weeks / dose 1 to if administered too soon

 Interrupted schedules: If vaccination set series does not need to be restarted. No additional dose recommended wher

been completed using the recommende Special situations Immunocompromising conditions, in

3-dose series, even for those who initiate 14 years

History of sexual abuse or assault: Sta

• Age 9-16 years living in dengue endemic areas AND have laboratory confirmation of previous dengue infection - 3-dose series administered at 0.6, and 12 months

Dengue vaccination

Routine vaccination

(minimum age: 9 years)

 Endemic areas include Puerto Rico, American Samoa, US Virgin Islands. Federated States of Micronesia, Republic of Marshall Islands, and the Republic of Palau. For updated guidance on dengue endemic areas and pre-vaccination laboratory testing see www.cdc.gov/mmwr/ volumes/70/rr/rr7006a1.htm?s_cid=rr7006a1_w and www.cdc.gov/ dengue/vaccine/hcp/index.html

Diphtheria, tetanus, and pertussis (DTaP) vaccination (minimum age: 6 weeks [4 years for Kinrix[®] or Ouadracel[®]])

Routine vaccination

• 5-dose series at age 2, 4, 6, 15-18 months, 4-6 years - Prospectively: Dose 4 may be administered as early as age 12 months if at least 6 months have elapsed since dose 3.

 Retrospectively: A 4th dose that was inadvertently administered as early as age 12 months may be counted if at least 4 months have

elapsed since dose 3. Catch-up vaccination

 Dose 5 is not necessary if dose 4 was administered at age 4 years or older and at least 6 months after dose 3. For other catch-up guidance, see Table 2.

Special situations

· Wound management in children less than age 7 years with history of 3 or more doses of tetanus-toxoid-containing vaccine: For all wounds except clean and minor wounds, administer DTaP if more than 5 years since last dose of tetanus-toxoid-containing vaccine. For detailed information, see www.cdc.gov/mmwr/volumes/67/rr/rr6702a1.htm.

Haemophilus influenzae type b vaccination (minimum age: 6 weeks)

Routine vaccination

 ActHIB[®], Hiberix[®], Pentacel[®], or Vaxelis[®]: 4-dose series (3 dose primary series at age 2, 4, and 6 months, followed by a booster dose* at age 12-15 months)

- *Vaxelis* is not recommended for use as a booster dose. A different Hib-containing vaccine should be used for the booster dose.
- PedvaxHIB[®]: 3-dose series (2-dose primary series at age 2 and 4 months, followed by a booster dose at age 12-15 months)

Catch-up vaccination

• Dose 1 at age 7-11 months: Administer dose 2 at least 4 weeks later and dose 3 (final dose) at age 12-15 months or 8 weeks after dose 2 (whichever is later)

• Dose 1 at age 12-14 months: Administer dose 2 (final dose) at least 8 weeks after dose 1

Dose 1 before age 12 months and dose 2 before age 15 months: Administer dose 3 (final dose) at least 8 weeks after dose 2. * 2 doses of PedvaxHIB* before age 12 months: Administer dose 3 (final dose) at 12-59 months and at least 8 weeks after dose 2.

17

needed

For other catch-up guidance, see Table 2. Vaxelise can be used for catchup vaccination in children less than age 5 years. Follow the catch-up schedule even if Vaxelis® is used for one or more doses. For detailed information on use of Vaxelis* see www.cdc.gov/mmwr/volumes/69/ wr/mm6905a5.htm

Special situations

Chemotherapy or radiation treatment: Age 12-59 months

- Unvaccinated or only 1 dose before age 12 months: 2 doses, 8 weeks apart

- 2 or more doses before age 12 months: 1 dose at least 8 weeks after previous dose

Doses administered within 14 days of starting therapy or during therapy should be repeated at least 3 months after therapy completion.

Hematopoietic stem cell transplant (HSCT);

- 3-dose series 4 weeks apart starting 6 to 12 months after successful transplant, regardless of Hib vaccination history

* Anatomic or functional asplenia (including sickle cell disease): Age 12-59 months

- Unvaccinated or only 1 dose before age 12 months: 2 doses, 8 weeks apart

- 2 or more doses before age 12 months: 1 dose at least 8 weeks after previous dose

Unvaccinated* persons age 5 years or older

Elective splenectomy:

- 1 dose

Unvaccinated* persons age 15 months or older - 1 dose (preferably at least 14 days before procedure)

HIV infection: Age 12–59 months

- Unvaccinated or only 1 dose before age 12 months: 2 doses,

8 weeks apart

 2 or more doses before age 12 months: 1 dose at least 8 weeks after previous dose Unvaccinated* persons age 5-18 years

- 1 dose

no doses (age 15 months or older)

 Immunoglobulin deficiency, early component complement deficiency:

Age 12–59 months

 Unvaccinated or only 1 dose before age 12 months: 2 doses, 8 weeks apart

- 2 or more doses before age 12 months: 1 dose at least 8 weeks after

previous dose *Unvaccinated = Less than routine series (through age 14 months) OR

* 1 dose administered at age 15 months or older: No further doses

• Unvaccinated at age 15-59 months: Administer 1 dose. Previously unvaccinated children age 60 months or older who are not considered high risk: Do not require catch-up vaccination

Appendix	Recommended Child an	nd Adolescent Immuniz	ation Sched	ule for ages 18 years or younger, United States, 2022			
Vaccine	Contraindications ¹			Precautions ²			
Dengue (DEN4CYD)	 Severe allergic reaction (e.g., anaphylaxis) af Severe immunodeficiency (e.g., hematologi immunodeficiency, long- term immunosup immunocompromised) 	ter a previous dose or to a vaccine compone c and solid tumors, receipt of chemotherapy pressive therapy or patients with HIV infection	ent ^a , congenital on who are severely	Pregnancy HIV infection without evidence of severe immunosuppression Moderate or severe acute illness with or without fever			
Diphtheria, tetanus, pertussis (DTaP) Tetanus, diphtheria (DT)	Severe allergic reaction (e.g., anaphylaxis) af For DTaP only: Encephalopathy (e.g., coma, attributable to another identifiable cause w	fter a previous dose or to a vaccine compone decreased level of consciousness, prolonged ithin 7 days of administration of previous do	ent ³ I seizures) not se of DTP or DTaP	Guillain-Barré syndrome (GBE) within 6 weeks after previous dose of tetanus-toxoidontaining vaccine History of Arthus-type hypersensitivity reactions after a previous dose of diphtheria-toxoid— containing or tetanus-trunid containing vaccine defer vaccination until at least 10 years have elansed since the la			
Haemophilus influenzae type b (Hib)	 Severe allergic reaction (e.g., anaphylaxis) i For Hiberix, ActHib, and PedvaxHIB only: H Less than ane 6 weeks 	Appendix Guide to Contraindicatio	ons and Preca	utions to Commonly Used Vaccines	ute for ages 18 years or younger, United States, 2022		
Hepatitis A (HepA)	Severe allergic reaction (e.g., anaphylaxis) a neomycin	recs/general-recs/contraindig	cations.html an	d ACIP's Recommendations for the Prevention and Control of 2021-22 seasonal i	influenza with Vaccines available at www.cdc.gov/mwr/volumes/70/rr/rr7005a1.htm.		
Hepatitis B (HepB)	Severe allergic reaction (e.g., anaphylaxis) = For Heplisav-B only: Pregnancy	Interim clinical consid	erations for u	se of COVID-19 vaccines including contraindications and precau	tions can be found at		
Hepatitis A- Hepatitis B vaccine [HepA-HepB, (Twinrix*)]	Severe allergic reaction (e.g., anaphylaxis) a neomycin and yeast	www.cdc.gov/vaccines/	covid-19/clini	cal-considerations/covid-19-vaccines-us.html			
Human papillomavirus (HPV)	Severe allergic reaction (e.g., anaphylaxis) a	Vaccino	Controlndicat	long1	Dragoutions?		
Measles, mumps, rubella (MMR)	Severe allergic reaction (e.g., anaphylaxis) : Severe immunodeficiency (e.g., hematolog immunodeficiency, long-term immunosus immunocompromised) Pregnancy Family history of altered immunocompete immunocompetent	Influenza, egg-based, inactivated injectable (IIV4)	 Severe allergi any egg-base Severe allergi 	ons ^s c reaction (e.g., anaphylaxis) after previous dose of any influenza vaccine (i.e., d IIV, ccIIV, RIV, or LAIV of any valency) c 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 Persons with egg allergy with symptoms other than hives (e.g., angioedema, respiratory distress) or required epinephrine or another emergency medical intervention: Any influenza vaccine appropriate for age and health status may be administered. If using egg-based IIV4, administer in medical setting under supervision of health care provider 		
Meningococcal ACWY (MenACWY) [MenACWY-CRM (Menveo"); MenACWY-D (Menactra");	 Severe allergic reaction (e.g., anaphylaxis) i For MenACWY-D and Men ACWY-CRM only containing vaccine 				who can recognize and manage severe allergic reactions. May consult an allergist. • Moderate or severe acute illness with or without fever		
MenACWY-TT (MenQuadfi*)] Meningococcal B (MenB) [MenB-4C (Bexsero*); MenB-FHbp (Trumenba*)]	For MenACWY-TT only: severe allergic reac Severe allergic reaction (e.g., anaphylaxis) a	Influenza, cell culture-based inactivated injectable [(ccllV4), Flucelvax® Quadrivalent]	 Severe allergi of ccIIV4 	c reaction (e.g., anaphylaxis) to any ccllV of any valency, or to any component ^a	 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 ccIV4, administer in medical catting under supervision of health care providen who care proceeding and manage causes 		
Pneumococcal conjugate (PCV13)	 Severe allergic reaction (e.g., anaphylaxis) ; Severe allergic reaction (e.g., anaphylaxis) t 				allergic reactions. May consult an allergist. • Moderate or severe acute illness with or without fever		
Pneumococcal polysaccharide (PPSV23)	Severe allergic reaction (e.g., anaphylaxis)	Influenza, recombinant injectable	Severe allergi RIV4	c reaction (e.g., anaphylaxis) to any RIV of any valency, or to any component ³ of	 Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine 		
Poliovirus vaccine, inactivated (IPV) Rotavirus (RV) [RV1 (Rotarix*), RV5 (RotaTeq*)]	Severe allergic reaction (e.g., anaphylaxis) ; Severe allergic reaction (e.g., anaphylaxis) ; Severe combined immunodeficiency (SCIE History of intussusception	[(ŘIV4), Flublok* Quadrivalent]			 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 		
Tetanus, diphtheria, and acellular pertussis (Tdap) Tetanus, diphtheria (Td)	Severe allergic reaction (e.g., anaphylaxis) ; For Tdap only: Encephalopathy (e.g., coma, attributable to another identifiable cause v Tdap	Influenza, live attenuated [LAIV4, Flumist* Quadrivalent]	 Severe allergi any egg-base Severe allergi Children age Anatomic or f Immunocom infection 	c reaction (e.g., anaphylaxis) after previous dose of any influenza vaccine (i.e., d IIV, ccIIV, RIV, or LAIV of any valency) c reaction (e.g., anaphylaxis) to any vaccine component ^a (excluding egg) 2 – 4 years with a history of asthma or wheezing unctional asplenia oromised due to any cause including, but not limited to, medications and HIV	 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 egg allergy with symptoms other than hives (e.g., angioedema, respiratory distress) or required epinephrine or another emergency medical intervention: Any influenza vaccine appropriate for age and health status may be administered. If using LAIV4 (which is egg based), administer in medical setting under supervision of health 		
Varicella (VAR) 1. When a contraindication is preset 2. When a precaution is present var	Severe allergic reaction (e.g., anaphylaxis) : Severe immunodeficiency (e.g., hematolog immunocedficiency, long-term immunosu immunocompromised) Pregnancy Family history of altered immunocompete immunocempetent nt, a vaccine should not be administered. K cination should generally be deferred but of the store of		Close contact environment Pregnancy Cochlear imp Active comm nasopharynx, Children and Received influ- hours, peram	s or caregivers of severely immunosuppressed persons who require a protected ant unication between the cerebrospinal fluid (CSF) and the oropharynx, nose, ear or any other cranial CSF leak adolescents receiving aspirin or salicylate-containing medications ienza antiviral medications oseltamivir or zanamivir within the previous 48 vir within the previous 5 days, or baloxavir within the previous 17 days	care provider who can recognize and manage severe allergic reactions. May consult an allergist. 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 preclation is present, val Guidelines for Immunization. ww Vaccination providers should che www.fda.gov/vaccines-blood-bio 	w.cdc.gov/vaccines/hcp/acip-recs/general-r ck FDA-approved prescribing information fc logics/approved-products/vaccines-license	1. When a contraindication is	present, a vaccir	ne should NOT be administered. Kroger A, Bahta L, Hunter P. ACIP General Best Pr	actice Guidelines for Immunization. www.cdc.gov/vaccines/hcp/acip-recs/general-recs/		

1. When a contraindication is present, a vaccine should NOT be administered. Kroger A, Bahta L, Hunter P. ACIP General Best Practice Guidelines for Immunization. www.cdc.gov/vaccines/hcp/acip-recs/general-recs/ contraindications.html

2. When a precaution is present, vaccination should generally be deferred but might be indicated if the benefit of protection 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

Clinical Scenario



Image courtesy of CDC/NCIRD

Medical and Vaccination History

- 14-year-old female (Date of birth-03/03/2008)
- Medical history: none
- Vaccination record shows the following:
 - DTaP on 6/29/2008 and 4/20/2013
 - Tdap on 7/20/2019
 - IPV on 6/29/2008 and 4/20/2013
 - Hepatitis B vaccine on 3/03/2008 and 6/29/2008

Parents report she received other vaccines and is up-to-date but have no record of those vaccinations

Vaccination History ¹

- Healthcare providers should only accept written, dated records as evidence of vaccination except for the following vaccines:
 - Influenza vaccine
 - Pneumococcal 23-valent polysaccharide vaccine (PPSV23)
- Self-reported doses of influenza vaccine and PPSV23 are acceptable because:
 - Influenza vaccine: the time period of recall is one year or less, making it very likely that correct recall will occur
 - PPSV23: a high frequency of vaccination leads to an increased rate of local reactions due to the reactogenicity of this vaccine.

1. Kroger A, Bahta L, Hunter P. General Best Practice Guidelines for Immunization. Best Practices Guidance of the Advisory Committee on Immunization Practices (ACIP). www.cdc.gov/vaccines/hcp/acip-recs/general-recs/downloads/generalrecs.pdf.



1. <u>Documented vaccination</u> <u>history</u>

- HepB, IPV, and DTaP/Tdap
- 2. <u>Routine vaccinations</u> <u>needed</u>
 - Influenza
 - HepB
 - IPV
 - MMR
 - VAR
 - HepA
 - Tdap
 - HPV
 - MenACWY
- 3. Other vaccinations needed
 - COVID-19



Recommended Child and Adolescent Immunization Schedule by Medical Indication, United States, 2022

Always use this table in conjunction with Table 1 and the Notes that follow.

					IN	DICATION				
			HIV infection	n CD4+ count ¹						
VACCINE	Pregnancy	Immunocom- promised status (excluding HIV infection)	<15% or total CD4 cell count of <200/mm ³	≥15% and total CD4 cell count of ≥200/mm ³	Kidney failure, end-stage renal disease, or on hemodialysis	Heart disease or chronic lung disease	CSF leak or cochlear implant	Asplenia or persistent complement component deficiencies	Chronic liver disease	Diabetes
Hepatitis B										
Rotavirus		SCID ²								
Diphtheria, tetanus, and acellular pertussis (DTaP)										
Haemophilus influenzae type b										
Pneumococcal conjugate										
Inactivated poliovirus										
Influenza (IIV4)										
Influenza (LAIV4)						Asthma, wheezing: 2–4yrs ³				
Measles, mumps, rubella	*									
Varicella	*									
Hepatitis A										
Tetanus, diphtheria, and acellular pertussis (Tdap)										
Human papillomavirus	*									
Meningococcal ACWY										
Meningococcal B										
Pneumococcal polysaccharide										
Dengue										
Vaccination according to routine schedule recommended	o the	Recommended for persons with an additio factor for which the vac would be indicated	nal risk a ccine r	/accination is recomi and additional doses necessary based on r condition or vaccine.	mended, F may be r medical o See Notes. o	Precaution—vaccine night be indicated if benefit of protection outweighs risk of adverse reaction	Contraindig recommen not be adm *Vaccinate	cated or not ded—vaccine should ninistered after pregnancy	No recomme applicable	endation/not

1 For additional information regarding HIV laboratory parameters and use of live vaccines, see the General Best Practice Guidelines for Immunization, "Altered Immunocompetence," at

www.cdc.gov/vaccines/hcp/acip-recs/general-recs/immunocompetence.html and Table 4-1 (footnote J) at www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html. 2 Severe Combined Immunodeficiency

³ LAIV4 contraindicated for children 2-4 years of age with asthma or wheezing during the preceding 12 months

Hepatitis B Vaccination Catch-up

Hepatitis B vaccination (minimum age: birth)

Birth dose (monovalent HepB vaccine only) Mother is HBsAg-negative:

 All medically stable infants ≥2,000 grams: 1 dose within 24 hours of birth

 Infants <2,000 grams: Administer 1 dose at chronological age 1 month or hospital discharge (whichever is earlier and even if weight is still <2,000 grams).

Mother is HBsAg-positive:

 Administer HepB vaccine and hepatitis B immune globulin (HBIG) (in separate limbs) within 12 hours of birth, regardless of birth weight. For infants <2,000 grams, administer 3 additional doses of vaccine (total of 4 doses) beginning at age 1 month. Test for HBsAg and anti-HBs at age 9-12 months. If HepB series is

delayed, test 1-2 months after final dose. Mother's HBsAg status is unknown:

 Administer HepB vaccine within 12 hours of birth, regardless of birth weight.

- For infants <2.000 grams, administer HBIG in addition to HepB vaccine (in separate limbs) within 12 hours of birth. Administer 3 additional doses of vaccine (total of 4 doses) beginning at age 1 month
- Determine mother's HBsAg status as soon as possible. If mother is HBsAg-positive, administer HBIG to infants ≥2,000 grams as soon as possible, but no later than 7 days of age.

Routine series

- 3-dose series at age 0, 1–2, 6–18 months (use monovalent Hep8) vaccine for doses administered before age 6 weeks)
- Infants who did not receive a birth dose should begin the series as soon as feasible (see Table 2).

- Administration of 4 doses is permitted when a combination vaccine containing HepB is used after the birth dose.
- Minimum age for the final (3rd or 4th) dose: 24 weeks
- Minimum intervals: dose 1 to dose 2: 4 weeks / dose 2 to dose 3: 8 weeks / dose 1 to dose 3: 16 weeks (when 4 doses are administered,

Catch-up vaccination

- Unvaccinated persons should complete a 3-dose series at 0, 1–2, 6 months.
- Adolescents age 11–15 years may use an alternative 2-dose schedule with at least 4 months between doses (adult formulation Recombivax HB* only).
- Adolescents age 18 years or older may receive a 2-dose series of Hep8 (Heplisav-B*) at least 4 weeks apart.
- Adolescents age 18 years or older may receive the combined HepA. and HepB vaccine, Twinrix*, as a 3-dose series (0, 1, and 6 months) or 4-dose series (3 doses at 0, 7, and 21-30 days, followed by a booster dose at 12 months).
- * For other catch-up guidance, see Table 2.

Special situations

 Revaccination is not generally recommended for persons with a normal immune status who were vaccinated as infants, children, adolescents, or adults.

- Post-vaccination serology testing and revaccination (if anti-HBs < 10mlU/mL) is recommended for certain populations, including:
- Infants born to HBsAg-positive mothers
- Hemodialysis patients
- Other immunocompromised persons

For detailed revaccination recommendations, see www.cdc.gov/

fluenza vaccination ninimum age: 6 months (IIV), 2 years (LAIV4), 3 years (recombinant influenza vaccine, RIV4))

Hepatitis B Vaccination Catch-up

Appendix Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2022

Vaccine	Contraindications ¹	Precautions ^a
Dengue (DEN4CYD)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of chemotherapy, congenital immunodeficiency, long-term immunosuppressive therapy or patients with HW infection who are severely immunocompromised) 	Pregnancy HV infection without evidence of severe immunosuppression Moderate or severe acute illness with or without fever
Diphtheria, tetanus, pertussis (DTaP) Tetanus, diphtheria (DT)	 Seven allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁴ For DTaP only: Encephalopathy (e.g., coma, decreased level of consciousness, prolonged seizures) not attributable to another identifiable cause within 7 days of administration of previous dose of DTP or DTaP 	 Gallain-Barré syndrome (GBS) within 6 veeks after previous dose of tetanus-toxida-containing vaccine History of Arthus-type hypersensitivity reactions after a previous dose of diphtheria-toxid or tetanus-toxida-containing vaccine, defer vaccination until at least 10 years have elapsed since the last tetanus-toxida-containing vaccine for DTa² only.¹Progressive neurologic disorder, including infantile spasms, uncontrolled epilepsy, progressive encrease neurologic disorder, including infantile spasms, uncontrolled epilepsy, progressive encrease illness with or without fever
Haemophilus influenzae type b (Hib)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ For Hiberity, ActHila, and PedvaadHB only: History of severe allergic reaction to dry natural latex Less than age 6 weeks 	Moderate or severe acute illness with or without fever
Hepatitis A (HepA)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ¹ including	Moderate or severe acute illness with or without fever
Hepatitis B (HepB)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component^a including yeast 	Moderate or severe acute illness with or without fever
Hepatitis A-Hepatitis B vaccine [HepA-HepB, (Twinrix*)]	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component^a including neomycin and yeast 	Moderate or severe acute illness with or without fever
Human papillomavirus (HPV)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ 	Moderate or severe acute illness with or without fever
Measles, mumps, rubella (MMR)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ⁴ Severe immunodeficiency (e.g., hematologic and solid rumors, receipt of chemotherapy, congenital immunochechecy, long-term immunosuppressive therapy or patients with HIV infection who are severely immunochecky in the severely immunochecky of altered imm	Recent (s11 months) receipt of antibody-containing blood product (specific interval depends on product) History of thrombocytopenia or thrombocytopenic purpura Need for tubercuin shin testing or interference agring a release assay (IGRA) testing Moderate or severe acute illness with or without fever
Meningococcal ACWY (MenACWY) (MenACWY-CRM (Menveo"); MenACWY-D (Menactra"); MenACWY-TT (MenQuadh")]	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ For MenACWV-D and Men ACWV-CRM only: severe allergic reaction to any diphtheria toxoid- or CRM197– containing vaccine For MenACWV-T only: severe allergic reaction to a tetanus toxoid-containing vaccine 	For MenACWY-CRM only: Preterm birth if less than age 9 months Moderate or severe acute illness with or without fever
Meningococcal B (MenB) (MenB-4C (Bessero*); MenB-FHbp (Trumenba*))	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component?	Pregnancy For MenB-4C only: Latex sensitivity Moderate or severe acute illness with or without fever
Pneumococcal conjugate (PCV13)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ Severe allergic reaction (e.g., anaphylaxis) to any diphtheria-toxoid- containing vaccine or its component¹ 	Moderate or severe acute illness with or without fever
Pneumococcal polysaccharide (PPSV23)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ^a	Moderate or severe acute illness with or without fever
Poliovirus vaccine, inactivated (IPV)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component*	Pregnancy Moderate or severe acute illness with or without fever
Rotavirus (RV) [RV1 (Rotarix*), RV5 (RotaTeq*)]	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ^s Severe combined immunodeficiency (SCID) History of intussusception	Altered immunocompetence other than SCID Chronic gastrointestinal disease RVI only: Spina bifda or bladder exstrophy Moderate or severe acute illness with or without fever
Tetanus, diphtheria, and acellular pertussis (Tdap) Tetanus, diphtheria (Td)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁸ For Tdap only: Encephalopathy (e.g., coma, decreased level of consciousness, prolonged seizures) not attributable to another identifiable cause within 7 days of administration of previous dose of DTP, DTaP; or Tdap 	 Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of tetanus-toxoid-containing vaccine History of Arthus type hypersensibility in nactions after a previous dose of diphthetia-toxoid—containing or tetanus-toxoid—containing vaccine, defor vaccination until at loast 10 years have elapsed since the last tetanus-toxoid—containing vaccine, defor vaccination until at loast 10 years have elapsed since the last tetanus-toxoid—containing vaccine, defore neurological disorder, uncontrolled seizures, or progressive encephalopathy until a treatment regimen has been established and the condition has stabilized Moderate or severe acute lines with or without fever
Varicella (VAR)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ⁸ Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of chemotherapy, congenital immunocempromised) environment of the severely immunocempromised) Pregnancy Family history of altered immunocompetence, unless verified clinically or by laboratory testing as	Recent (s11 months) receipt of antibody-containing blood product (specific interval depends on product) Receipt of specific antiviral drugs (acyclovir, famcidovir, or valacyclovir) 24 hours before vaccination (avoid use of these antiviral drugs for 14 days after vaccination) Use of aspirin or aspirin-containing products Moderate or severe acute illness with or without fever

Date of birth-03/03/2008

- Hepatitis B vaccination history • Dose 1: 03/03/2008
 - Dose 2: 6/29/2008

Plan

Give dose 3 today as final dose if no contraindication or precaution.

I. When a contraindication is present, a vaccine should NOT be administered. Kroger A, Bahta L, Hunter P. ACIP General Best Practice Guidelines for Immunization. www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html
I. When a procaution is present, vaccination should generally be deferred but might be indicated if the benefit of protection from the vaccine outweights 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

 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 bear available a www.fda.gov/vaccines-biodoc/biologics/approved-products/vaccines-licensed-use-united-states.

Poliovirus Vaccination Catch-up



The table below provides catch-up schedules and minimum intervals between doses for children whose vaccinations have been delayed. A vaccine series does not need to be restarted, regardless of the time that has elapsed between doses. Use the section appropriate for the child's age. Always use this table in conjunction with Table 1 and the Notes that follow.

darding	Minimum Ann for		Minimum Internet Patrony Decar		
Aaccine	Dose 1	Dave 1 to Dave 2	Minimum interval between Doses	Deve 3 to Deve 4	Davis data Davis
lepatitis B	Birth	4 weeks	Dose 2 to Dose 3 8 weeks and at least 16 weeks after first dose	Dose 3 to Dose 4	Dose 4 to Dos
totavirus	6 weeks Maximum age for first	4 weeks	mentium age for the tinal dose is 24 weeks 4 weeks maximum age for final dose is 8 months, 0 days		
iphtheria, tetanus, and reliefar nortuosis	dose is 14 weeks, 6 days. 6 weeks	4 weeks	4 weeks	6 months	6 months
ype b	6 weeks	No further doses needed if first dose was administered at age 15 months or older. 4 weeks if first dose was administered before the 1° birthdose. 8 weeks (as final dose) if first dose was administered at age 12 through 14 months.	No further doses needed If previous dose was administered at age 15 months or older 4 weeks 1 previous dose was PRP-T (ActHild', Prentacel', Hibert'), Vaaels' or unknown 8 weeks and go 12 through 59 months (and first dose was administered at younger than age 7 months and at least 1 previous dose was PRP-T (ActHild', Prentacel', Hibert'), Vaaels' or unknown 8 weeks and go 12 through 59 months (and first dose was administered at age 7 through 11 months; OR If current age is younger than 12 months and first dose was administered before the 1" birthday and second dose was administered at younger than 15 months; OR If both doses were PedvasHB* and were administered before the 1st birthday	8 weeks (as final dose) This dose only necessary for chalsen age 12 through 59 months who received 3 doses before the 1+ birthday.	
neumococcal conjugate	6 weeks	No further doses needed for healthy children if first dose was administened at age 24 months or older 4 weeks 1° birthday 8 weeks (as final dose for healthy children) 8 first dose was administered at the 1° birthday or after	No further doses needed for healthy children if previous dose was administered at age 24 months or older 4 weeks if current age is younger than 12 months and previous dose was administered at <7 months old 8 weeks (as final dose for healthy children) 10 previous dose was administered battween 7-11 months (wait until at least 12 months old); OR if current age is 12 months or older and at least 1 dose was administered before age 12 months	B weeks (as final dose) This dose only necessary for children age 12 through 59 months who received 3 dose before age 12 months or for children at high risk who received 3 doses at any age.	
sactivated poliovirus	6 weeks	4 weeks	4 weeks If current age is <4 years 6 months (as final dose) If current age is 4 years or older	6 months (minimum age 4 years for final dose)	
leasles, mumps, rubella	12 months	4 weeks			
aricella	12 months	3 months			
orotitis A	12 months	6 months			
leningococcal ACWY	2 months MenACWY-CRM 9 months MenACWY-D 2 years MenACWY-TT	8 weeks	See Notes	See Notes	
			Children and adolescents age 7 through 18 years		
leningococcal ACWY	Not applicable (N/A)	8 weeks			
etanus, diphtheria; etanus, diphtheria, and cellular pertussis	7 years	4 weeks	4 weeks If first does of DTaP/DT was administered before the 1 st birthday 6 months (as final dose) If first dose of DTaP/DT or Tdap/Id was administered at or after the 1 st birthday	6 months if first dose of DTaP/DT was administered before the 1° birthday	
uman papillomavirus	9 years	Routine dosing intervals are recommended.			
epatitis A	N/A	6 months			
epatitis B	N/A	4 weeks	8 weeks and at least 16 weeks after first dose		
activated poliovirus	N/A	4 weeks	6 months A fourth dose is not necessary if the third dose was administered at age 4 years or older and at least 6 months after the previous dose.	A fourth dose of IPV is indicated if all provious doses were administered at <4 years or if the third dose was administered <6 months after the second dose.	
Varicella	N/A	3 months if younger than age 13 years. 4 weeks if age 13 years or older			

Date of birth-03/03/2008

ullet

Inactivated Poliovirus (IPV) vaccination history O Dose 1: 4/25/2008 O Dose 2: 4/20/2013

IPV Vaccination Catch-up

Notes

Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2022

Chronic liver disease, alcoholism

ign 6 The years No history of PPSV23 T dose PPSV23 (at least 5 weeks after any prior. PCV 13 dose

incomplete sines – Not having received all doses in either the tecommended vertex or an age-appropriate catch-up series See Tables 8.9, and 11 in the AGP preumococcial vaccine recommendations (Newsord Graph Immini pdf. mm5911 pdf. (Se complete schedule details.

Poliovirus vaccination (minimum age: 6 weeks)

Routine vaccination

 4-dose series at ages 2, 4, 6–18 months, 4–6 years; administer the final dose on or after age 4 years and at least 6 months after the previous dose.

 4 or more doses of IPV can be administered before age 4 years when a combination vaccine containing IPV is used. However, a dose is still recommended on or after age 4 years and at least 6 months after the

Catch-up vaccination

 In the first 6 months of life, use minimum ages and intervals only for travel to a polio-endemic region or during an outbreak.

 IPV is not routinely recommended for U.S. residents age 18 years or older.

Series containing oral polio vaccine (OPV), either mixed OPV-IPV or OPV-only series:

 Total number of doses needed to complete the series is the same as that recommended for the U.S. IPV schedule. See www.cdc.gov/ mmwr/volumes/66/wr/mm6601a6.htm?s_%20cid=mm6601a6_w.

 Only trivalent OPV (tOPV) counts toward the U.S. vaccination requirements.

 Doses of OPV administered before April 1, 2016, should be counted (unless specifically noted as administered during a campaign).

 Doses of OPV administered on or after April 1, 2016, should not be counted.

 For guidance to assess doses documented as "OPV," see www.cdc.gov/mmwr/volumes/66/wr/mm6606a7.htm?s_ cid=mm6606a7_w.

For other catch-up guidance, see Table 2.

minimum age: 6 week

Routine vaccination

Rotarix*: 2 dose veries at age 2 and 4 months;

· RotaTeg': 3-dose series at age 2, 4, and 6 ma

 If any dote in the series is either RotaTeq* or unknown, default to 1 dote series.

Catch-up vaccinatio

Do not start the series on or after age 15 weeks, 0 days
 The maximum age his the final doise is it mumbs, 0 days
 The other catch-colouidance, see Table 2.

Tetanus, diphtheria, and pertussis (Tdap) vaccination

(minimum age: 11 years for routine vaccination) 7 years for catch-up vaccination)

Routine vaccination

Adolescents age 11-12 years: % dase Tdap Pregnancy: 7 dose Tdap during each pregnancy, proferably and part of motioned age 27.2

Tdap may be administered regardless of the interval since the last tetahus, and diphthetia tisso discontaining vaccine.

atch-up vaccination

Adolescents age 13–18 years who have not received Tdap: Educe Tdap, then Td or Tdap booster every 10 years.

Persons age 7-18 years not fully vaccinated with DTaP : I dose Tdap as part of the catching series (prefectably the first dose) if additional dosenate needed, we Td or Tdap.

dap administered at age 7-10 years:

Children age 7-9 years who receive Tdap should receive the portine Tdap dose at age 11-12 years.

Children age 10 years who receive Tdat: do not need the routine dap dove at ane 11-12 years.

aP inadvertently administered on or after age 7 years

Children age 7-9 years. DTaP may count as part of catch-up

ieries. Administer toutine Idap dose al age 11-12 years.

Independent of the years, count dolle of this as the address and Independent

lot other catch-up guidance, cee Table

Special situation

Sound management in personance (?) years or older with hadovy 3 or more down of tetative toxold-tentaining vaccine. For cleans dimmor wounds, administer Tolgo or Tol it more than 10 years nee last dose of tetanus toxold-containing vaccine, for all other bands, administer Tolap or Tol if more than 6 years, since last dose of tenus, toxold-containing vaccine. Tolgo is preferred for persons age years or older who have not previously inclusive did to be tablicity is unknown, if a situations toxold-containing vaccine in dirusted for a previously inclusion to be done of the attent or a previously inclusion to be done of the attent or a previously inclusion of the teta of the second dirusted for a previously inclusion to be done of the second containing vaccine in dirusted for a previously inclusion.

For detailed internation, see www.u.ac.gov/mmes/volumes/binwr/ mmes/stats.html

Ny volucitathed – Si valid dipses of DTaP OR 4 valid doses of DTaP 8 ose 4 was administered at age 4 years or order

Varicella vaccination

(minimum age: 12 months)

Routine vaccination

* 2 dove venes at age 32-15 months, 4-6 years -* VAB or MMRV may be administered*

 Dove 2 may be administered as early as 3 months after dove 1 a dove madvertently administered after at least 4weeks may be counted as wild!

*Note: For doxe 1 in children age 12:47 months, it is recommended to administer WWR and varicelia vaccines separately. MWRV may be used if parents of caregovers express a preference.

Catch-up vaccination

 Ensure persons age 1.10 years without evidence of miniumity (see M2030at yourside gas ministicatif (consolid) pdf) have a 2-dose sector;

Age 7-12 years routine interval: 3 months (a done inadvertently sommarized after at lead 4 weeks may be counted as valid)

nteruzi a weeku

The musicisum age for use of MMRV is 12 years.

IPV Vaccination Catch-up

Appendix	Recommended Child and Adolescent Immunization Sched	ule for ages 18 years or younger, United States, 2022
Vaccine	Contraindications ¹	Precautions ²
Dengue (DEN4CYD)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of chemotherapy, congenital immunodeficiency, long- term immunosuppressive therapy or patients with HN infection who are severely immunocompromised) 	Pregnancy HV Infection without evidence of severe immunosuppression Moderate or severe acute illness with or without fever
Diphtheria, tetanus, pertussis (DTaP) Tetanus, diphtheria (DT)	 Seven allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁸ For DTaP only: Encephalopathy (e.g., coma, decreased level of consciousness, prolonged seizures) not attributable to another identifiable cause within 7 days of administration of previous dose of DTP or DTaP 	 Guillain-Band syndrome (GBS) within 6 weeks after previous dose of tetanus-toxoid-containing vaccine. History of Arthus-type hypersensitivity reactions after a previous dose of diphthenia-toxoid-containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid-containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid-containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid-containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid-containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid-containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid-containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid-containing vaccine; defer vaccination; defer
Haemophilus influenzae type b (Hib)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁴ For Hiberity, ActHilg, and PedvaadHB only: History of severe allergic reaction to dry natural latex Less than age 6 weeks 	Moderate or severe acute illness with or without fever
Hepatitis A (HepA)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ including neomycin 	Moderate or severe acute illness with or without fever
Hepatitis B (HepB)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ including yeast For Heplisav-B only: Pregnancy 	Moderate or severe acute illness with or without fever
Hepatitis A-Hepatitis B vaccine [HepA-HepB, (Twinrix")]	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ including neomycin and yeast 	Moderate or severe acute illness with or without fever
Human papillomavirus (HPV)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁴ 	Moderate or severe acute illness with or without fever
Measles, mumps, rubella (MMR)	Seven interminedeficiency (e.g., nanaphylaxis) after a previous dose or to a vaccine component ⁴ Seven immunodeficiency, (e.g., hermatologic and solid tumors, receipt of chemotheragy, congenital immunocomponised) Pregnancy Previous and the sevent of the seven	elseent (s11 moorths) receipt of antibody-containing blood product (specific interval depends on product) History of thrombocytopenia or thrombocytopenia purpura Need for tuberculin skin testing or interferon-gamma release assay (IGRA) testing Moderate or severe acute illness with or without fever
Meningococcal ACWY (MenACWY) [MenACWY-CRM (Menveo*); MenACWY-D (Menactra*); MenACWY-TT (MenQuadfi*)]	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ For MenACWr-D and Men ACWP-CRM only: severe allergic reaction to any dipitheria toxoid- or CRM197– containing vaccine For MenACWP-TT only: severe allergic reaction to a tetanus toxoid-containing vaccine 	For MenACWY-CRM only: Preterm birth if less than age 9 months Moderate or severe acute illness with or without fever
Meningococcal B (MenB) [MenB-4C (Bexsero*); MenB-FHbp (Trumenba*)]	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component?	Pregnancy For MenB-4C only: Latex sensitivity Moderate or severe acute illness with or without fever
Pneumococcal conjugate (PCV13)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ Severe allergic reaction (e.g., anaphylaxis) to any diphtheria-toxoid- containing vaccine or its component¹ 	Moderate or severe acute illness with or without fever
Pneumococcal polysaccharide	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ 	Moderate or severe acute illness with or without fever
Poliovirus vaccine, inactivated (IPV)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component^s 	Pregnancy Moderate or severe acute illness with or without fever
Rotavirus (RV) [RV1 (Rotarix*), RV5 (RotaTeq*)]	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ^s Severe combined immunodeficiency (SCID) History of intussusception	Altered immunocompetence other than SCID Chronic gastrointestinal disease RVI only: Spina bifida or bladder exetrophy Moderate or severe acute illness with or without fever
Tetanus, diphtheria, and acellular pertussis (Tdap) Tetanus, diphtheria (Td)	 Seven allergic maction (e.g., anaphyliatis) after a previous dose or to a vaccine component⁸ For Tidap only: Encephalopathy (e.g., coma, decreased level of consciousness, prolonged seizures) not attributable to another identifiable cause within 7 days of administration of previous dose of DTP, DTaP, or Tidap 	 Gelfain-Band syndrome (GRS) within 6 weeks after a previous does of tetranus-travid-containing vaccine History of Arthus type hypersensitivity reactions after a previous does of diphthetia-toxoid— containing or tetanus-toxoid— containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid— containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid—containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid—containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid—containing vaccine; defer vaccination until at least 10 years have elapsed rortGap only. Progressive or unstable neurological disorder, uncontrolled seizures, or progressive encephalopathy until a treatment regimen has been established and the condition has stabilized - Moderater or severe acute liness with or without fiver
Varicella (VAR)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component* Severe immunodeficiency (e.g., hematologic and solid rumors, receipt of chemotherapy, congenital immunodeficiency, long-term immunosuppressive therapy or patients with HV infection who are severely immunodeficiency and the severely immunocomponentsed) Fegnancy of altered immunocompetence, unless verified clinically or by laboratory testing as immunocompetent	Recent (s 11 months) receipt of antibody containing blood product (specific interval depends on product) Receipt of specific antiviral drugs (acyclovit, famicidovit, or valacyclovit) 24 hours before vaccination (avoid use of these antiviral drugs for 14 days after vaccination) Use of aspirin or aspirin-containing products Moderate or severe acute illness with or without fever

1. When a contraindication is present, a vaccine should NOT be administered. Kroger A, Bahta L, Hunter P. ACIP General Best Practice Guidelines for Immunization, www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html

 When a procaution is present, vaccination should generally be deferred but might be indicated if the benefit of protection from the vaccine outweighs the risk for an adverse reaction. Kroger A, Bahta¹L, Hunter P. ACIP General Best Practice Control the rest for an adverse reaction.

Guidelines for Immunization. www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html

 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.lda.gov/vaccines-blood-blologics/approved-products/vaccines-licensed-use-united-states.

- Date of birth-03/03/2008
- IPV vaccination history

 Dose 1: 4/25/2008
 Dose 2: 4/20/2013

<u>Plan</u>

Give dose 3 today as final dose if no contraindication or precaution.

Tdap Vaccination Catch-up

Table 2 Recommended Catch-up Immunization Schedule for Children and Adolescents Who Start Late or Who Are More than 1 Month Behind, United States, 2022

the table below provides catch-up schedules and minimum intervals between doses for children whose vaccinations have been delayed. A vaccine series does not need to be restarted, regardless of the time that has apsed between doses. Use the section appropriate for the child's age. Always use this table in conjunction with Table 1 and the Notes that follow.

			Children age 4 months through 6 years		
Vaccine	Minimum Age for		Minimum Interval Between Doses		
	Dose 1	Dose 1 to Dose 2	Dose 2 to Dose 3	Dose 3 to Dose 4	Dose 4 to Dose !
Hepatitis B	Birth	4 weeks	8 weeks and at least 16 weeks after first dose minimum age for the final dose is 24 weeks		
Rotavirus	6 weeks Maximum age for first dose is 14 weeks, 6 days.	4 weeks	4 weeks maximum age for final dose is 8 months, 0 days		
Diphtheria, tetanus, and acellular pertussis	6 weeks	4 weeks	4 weeks	6 months	6 months
Haemophilus influenzae type b	6 weeks	No further dose needed if first dose was administered at age 15 months or older. 4 weeks if first dose was administered before the 1+ birthday. 8 weeks (as final dose) if first dose was administered at age 12 through 14 months.	No further doses needed If previous dose was administered at age 15 months or older 4 weeks If current age is younger than 12 months end first dose was administered at younger than age 7 months end at least 1 previous dose was PBP-T (ActHib [*] , Pentacel [*] , Hiberix [*]), Vaselis [*] or unknown 8 weeks and age 12 through 59 months (as final dose) If current age is younger than 12 months end first dose was administered at age 7 through 11 months; OR If current age is 12 through 59 months end first dose was administered before the 1 st birthday and second dose was administered at younger than 15 months; OR If both doses were PedvaaHB [*] and were administered before the 1st birthday	8 weeks (as final dose) This dose only necessary for childen age 12 through 59 months who received 3 doses before the 1* birthday.	
Pneumococcal conjugate	6 weeks	No further doses needed for healthy children if first dose was administered at age 24 months or older 4 weeks 14 first dose was administered before the 1+ birthday 8 weeks (as final dose for healthy children) 14 birthday or after	No further doses needed for healthy children if previous dose was administered at age 24 months or older 4 weeks if current age is younger than 12 months and previous dose was administered at <7 months old 8 weeks (as final dose for healthy children) if previous dose was administered between 7–11 months (wait until at least 12 months old); OR if current age is 12 months or older and at least 1 dose was administered before age 12 months	8 weeks (as final dose) This dose only necessary for childen age 12 through 59 months who received 3 doses before age 12 months or for childen at high risk who received 3 doses at any age.	
Inactivated poliovirus	6 weeks	4 weeks	4 weeks 8 current age is <4 years 6 months (as final dose) 8 current age is 4 years or older	6 months (minimum age 4 years for final dose)	
Measles, mumps, rubella	12 months	4 weeks			
Varicella	12 months	8 months			
Heratitic A	12 months	6 months			
Meningococcal ACWY	2 months MenACWY-CRM 9 months MenACWY-D 2 years MenACWY-TT	8 weeks	See Notes	See Notes	
	Contraction of the local division of the loc		Children and adolescents age 7 through 18 years		
Meningococcal ACWY	Not applicable (N/A)	8 weeks			-
etanus, diphtheria; etanus, diphtheria, and cellular pertussis	7 years	4 weeks	4 weeks if first dose of DTaP/DT was administered before the 1 st birthday 6 months (as final dose) if first dose of DTaP/DT or Totao/Nd was administered at or after the 1 st birthday	6 months If first dose of DTaP/DT was administered before the 1° birthday	
Human papilomawirus	9 years	Routine dosing intervals are recommended.			
Hepatitis A	N/A	6 months			
Hepatitis B	N/A	4 weeks	8 weeks and at least 16 weeks after first dose		
inactivated poliovirus	N/A	4 weeks	6 months A fourth dose is not necessary if the third dose was administered at age 4 years or older and at least 6 months after the previous dose.	A fourth dose of IPV is indicated if all previous doses were administered at <4 years or if the third dose was administered <6 months after the second dose.	
Measles, mumps, rubella	N/A	4 weeks			
Varicella	N/A	3 months if younger than age 13 years. 4 weeks if age 13 years or older			
Dengue	9 years	6 months	6 months		

Date of birth-03/03/2008

DTaP/Tdap vaccination history

- Dose 1: 6/29/2008 (DTaP)
- Dose 2: 4/20/2013 (DTaP)
- Dose 3: 7/20/2019 (Tdap)

Tdap Vaccination Catch-up



Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2022

Chronic liver disease, alcoholism,

Aget 19 years • No history of PPSV23: 1 date PPSV23 (at least 8 weeks after any prior • PCV13 dose

Incomplete sames — Not having received all doves in either the tecommended series of an age appropriate catch-up series. See Tables 8, 9 and 11 in the ACP pneumococcial vaccine recommendations inswire dagos throng pdf. (hor complete ochecture details.)

Poliovirus vaccination (minimum age: 6 weeks)

Routine vaccination

- 4 4-dose series at ages 2, 4, 6, 18 months, 4, 6 years, administer the Final dose on or alter age 4 years and at least 6 months after the previous dose.
- 4 or more dayse, of IPX can be administered before age 4 years when an combination success containing IPV is used. However, a days is still recommended on or after age 4 years and at least 6 months after the previous dose.

Catch-up vaccination

- In the first 6 months of life, use minimum ages and intervals drive for travel to a policil endemy, region or during an outbreak
- IPV is not notlinely recommended for U.S. residents age 18 years or older

Series containing oral polio vaccine (OPV), either mixed OPV (PV o OPV only series

- Initial mumber of doves treated to complete the same, a the same as that recommended for the U.S. IPV schedule See www.ide.gov mmwr.velumes.col. or primition schemich, in 2016 mmHR01ab.
- Only Invalent, OPV (OPV) counts toward the U.S. vaccination inquinements.
- Quies of OPV administered before April 1, 2016, should be counted uplies, specifically noted as administered during a campaignt.

Doses of OPV administered on or after April 3, 2016, should no be counted

- For guidance to assess documented as "OPV" see www.tdc.gov.mmwv.stillum.in/GG/Willimmt60ball.html
- . The others is him multiple

Rotavirus vaccination

Routine vaccination

- Rotarix1:2 dose Series at age 2 and 4 months
- If any dove in the series is either RotaTeq* or unknown, default to Area series

atch-up vaccination

Do not start the series on or after age 15 weeks, 0 days. The maximum age for the final dose is it months, it days for other catch sor guidance, see Table 2.

Tetanus, diphtheria, and pertussis (Tdap) vaccination

(minimum age: 11 years for routine vaccination, 7 years for catch-up vaccination)

Routine vaccination

 Adolescents age 11–12 years: 1 dose Tdap
 Pregnancy: 1 dose Tdap during each pregnancy, preferably in early part of gestational weeks 27–36.

* Tdap may be administered regardless of the interval since the last

Catch-up vaccination

 Adolescents age 13–18 years who have not received Tdap: 1 dose Tdap, then Td or Tdap booster every 10 years

 Persons age 7–18 years not fully vaccinated' with DTaP: 1 dose Tdap as part of the catch-up series (preferably the first dose); if additional doses are needed, use Td or Tdap.

- Tdap administered at age 7–10 years:
- Children age 7–9 years who receive Tdap should receive the routine Tdap dose at age 11–12 years.

 Children age 10 years who receive Tdap do not need the routine Tdap dose at age 11–12 years.

• DTaP inadvertently administered on or after age 7 years: - Children age 7–9 years: DTaP may count as part of catch-up

series. Administer routine Tdap dose at age 11–12 years. Children age 10–18 years: Count dose of DTaP as the adolescent Tdap booster.

For other catch-up guidance, see Table 2.

Special situations

 Wound management in persons age 7 years or older with history of 3 or more doses of tetanus-toxoid-containing vaccine: For clean and minor wounds, administer Tdap or Td if more than 10 years since last dose of tetanus-toxoid-containing vaccine; for all other wounds, administer Tdap or Td if more than 5 years since last dose of tetanus-toxoid-containing vaccine. Tdap is preferred for persons age 11 years or older who have not previously received Tdap or whose Tdap history is unknown. If a tetanus-toxoid-containing vaccine is indicated for a pregnant adolescent, use Tdap.

 For detailed information, see www.cdc.gov/mmwr/volumes/69/wr/ mm6903a5.htm.

*Fully vaccinated = 5 valid doses of DTaP OR 4 valid doses of DTaP if dose 4 was administered at age 4 years or older

Varicella vaccination

Boutine vaccination

+2 dose series at age 12. (5 months 4: 6 years

We or WMRV may be administered.

 Dose 2 may be administered as early as 3 months after dose 1 a dose madvertently administered after at least tweeky may be counted as word.

«Note: For dow! Linichidnen app 12:42 months, it is recommended to judminister MMR and varicelia vaccinet separately. IMRV may be used, if parents of categories express a preference.

atch-up vaccination

 Ensure personscape 1: 10 years without evidence of minutery see MMRRar (possible gas minus bull (terrision per) have a 2-dose series.

Age 7-12 years including interval 3 months is done inadvertently administered after at least 4 weeks may be educted as valid Age 13 years and older to the interval 4, il weeks intervent

riferan Aweeku

The musicinum age for use of MMRV is 12 years.

Tdap Vaccination Catch-up

Vaccine	Contraindications ¹	Precautions ²					
Dengue (DEN4CYD)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁴ Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of chemotherapy, congenital immunodeficiency, long- term immunosuppressive therapy or patients with HV infection who are severely immunocompromised) 	Pregnancy HV infection without evidence of severe immunosuppression Moderate or severe acute illness with or without fever					
Xiphtherla, tetanus, pertussis (DTaP) letanus, diphtherla (DT)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁴ For DTaP only: Encephalopathy (e.g., coma, decreased level of consciousness, prolonged seizures) not attributable to another identifiable cause within 7 days of administration of previous dose of DTP or DTaP 	 Guillain-Baine syndrome (GBS) within 6 weeks after previous dose of tistanus toxoid—containing vaccine. History of Arthus-type hypersensitivity reactions after a previous dose of dipthenia-toxoid—containing vaccine; defer vaccination until at least 10 years have elapsed since the l transit toxoid—containing vaccine; defer vaccination until at least 10 years have elapsed since the l transit toxoid—containing vaccine; defer vaccination until at least 10 years have elapsed since the l transit toxoid—containing vaccine; defer vaccination until at least 10 years have elapsed since the l transit toxoid—containing vaccine; defer vaccination until at least 10 years have elapsed since the l transit toxoid—containing vaccine; defer to the loss of the last since the last loss of the last since the loss of the loss of the last los					
laemophilus influenzae type b (Hib)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁸ For Hiberix, ActHeb, and PedvadHB only: History of severe allergic reaction to dry natural latex Less than age 6 weeks 	Moderate or severe acute illness with or without fever					
epatitis A (HepA)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ including neomycin 	Moderate or severe acute illness with or without fever					
epatitis B (HepB)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ¹ including yeast For Heplisav-Bonly: Pregnancy	Moderate or severe acute illness with or without fever					
epatitis A- Hepatitis B vaccine lepA-HepB, (Twinrix*)]	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ including neomycin and yeast 	Moderate or severe acute illness with or without fever					
uman papillomavirus (HPV)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ 	Moderate or severe acute illness with or without fever					
Aeasles, mumps, rubella (MMR)	Severe allergic reaction (e.g., anaphylaski) after a previous dose or to a vaccine component ⁴ Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of d'hemotherapy, congenital immunodeficiency, long-term immunosuppressive therapy or patients with HIV infection who are severely immunocompromised) Pregnancy Family history of altered immunocompetence, unless verified clinically or by laboratory testing as immunocompetent	Recent (s11 months) receipt of antibody-containing blood product (specific interval depends on product) History of thrombocytopenic or thrombocytopenic gumpura Need for tuberculin skin testing or interferon-gamma release assay (IGRA) testing Moderate or severe acute illness with or without fever					
leningococcal ACWY (MenACWY) MenACWY-CRM (Menveo*); IenACWY-D (Menactra*); IenACWY-TT (MenQuadfi*)]	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ For MenACW/O and Men ACWY-CRM only: severe allergic reaction to any diphtheria toxoid – or CRM197– containing vaccine For MenACW/T only: severe allergic reaction to a tetanus toxoid-containing vaccine 	For MenACWI-CRM only: Preterm birth if less than age 9 months Moderate or severe acute illness with or without fever					
leningococcal B (MenB) AenB-4C (Bessero"); lenB-FHbp (Trumenba*)]	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁸ 	Pregnancy For MenB-4C only: Latex sensitivity Moderate or severe acute illness with or without fever					
neumococcal conjugate (PCV13)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ Severe allergic reaction (e.g., anaphylaxis) to any diphtheria-toxoid containing vaccine or its component¹ 	Moderate or severe acute illness with or without fever					
neumococcal polysaccharide PSV23)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ 	Moderate or severe acute illness with or without fever					
oliovirus vaccine, inactivated (IPV)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁴ 	Pregnancy Moderate or severe acute illness with or without fever					
otavirus (RV) [RV1 (Rotarix*), V5 (RotaTeq*)]	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁴ Severe combined immunodeficiency (SCID) History of intussusception 	Altered immunocompetence other than SCID Chronic gastrointestinal disease W1 only: Spina bifda or bladder exetrophy Moderate or severe acute illness with or without fever					
etanus, diphtheria, and acellular ertussis (Tdap) 'etanus, diphtheria (Td)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁴ For Tdap only: Encephalopathy (e.g., coma, decreased level of consciousness, prolonged seizures) not attributable to another identifiable cause within 7 days of administration of previous dose of DTP, DTaP, or Tdap 	 Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of tetanus-toxioid-containing vaccine History of Arthis type hypersensitivity reactions after a previous dose of diphtheria-toxoid—containing or tetanus-toxioid—containing vaccine; der tranus-toxoid-containing vaccine; for Tdap only: Progressive or unstable neurological disorder, uncontrolled seizures, or progressive encephalopathy until a treatment regimen has been established and the condition has stabilized Moderate or severe acute illness with or without fever 					
ricella (VAR)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ⁶ Severe inmunodeficiency (e.g., hematologic and solid tumors, receipt of chemotherapy, congenital immunodeficiency, long-term inmunosuppressive therapy or patients with HV infection who are severely immunocompromised) Pregnancy Family history of altered immunocompetence, unless verified clinically or by laboratory testing as	Recent (<11 months) receipt of antibody-containing blood product (specific interval depends on product Receipt of specific antiviral drugs (acyclovir, famcidovir, or valacyclovir) 24 hours before vaccination (avoid use of these antiviral drugs for 14 days after vaccination) Use of aspirin or aspirin-containing products Moderate or severe acute illness with or without fever					

- Date of birth-03/03/2008
- DTaP/Tdap vaccination history
 - Dose 1: 6/29/2008 (DTaP)
 - Dose 2: 4/20/2013 (DTaP)
 - Dose 3: 7/20/2019 (Tdap)

<u>Plan</u>

Give dose 4 (Tdap) today if no contraindication or precaution.

Guidelines for Immunization, www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.htr

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.

Catch-up Vaccination Job Aid

Vaccine Catch-Up Guidance

CDC has developed catch-up guidance job aids to assist health care providers in interpreting Table 2 in the child and adolescent immunization schedule.

- <u>Pneumococcal Conjugate Vaccine (PCV) Catch-Up</u> <u>Guidance for Children 4 Months through 4 Years of</u> <u>Age</u> [3 pages]
- *Haemophilus influenzae* type b-Containing Vaccines Catch-Up Guidance for Children 4 Months through 4 Years of Age
 - <u>Hib vaccine products: ActHIB, Pentacel, Hiberix,</u> or unknown [3 pages]
 - <u>Hib vaccine products: PedvaxHIB vaccine only</u>
 [2 pages]
- <u>Diphtheria-, Tetanus-, and Pertussis-Containing</u>
 <u>Vaccines Catch-Up Guidance for Children 4 Months</u>
 <u>through 6 Years of Age</u> [2 pages]

- Inactivated Polio Vaccine (IPV) 📕 [2 pages]
- <u>Tetanus-, Diphtheria-, and Pertussis-Containing</u> <u>Vaccines Catch-Up Guidance for Children 7 through</u> <u>9 Years of Age</u> [2 pages]
- <u>Tetanus-, Diphtheria-, and Pertussis-Containing</u>
 <u>Vaccines Catch-Up Guidance for Children 10 through</u>
 <u>18 Years of Age</u>

*Child and adolescent Immunization schedule <u>https://www.cdc.gov/vaccines/schedules/hcp/schedule-changes.html</u>

Tdap Catch-up Guidance for Ages 10-18 years

Catch-Up Guidance for Children 10 through 18 Years of Age

Tetanus-, Diphtheria-, and Pertussis-Containing Vaccines: Tdap/Td

IF current age is	AND # of previous doses of DTaP, DT, Td, or Tdap is	AND	AND	AND	THEN	Next dose due		
10 through 18 years	3	Dose 1 was given before 12 months of age Dose 1 was given at 12 months of age or older	<mark>lt has been at</mark> least 6 calendar	Any dose was Tdap ¹	Give Dose 4 (Td or Tdap) today ²	Give Td or Tdap 10 years		
			Dose 3	No dose was Tdap ³	Give Dose 4 (Tdap) today	after Dose 4		
			It has not been 6 calendar	Any dose was Tdap ¹	No dose today	Give Dose 4 (Td or Tdap) at least 6 calendar months after Dose 3 ²		
			months since Dose 3	No dose was Tdap ³	No dose today	Give Dose 4 (Tdap) at least 6 calendar months after Dose 3		
			No dose was Tdap¹	→	Give Dose 4 (Tdap) today	Give Td or Tdap 10 years after Dose 4		
			Any dose was Tdap²	→	No dose today	Give Td or Tdap 10 years after Dose 3		
			No Tdap was given after 7 th birthday	→	Give a dose of	Give Td or Tdap 10 years		
	4	→	Any dose of Tdap was given	No Tdap was given after 10 th birthday	Tdap today⁴	after Tdap dose		
			at age 7 years	Tdap was				

*Child and adolescent Immunization schedule Immunization Schedule Changes | CDC

DTaP/Tdap vaccination history

- Dose 1 (DTaP): 6/29/2008
- Dose 2 (DTaP): 4/20/2013
- Dose 3 (Tdap): 7/20/2019

<u>Plan</u>

Give dose 4 (Tdap) today if no contraindication or precaution.

Other Catch-up Vaccinations Needed



Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2022

These recommendations must be read with the notes that follow. For those who fall behind or start late, provide catch-up vaccination at the earliest opportunity as indicated by the green bars. To determine minimum intervals between doses, see the catch-up schedule (Table 2).

Vaccine	Birth	1 mo	2 mos	4 mos	6 mos	9 mos	12 mos	15 mos	18 mos	19–23 mos	2-3 yrs	4–6 yrs	7-10 yrs	11-12 y	13-15 yrs	16 yrs	17–18 yrs
Hepatitis B (HepB)	1ª dose	4 2 nd	dose •		•		3 st dose										
Rotavirus (RV): RV1 (2-dose series), RV5 (3-dose series)			1 ^e dose	2 nd dose	See Notes												
Diphtheria, tetanus, acellular pertussis (DTaP <7 yrs)			1ª dose	2 ^{ed} dose	3 st dose			∢ 4 [®] d	iose•			5 th dose					
Haemophilus influenzae type b (Hib)			1*dose	2 ^{wl} dose	See Notes		def def all all all all all all all all all al	* dose									
Pneumococcal conjugate (PCV13)			1 ^e dose	2 nd dose	3 ^{si} dose		4 4 th (dose>									
Inactivated poliovirus (IPV <18 yrs)			1ª dose	2 ^{nt} dose	•		— 3 ^{ul} dose -					4ª dose					
Influenza (IIV4)							,	Innual vacci	ination 1 or	2 doses			_	Ann	al vaccination	dose on	y .
Influenza (LAIV4)											Annua 1 c	l vaccinatio r 2 doses	n (Ann	alvaccination	dose on	y.
Measles, mumps, rubella (MMR)					Seel	Notes	← 1°	dose>				2 ^{se} dose					
Varicella (VAR)							<1 ⁰	iose•				2 st dose			9.0		
Hepatitis A (HepA)					Seel	Notes		2-dose serie	rs, See Note	s							
Tetanus, diphtheria, acellular pertussis (Tdap ≥7 yrs)														1 dose			
Human papillomavirus (HPV)														See Notes			
Meningococcal (MenACWY-D ≥9 mos, MenACWY-CRM ≥2 mos, MenACWY-TT ≥2years)								See Notes	11 1					1º dos		f" dose	
Meningococcal B (MenB-4C, MenB- FHbp)															See No	5	
Pneumococcal polysaccharide (PPSV23)														See Not			
Dengue (DEN4CYD; 9-16 yrs)													Se	eropositiv	in endemic a See Notes)	as only	
Range of recommended ages for all children	Range of r	ecommend up vaccinati	ed ages on	Rar for	nge of recor	nmended a h-risk group	ges s	Recommission	mended vac in in this ag	cination ge group	R	ecommende n shared clin	ed vaccinatio	on based n-making	Nor	ecommer	ndation/

- 1. <u>Documented vaccination</u> <u>history</u>
 - HepB, IPV, and DTaP/Tdap
- 2. <u>Routine vaccinations</u> <u>needed</u>



Image courtesy of CDC/NCIRD

Other Catch-up Vaccinations Needed

			Children age 4 months through 6 years		
Vaccine	Minimum Anofes		Children age 4 months un ough o years		
vaccine	Dose 1	Date 1 to Date 2	Does 2 to Does 2	Dote 2 to Doce 4	Doce 4 to Doce 5
Hepatitis B	Birth	4 weeks	8 weeks and at least 16 weeks after first dose minimum age for the final dose is 24 weeks		
Rotavirus	6 weeks Maximum age for first dose is 14 weeks, 6 days.	4 weeks	4 weeks maximum age for final dose is 8 months, 0 days		
Diphtheria, tetanus, and acellular pertussis	6 weeks	4 weeks	4 weeks	6 months	6 months
Haemophilus influenzae type b	6 weeks	No further doses needed if first dose was administered at age 15 months or older. 4 weeks if first dose was administered before the 1 th birthday. 8 weeks (as final dose) if first dose was administered at age 12 through 14 months.	No further doses needed If previous dose was administered at age 15 months or older 4 weeks If current age is younger than 12 months and first dose was administered at younger than age 7 months and at least 1 previous dose was PRP-T (AcHib ⁺ , Pentacel ⁺ , Hiberix ⁺), Vasels ⁺ or unknown 8 weeks and age 12 through 59 months (as final dose) If current age is younger than 12 months and first dose was administered at age 7 through 11 months; OR If current age is 12 through 59 months and first dose was administered before the 1 st birthday and second dose was administered at younger than 15 months; OR If both doses were PedvasHB ^a and were administered before the 1st birthday	8 weeks (as final dose) This dose only necessary for childen age 12 through 59 months who received 3 doses before the 14 birthday.	
Pheumoooccal conjugate	6 weeks	No further doses needed for healthy children if first dose was administered at age 24 months or older 4 weeks 14 first dose was administered before the 14 birthday 8 weeks (as final dose for healthy children) if hist dose was administered at the 14 birthday or after	No further doses needed for healthy children if previous dose was administered at age 24 months or older 4 weeks 4 current age is younger than 12 months and previous dose was administered at <7 months old 8 weeks (as final dose for healthy children) 8 previous close was administered between 7–11 months (wait until at least 12 months old); OR 8 current age is 12 months or older and at least 1 dose was administered before age 12 months	8 weeks (as final dose) This dose only necessary for childen age 12 through 59 months who received 3 doses before age 12 months or for childen at high risk who received 3 doses at any age.	
Inactivated policyirus	6 weeks	4 weeks	4 weeks if current age is <4 years 6 months (as final dose) if current age is 4 waars or older	6 months (minimum age 4 years for final dose)	
Measles, mumps, rubella	12 months	4 weeks			
Varicella	12 months	3 months			
Hepatitis A	12 months	6 months			
Meningococcal ACWY	2 months MenACWY-CRM 9 months MenACWY-D 2 years MenACWY-TT	8 weeks	See Note:	See Notes	
	and the second second second		Children and adolescents age 7 through 18 years		
Meningococcal ACWY	Not applicable (N/A)	8 weeks			
tetanus, diphonena; tetanus, diphtheria, and acellular pertussis	/ years	4 Webbs	4 weeks if first dose of DTaP/DT was administered before the 1 st birthday 6 months (as final dose) If first drives of DTaP/DT or Transford was administered at or after the 1 st birthday	6 months If first dose of DTaP/DT was administered before the 1 st hierbeby	
Human papillomavirus	9 years	Routine dosing intervals are			
Hepatitis A	N/A	6 months			
Hepatitis B	N/A	4 weeks	8 weeks and at least 16 weeks after first dose		
Inactivated poliovirus	N/A	4 weeks	6 months A fourth dose is not necessary if the third dose was administered at age 4 years or older and at least 6 months after the previous dose.	A fourth dose of IPV is indicated if all previous doses were administered at <4 years or if the third dose was administered <6 months after the second dose.	
Measles, mumps, rubella	N/A	4 weeks			
Varicella	N/A	3 months if younger than age 13 years. 4 weeks if age 13 years or older			
Destaura :	Builder	6 months	Exceptes		

Table 2 Recommended Catch-up Immunization Schedule for Children and Adolescents Who Start Late or Who Are More

35

Image courtesy of CDC/NCIRD

Notes Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2022

Adolescent vaccination of children who received MenACWY prior to

ongoing increased risk of meningococcal disease (e.g., those with complement deficiency, HIV, or asplenia): Follow the booster schedule

Children for whom boosters are not recommended (e.g., a healthy

meningococcal disease is endemic): Administer MenACWY according

Children for whom boosters are recommended because of an

child who received a single dose for travel to a country where

Meningococcal serogroup A,C,W,Y vaccination (minimum age: 2 months [MenACWY-CRM, Menveo], 9 months [MenACWY-D, Menactra], 2 years [MenACWY-TT, MenQuadfi])

Routine vaccination

2-dose series at age 11–12 years; 16 years

Catch-up vaccination Age 13–15 years: 1 dose now and I

(minimum interval: 8 weeks) Age 16–18 years: 1 dose

Special situations Anatomic or functional asplenia **HIV infection, persistent complet** complement inhibitor (e.g., eculi:

 Menveo - Dose 1 at age 2 months: 4-dose

and 12 months) Dose 1 at age 3-6 months: 3- or 4

after dose 1 and after age 12 mor

- Dose 1 at age 24 months or older

Persistent complement compo

Age 9-23 months: 2-dose series

Age 9-23 months: Not recomm

Age 24 months or older: 2-dose

Menactra® must be administer

- Dose 1 at age 24 months or older

Travel in countries with hyperend

disease, including countries in the

Children less than age 24 months:

Menveo[®] (age 2-23 months)

· Dose 1 at age 2 months: 4-dose

Dose 1 at age 3-6 months: 3- or

3 if applicable] at least 8 weeks a

received at age 7 months or old

least 12 weeks later and after ad

after dose 1 and after age 12 mc

· 2-dose series (dose 2 at least 12

administered as early as 8 week

Dose 1 at age 7–23 months: 2-d

Menactra® (age 9-23 months)

Children age 2 years or older: 1 do:

First-year college students who liv

previously vaccinated at age 16 ye * 1 dose Menveo[®], Menactra[®], or N

the Hajj (www.cdc.gov/travel/):

Menactra

inhibitor use:

infection:

MenQuadfi[®]

MenQuadfi

of PCV13 series.

and 12 months)

2-dose series (minimum interval: 6 months) at age 12–23 months 3 if applicable] at least 8 weeks af

Catch-up vaccination received at age 7 months or olde Unvaccinated persons through age 18 years should complete a least 12 weeks later and after age Dose 1 at age 7–23 months: 2-do 2-dose series (minimum interval: 6 months)

 Persons who previously received 1 dose at age 12 months or older should receive dose 2 at least 6 months after dose 1.

(minimum age: 12 months for routine vaccination)

 Adolescents age 18 years or older may receive the combined HepA and HepB vaccine, Twinrix®, as a 3-dose series (0, 1, and 6 months) or 4-dose series (3 doses at 0, 7, and 21-30 days, followed by a booster

dose at 12 months). Age 24 months or older: 2-dose Anatomic or functional aspleni International travel

Notes

Hepatitis A vaccination

Routine vaccination

· Persons traveling to or working in countries with high or intermediate endemic hepatitis A (www.cdc.gov/travel/):

Infants age 6-11 months: 1 dose before departure; revaccinate with 2 doses, separated by at least 6 months, between age 12-23 months.

age 10 years:

for persons at increased risk.

Unvaccinated age 12 months or older: Administer dose 1 as soon as travel is considered

Hepatitis B vaccination (minimum age: birth)

Birth dose (monovalent HepB vaccine only)

 Mother is HBsAg-negative: - All medically stable infants ≥2,000 grams: 1 dose within 24 hours of

 Infants <2,000 grams: Administer 1 dose at chronological age 1 month or hospital discharge (whichever is earlier and even if weight is still <2,000 grams).

Mother is HBsAg-positive

Administer HepB vaccine and hepatitis B immune globulin (HBIG) (in separate limbs) within 12 hours of birth, regardless of birth weight. For infants <2,000 grams, administer 3 additional doses of vaccine (total of 4 doses) beginning at age 1 month.

- Test for HBsAg and anti-HBs at age 9–12 months. If HepB series is delayed, test 1-2 months after final dose.

• Mother's HBsAg status is unknown:

- Administer HepB vaccine within 12 hours of birth, regardless of birth weight.
- For infants <2,000 grams, administer HBIG in addition to HepB vaccine (in separate limbs) within 12 hours of birth. Administer 3 additional doses of vaccine (total of 4 doses) beginning at age 1 month.
- Determine mother's HBsAg status as soon as possible. If mother is HBsAg-positive, administer HBIG to infants ≥2,000 grams as soon as possible, but no later than 7 days of age.

Routine series

- 3-dose series at age 0, 1–2, 6–18 months (use monovalent HepB) vaccine for doses administered before age 6 weeks)
- . Infants who did not receive a birth dose should begin the series as soon as feasible (see Table 2).

containing HepB is used after the birth d Minimum age for the final (3rd or 4th) dc Notes Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2022 Minimum intervals: dose 1 to dose 2:4

mellitus:

Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2022

Special situations

be administered during same visit.

months. Adolescents age 11–15 years may use ar

schedule with at least 4 months between Recombivax HB^e only) · Adolescents age 18 years or older may r (Heplisav-B*) at least 4 weeks apart.

 Adolescents age 18 years or older may re and HepB vaccine, Twinrix®, as a 3-dose 4-dose series (3 doses at 0, 7, and 21-30 dose at 12 months)

For other catch-up guidance, see Table 2

Special situations

 Revaccination is not generally recommendation normal immune status who were vaccin adolescents, or adults.

> Post-vaccination serology testing and 10mlU/mL) is recommended for certain Infants born to HBsAg-positive motified

> vaccines/hcn/acin-recs/vacc-specific/hepl

Hemodialysis patients Other immunocompromised person For detailed revaccination recommendation

Human papillomavirus vacci (minimum age: 9 years)

Routine and catch-up vaccinat HPV vaccination routinely recommende start at age 9 years) and catch-up HPV all persons through age 18 years if not a 2- or 3-dose series depending on age at

Age 9–14 years at initial vaccination months (minimum interval: 5 months; too soon)

Age 15 years or older at initial vaccir months, 6 months (minimum intervals dose 2 to dose 3: 12 weeks / dose 1 to if administered too soon)

 Interrupted schedules: If vaccination set series does not need to be restarted. No additional dose recommended wher been completed using the recommende

Special situations Immunocompromising conditions, in 3-dose series, even for those who initiate

14 years. History of sexual abuse or assault: Sta

Administration of 4 doses is permitted when a combination vaccine
 Pregnancy: Pregnancy: testing not needed before vaccination: HPV

Underlying conditions below: When both PCV13 and PPSV23 are

indicated, administer PCV13 first. PCV13 and PPSV23 should not

Chronic heart disease (particularly cyanotic congenital heart disease and cardiac failure); chronic lung disease (including

asthma treated with high-dose, oral corticosteroids); diabetes

8 weeks / dose 1 to dose 3: 16 weeks (wh

substitute "dose 4" for "dose 3" in these ca Catch-up vaccination Unvaccinated persons should complete

the Recommended Adult Immunization Schedule, 2022.

For vaccination recommendations for persons ages 19 years or older, see

Additional information

COVID-19 Vaccination COVID-19 vaccines are recommended for use within the scope of the Emergency Use Authorization or Biologics License

Application for the particular vaccine. ACIP recommendations for the use of COVID-19 vaccines can be found at www.cdc.gov vaccines/hcp/acip-recs/vacc-specific/covid-19.html CDC's interim clinical considerations for use of COVID-19

vaccines can be found at www.cdc.gov/vaccines/covid-19/clinica considerations/covid-19-vaccines-us.html.

 Consult relevant ACIP statements for detailed recommendations at www.cdc.gov/vaccines/hcp/acip-recs/index.html.

 For calculating intervals between doses, 4 weeks = 28 days. Intervals of ≥4 months are determined by calendar months.

Within a number range (e.g., 12–18), a dash (–) should be read as "through."

Vaccine doses administered ≤4 days before the minimum age or interval are considered valid. Doses of any vaccine administered ≥5 days earlier than the minimum age or minimum interval should not be counted as valid and should be repeated as age appropriate. The repeat dose should be spaced after the invalid dose by the recommended minimum interval. For further details, see Table 3-1, Recommended and minimum ages and intervals between vaccine doses, in General Best Practice Guidelines for Immunization at www.cdc.gov/vaccines/hcp/acip-recs/general-recs/timing.html.

 Information on travel vaccination requirements and recommendations is available at www.cdc.gov/travel/.

 For vaccination of persons with immunodeficiencies, see Table 8-1, Vaccination of persons with primary and secondary immunodeficiencies, in General Best Practice Guidelines for Immunization at www.cdc.gov/vaccines/hcp/acip-recs/general-recs/ mmunocompetence.html, and Immunization in Special Clinical Circumstances (In: Kimberlin DW, Brady MT, Jackson MA, Long SS, eds. Red Book: 2018 Report of the Committee on Infectious Diseases. 31st ed. Itasca, IL: American Academy of Pediatrics; 2018:67-111).

· For information about vaccination in the setting of a vaccinepreventable disease outbreak, contact your state or local health department.

The National Vaccine Injury Compensation Program (VICP) is a no-fault alternative to the traditional legal system for resolving vaccine injury claims. All routine child and adolescent vaccines are covered by VICP except for pneumococcal polysaccharide vaccine (PPSV23). For more information, see www.hrsa.gov/vaccinecompensation/index.html.

(minimum age: 9 years)

Routine vaccination

Dengue vaccination

• Age 9-16 years living in dengue endemic areas AND have laboratory confirmation of previous dengue infection - 3-dose series administered at 0, 6, and 12 months

 Endemic areas include Puerto Rico, American Samoa, US Virgin Islands. Federated States of Micronesia, Republic of Marshall Islands, and the Republic of Palau. For updated guidance on dengue endemic areas and pre-vaccination laboratory testing see www.cdc.gov/mmwr/ volumes/70/rr/rr7006a1.htm?s_cid=rr7006a1_w and www.cdc.gov/ dengue/vaccine/hcp/index.html

Diphtheria, tetanus, and pertussis (DTaP) vaccination (minimum age: 6 weeks [4 years for Kinrix[®] or Quadracel[®]])

Routine vaccination

• 5-dose series at age 2, 4, 6, 15-18 months, 4-6 years - Prospectively: Dose 4 may be administered as early as age 12 months if at least 6 months have elapsed since dose 3.

 Retrospectively: A 4th dose that was inadvertently administered as early as age 12 months may be counted if at least 4 months have

elapsed since dose 3. Catch-up vaccination

 Dose 5 is not necessary if dose 4 was administered at age 4 years or older and at least 6 months after dose 3. For other catch-up guidance, see Table 2.

Special situations

· Wound management in children less than age 7 years with history of 3 or more doses of tetanus-toxoid-containing vaccine: For all wounds except clean and minor wounds, administer DTaP if more than 5 years since last dose of tetanus-toxoid-containing vaccine. For detailed information, see www.cdc.gov/mmwr/volumes/67/rr/rr6702a1.htm.

Haemophilus influenzae type b vaccination (minimum age: 6 weeks)

Routine vaccination

 ActHIB[®], Hiberix[®], Pentacel[®], or Vaxelis[®]: 4-dose series (3 dose primary series at age 2, 4, and 6 months, followed by a booster dose* at age 12-15 months)

- *Vaxelis* is not recommended for use as a booster dose. A different Hib-containing vaccine should be used for the booster dose.
- PedvaxHIB[®]: 3-dose series (2-dose primary series at age 2 and 4 months, followed by a booster dose at age 12-15 months)

Catch-up vaccination

• Dose 1 at age 7-11 months: Administer dose 2 at least 4 weeks later and dose 3 (final dose) at age 12-15 months or 8 weeks after dose 2 (whichever is later)

• Dose 1 at age 12-14 months: Administer dose 2 (final dose) at least 8 weeks after dose 1

Dose 1 before age 12 months and dose 2 before age 15 months: Administer dose 3 (final dose) at least 8 weeks after dose 2. * 2 doses of PedvaxHIB* before age 12 months: Administer dose 3 (final dose) at 12-59 months and at least 8 weeks after dose 2.

36

* 1 dose administered at age 15 months or older: No further doses needed

• Unvaccinated at age 15-59 months: Administer 1 dose. Previously unvaccinated children age 60 months or older who are not considered high risk: Do not require catch-up vaccination

For other catch-up guidance, see Table 2. Vaxelise can be used for catchup vaccination in children less than age 5 years. Follow the catch-up schedule even if Vaxelis® is used for one or more doses. For detailed information on use of Vaxelis* see www.cdc.gov/mmwr/volumes/69/ wr/mm6905a5.htm

Special situations

Chemotherapy or radiation treatment: Age 12-59 months

- Unvaccinated or only 1 dose before age 12 months: 2 doses, 8 weeks apart

- 2 or more doses before age 12 months: 1 dose at least 8 weeks after previous dose

Doses administered within 14 days of starting therapy or during therapy should be repeated at least 3 months after therapy completion.

Hematopoietic stem cell transplant (HSCT);

- 3-dose series 4 weeks apart starting 6 to 12 months after successful transplant, regardless of Hib vaccination history

* Anatomic or functional asplenia (including sickle cell disease): Age 12-59 months

- Unvaccinated or only 1 dose before age 12 months: 2 doses, 8 weeks apart

- 2 or more doses before age 12 months: 1 dose at least 8 weeks after previous dose

Unvaccinated* persons age 5 years or older

Unvaccinated* persons age 5-18 years

no doses (age 15 months or older)

Unvaccinated* persons age 15 months or older

- 1 dose (preferably at least 14 days before procedure)

Unvaccinated or only 1 dose before age 12 months: 2 doses,

Immunoglobulin deficiency, early component complement

Unvaccinated or only 1 dose before age 12 months: 2 doses,

- 2 or more doses before age 12 months: 1 dose at least 8 weeks after

Image courtesy of CDC/NCIRD

*Unvaccinated = Less than routine series (through age 14 months) OR

2 or more doses before age 12 months: 1 dose at least 8 weeks after

Elective splenectomy:

HIV infection:

Age 12–59 months

8 weeks apart

previous dose

- 1 dose

deficiency:

Age 12–59 months

8 weeks apart

previous dose

- 1 dose
Other Catch-up Vaccinations Needed

Appendix	Recommended Child and Adolescent Immunization Sched	lule for ages 18 years or younger, United States, 2022		
Vaccine	Contraindications ¹	Precautions ²		
Dengue (DEN4CYD)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁴ Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of chemotherapy, congenital immunodeficiency, long- term immunosuppressive therapy or patients with HN infection who are severely immunocompromised) 	Pregnancy HtVInfection without evidence of severe immunosuppression Moderate or severe acute illness with or without fever		
Diphtheria, tetanus, pertussis (DTaP) Tetanus, diphtheria (DT)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component⁸ For DTaP only: Encephalopathy (e.g., coma, decreased level of consciousness, prolonged seizures) not attributable to another identifiable cause within 7 days of administration of previous dose of DTP or DTaP 	Guillain-Barré syndrome (GBS) within 6 weeks after previous dose of tetanus-toxoid-containing vaccine History of Arthus-type hypersensitivity reactions after a previous dose of diphtheria-toxoid—containing or tetanus toxoid-containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus toxoid-containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus toxoid-containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus toxoid-containing vaccine; defer location in the status classified and stabilized For DTaP only: Progressive neurologic disorder; including infantile spasms, uncontrolled epilepsy, progressive encephalopathy; defer DTaP until neurologic status clarified and stabilized Moderate or severe acute illness with or without fever		
Haemophilus influenzae type b (Hib)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component* For Hiberix, ActHib, and PedvaxHiB only: History of severe allergic reaction to dry natural latex Less than are 6 weeks			
Hepatitis A (HepA)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ including neomycin 	Moderate or severe acute illness with or without fever		
Hepatitis B (HepB)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ¹ including yeast Por Heplisav-B only: Pregnancy			
Hepatitis A- Hepatitis B vaccine [HepA-HepB, (Twinrix*)]	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ including neomycin and yeast 	Moderate or severe acute illness with or without fever		
Human papillomavirus (HPV)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ 	Moderate or severe acute illness with or without fever		
Measles, mumps, rubella (MMR)	 Server allergic reaction (e.g. anaphylaxid) after a previous dose or to a vaccine component⁴. Server immunodeficiency (e.g. hematologic and solid tumors, receipt of chemotherapy, congenital immunodeficiency, long-term immunosuppressive therapy or patients with HIV infection who are severely immunocomponised) Pregnancy Family history of altered immunocompetence, unless verified clinically or by laboratory testing as immunocompetent 	Recent (s11 months) receipt of antibody-containing blood product (specific interval depends on product) History of thrombocytopenia or thrombocytopenic purpura Need for tuberculin skin testing or interferon-gamma release assay (IGRA) testing Moderate or severe acute illness with or without fever		
Meningococcal ACWY (MenACWY) [MenACWY-CRM (Menveo"); MenACWY-D (Menactra"); MenACWY-TT (MenQuadfi")]	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ For MenACW/-D and Men ACWY-CRM only: severe allergic reaction to any diphtheria taxoid- or CRM197 containing vaccine For MenACW/-TT only: severe allergic reaction to a tetanus toxoid-containing vaccine 	For MenACWI-CRM only: Preterm birth if less than age 9 months Moderate or severe acute illness with or without fever		
Meningococcal B (MenB) (MenB-4C (Bexsero*); MenB-FHbp (Trumenba*)]	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ^a	Pregnancy For MenB-4C only: Latex sensitivity Moderate or severe acute illness with or without fever		
Pneumococcal conjugate (PCV13)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ Severe allergic reaction (e.g., anaphylaxis) to any diphtheria-toxoid- containing vaccine or its component¹ 	Moderate or severe acute illness with or without fever		
Pneumococcal polysaccharide (PPSV23)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ¹	Moderate or severe acute illness with or without fever		
Poliovirus vaccine, inactivated (IPV)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ 	Pregnancy Moderate or severe acute illness with or without fever		
Rotavirus (RV) [RV1 (Rotarix*), RV5 (RotaTeq*)]	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component^a Severe combined immunodeficiency (SCID) History of intussusception 	Altered immunocompetence other than SCID Chronic gastrointestinal disease RV1 only: Spina bifida or bladder exstrophy Moderate or severe acute illness with or without fever		
Tetanus, diphtheria, and acellular pertussis (Tdap) Tetanus, diphtheria (Td)	 Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component¹ For Tdap only: Encephalopathy (e.g., coma, decreased level of consciousness, prolonged seizures) not attributable to another identifiable cause within 7 days of administration of previous dose of DTP; DTaP; or Tdap 	Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of tetanus-toxoidcontaining vaccine History of Arthus-type hypersensitivity reactions after a previous dose of diphtheria-toxoid— containing or tetanus-toxoid containing vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid containing vaccine; For Tdap only: Progressive or unstable neurological disorder, uncontrolled seizures, or progressive encephalopathy until a treatment regimen has been established and the condition has stabilized Moderate or severe acute illness with or without fever		
Varicella (WAR)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ¹ Severe immunodeficiency (e.g., hematologic and solid turnors, receipt of chemotherapy, congenital immunodeficiency, (ang.) term immunosuppressive therapy or patients with HIV infection who are severely immunocompromised) Pregnancy Family history of altered immunocompetence, unless verified clinically or by laboratory testing as immunocompetent	accine component ⁴ * Recent (s11 months) receipt of antibody-containing blood product (specific interval depends on product) * Receipt of specific antiviral drugs (acyclovir, famciclovir, or valacyclovir) 24 hours before vaccination (avoid use of these antiviral drugs for 14 days after vaccination) Use of aspirin or aspirin-containing products * Moderate or severe acute illness with or without fever		

When a contraindication is present, a vaccine should NOT be administered. Kroger A, Bahta L, Hunter P. ACIP General Best Practice Guidelines for Immunization. www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html

 When a precaution is present, vaccination should generally be deferred but might be indicated if the benefit of protection 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

 Vaccination providers should check FDA² 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.

Case study

	Vaccines Needed
Catch-up routine vaccinations needed	 HepB: dose 3 IPV- dose 3 Tdap or Td -dose 4 MMR- dose 1 and 2 VAR- dose 1 and 2 HepA- dose 1 and 2 HPV- dose 1 and 2 MenACWY- dose 1 and 2
Other vaccinations	 COVID-19 vaccine (Pfizer; dose 1, 2, and booster)

Medical and Vaccination History

- 14-year-old female (Date of birth-03/03/2008)
- Medical history: none HIV diagnosed 2020
- Immunization record shows only:
 - DTaP on 6/29/2008 and 4/20/2013
 - Tdap on 7/20/2019
 - IPV on 6/29/2008 and 4/20/2013
 - Hepatitis B vaccine on 3/03/2008 and 6/29/2008

Parents report she received other vaccines and is up-to-date but have no records



Recommended Child and Adolescent Immunization Schedule by Medical Indication, United States, 2021

Always use this table in conjunction with Table 1 and the notes that follow.

					IN	DICATION				
VACCINE	Pregnancy	Immunocom- promised status (excluding HIV infection)	HIV infection <15% and total CD4 cell count of <200/mm ³	CD4+ count ¹ ≥15% and total CD4 cell count of ≥200/mm ³	Kidney failure, end-stage renal disease, or on hemodialysis	Heart disease or chronic lung disease	CSF leak or cochlear implant	Asplenia or persistent complement component deficiencies	Chronic liver disease	Diabetes
Hepatitis B										
Rotavirus		SCID ²								
Diphtheria, tetanus, and acellular pertussis (DTaP)										
Haemophilus influenzae type b										
Pneumococcal conjugate										
Inactivated poliovirus										
Influenza (IIV)	_									
Influenza (LAIV4)						Asthma, wheezing: 2–4yrs ²				
Measles, mumps, rubella	*									
Varicella	*									
Hepatitis A										
Tetanus, diphtheria, and acellular pertussis (Tdap)										
Human papillomavirus	*									
Meningococcal ACWY										
Meningococcal B										
Pneumococcal polysaccharide										
Vaccination according t routine schedule recommended	to the	Recommended for persons with an additio risk factor for which the vaccine would be indic	onal and e nec ated con	additional doses essary based on n dition. See Notes.	hended, Not maybe com nedical sho	recommended/ traindicated—vaccine uld not be administered. ccinate after pregnancy.	Precaution might be indi of protection of adverse rea	raccine cated if benefit outweighs risk action	commendat cable	ion/not

medical condition (HIV) Hib • PCV13 • PPSV23 • Vaccine that MIGHT be contraindicated due to medical condition (HIV) MMR • Varicella • Vaccine that requires additional dose due to medical condition (HIV)

Vaccine indicated due to

- HPV
- MenACWY

 For additional information regarding HIV laboratory parameters and use of live vaccines, see the General Best Practice Guidelines for Immunization, "Altered Immunocompetence," at www.cdc.gov/vaccines/hcp/acip-recs/general-recs/immunocompetence.html and Table 4-1 (footnote D) at www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html.
 Severe Combined Immunodeficiency

3 LAIV4 contraindicated for children 2-4 years of age with asthma or wheezing during the preceding 12 months

Case study

	Vaccines Needed
Catch-up routine vaccinations needed	 HepB: dose 3 IPV- dose 3 Tdap or -dose 4 MIMR- dose 1 and 2 (only if CD4≥15% <u>and</u> total CD4 cell count of ≥200/mm) VAR- dose 1 and 2 (only if CD4≥15% <u>and</u> total CD4 cell count of ≥200/mm) HepA- dose 1 and 2 HPV- dose 1, 2, and 3 MenACWY- dose 1 and 2; and booster doses Hib-dose 1 (final) PCV13- dose 1 (final) PPSV23- dose 1 and 2
Other vaccinations	 COVID-19 vaccine (Pfizer; dose 1, 2, additional dose and booster)

Co-administration of Vaccines

Co-administration of Vaccines

 Co-administration (simultaneous administration) of vaccines is defined as administering more than one vaccine on the same clinic day, at different anatomic sites, and not combined in the same syringe

TABLE 3-4. Guidelines for spacing of live and non-live antigens			
Antigen combination	Recommended minimum interval between doses		
Two or more non-live (a),(b)	May be administered simultaneously or at any interval between doses		
Non-live and live (c)	May be administered simultaneously or at any interval between doses		
Two or more live injectable ^(e)	28 days minimum interval, if not administered simultaneously		
Source: (83).			
^(a) Certain experts suggest a 28-day interva (Tdap) vaccine and tetravalent meningoco	al between tetanus toxoid, reduced diphtheria toxoid, and acellular pertussis occal conjugate vaccine if they are not administered simultaneously.		
^(b) In persons with functional or anatomic should be spaced by 4 weeks. Likewise for and PPSV23, PCV13 should be administer For persons 65 years old or older indicate should be administered 6-12 months later	asplenia, MCV-D and PCV13 should not be administered simultaneously and persons with immunosuppressive high-risk conditions indicated for PCV13 ed first, and PPSV23 should be administered no earlier than 8 weeks later. d for PCV13 and PPSV23, PCV13 should be administered first and PPSV23		

^(c) The live oral vaccines Ty21a typhoid vaccine and rotavirus vaccine may be administered simultaneously with or at any interval before or after non-live or live injectable vaccines.

Kroger A, Bahta L, Hunter P. General Best Practice Guidelines for Immunization. Best Practices Guidance of the Advisory Committee on Immunization Practices (ACIP). https://www.cdc.gov/vaccines/hcp/acip-recs/general-recs/timing.html

Coadministration of COVID-19 vaccines with other vaccines

COVID-19 vaccines and other vaccines may be administered without regard to timing.



Co-administration of COVID-19 with other vaccines. <u>https://www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html?CDC_AA_refVal=https%3A%2F%2Fwww.cdc.gov%2Fvaccines%2Fcovid-19%2Finfo-by-product%2Fclinical-considerations.html#Coadministration</u>

Vaccination resources for healthcare providers

CDC vaccination resources for healthcare providers

- Schedules App
 - <u>https://www.cdc.gov/vaccines/schedules/hcp/schedule-app.html</u>
- Child and Adolescent Vaccine Assessment Tool
 - https://www2.cdc.gov/vaccines/childquiz

- Storage and Handling Toolkit
 - <u>https://www.cdc.gov/vaccines/hcp/admin/storage/</u> toolkit/index.html



Key Takeaways

- CDC's child and adolescent immunization schedule is updated and published annually
- The immunization schedule is a useful tool for healthcare providers to determine which vaccines are indicated for each child/adolescent
- Reviewing all sections of the schedule is important when determining which vaccines are needed for each child/adolescent
- Children/adolescents with certain medical conditions need additional vaccines not typically recommended for age, or additional doses of a recommended vaccine
- COVID-19 vaccines can be co-administered with other vaccines on the same day or at any interval from other vaccines



Centers for Disease Control and Prevention. (2021). ACIP Vaccine Recommendations and Guidelines. https://www.cdc.gov/vaccines/hcp/acip-recs/index.html

Centers for Disease Control and Prevention. (2021). Child and adolescent immunization schedule.

https://www.cdc.gov/vaccines/schedules/hcp/imz/child-adolescent.html

Centers for Disease Control and Prevention. (2021). Interim Clinical Considerations for Use of COVID-19 Vaccines.

https://www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html

Kroger, A., Bahta, L., Hunter, P. General Best Practice Guidelines for Immunization. Best Practices Guidance of the Advisory Committee on Immunization

Practices (ACIP). (2021). <u>https://www.cdc.gov/vaccines/hcp/acip-recs/general-recs/timing.html</u>

Wodi, A.P., Ault, K., Hunter, P., McNally, V., Szilagyi, P.G., Bernstein, H. (2021). Advisory Committee on Immunization Practices Recommended Immunization

Schedule for Children and Adolescents Aged 18 Years or Younger — United States, 2021. Morbidity and Mortality Weekly Report, 70:189–192.

doi: http://dx.doi.org/10.15585/mmwr.mm7006a1external icon

For more information

For more information, contact CDC 1-800-CDC-INFO (232-4636) TTY: 1-888-232-6348 www.cdc.gov

https://www.cdc.gov/cdc-info/index.html

The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the Centers for Disease Control and Prevention.

Photographs and images included in this presentation are licensed solely for CDC/NCIRD online and presentation use. No rights are implied or extended for use in printing or any use by other CDC CIOs or any external audiences.



Pediatric COVID-19 Vaccination

Elisha Hall, PhD, RD Clinical Guidelines Lead Vaccine Task Force

Clinical Communities Speaker Series April 28, 2022





cdc.gov/coronavirus

Learning Objectives

At the conclusion of this session participants will be able to:

- Describe COVID-19 vaccination recommendations for children ages 5– 11 years and adolescents ages 12–17 years of age.
- 2. Review COVID-19 vaccine formulations for the pediatric population.

Overview of COVID-19 Vaccination

- Recommended for everyone ages 5 years and older in the United States for the prevention of COVID-19
- Effective in preventing serious outcomes of COVID-19
- Safe; known benefits continue to outweigh possible risks
- Critical to get people up to date on COVID-19 vaccination

COVID-19 Vaccination Coverage¹

Children ages 5–11 years



At least 1 dose

27.8% Fully vaccinated

Adolescents ages 12–17 years

68.5%

At least 1 dose

58.5% Fully vaccinated

COVID-19 Vaccination Coverage¹

Children ages 5–11 years



At least 1 dose

27.8% Fully vaccinated

Adolescents ages 12–17 years

68.5%

At least 1 dose

58.5%

Fully vaccinated

¹Data as of March 31, 2022, <u>https://covid.cdc.gov/covid-data-tracker/#vaccinations-cases-trends</u>

Pediatric COVID-19 Vaccination Recommendations





Degree of immune suppression

Age group

Pediatric COVID-19 Vaccination Recommendations

- Recommendations vary with degree of immune suppression.
- People with immunocompromising conditions or people who take immunosuppressive medications or therapies:
 - Are at increased risk for severe COVID-19
 - May be less likely to mount a protective immune response after initial vaccination
 - Have waning protection over time

People Who Are Moderately or Severely Immunocompromised

- Active treatment for solid tumor and hematologic malignancies
- Receipt of solid-organ transplant and taking immunosuppressive therapy
- Receipt of CAR-T-cell or hematopoietic stem cell transplant (HCT) (within 2 years of transplantation or taking immunosuppression therapy)
- Moderate or severe primary immunodeficiency (e.g., DiGeorge, Wiskott-Aldrich syndromes)
- Advanced or untreated HIV infection
- Active treatment with high-dose corticosteroids (i.e., ≥20mg prednisone or equivalent per day), alkylating agents, antimetabolites, transplant-related immunosuppressive drugs, cancer chemotherapeutic agents classified as severely immunosuppressive, TNF blockers, and other biologic agents that are immunosuppressive or immunomodulatory

https://www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html#vaccinationpeople-immunocompromised

People Who Are Moderately or Severely Immunocompromised

- People can self-attest.
- Patients do NOT need to provide documentation.
- Vaccinators should NOT deny COVID-19 vaccination to a person due to lack of documentation.
- People who are moderately or severely immunocompromised can discuss what is appropriate for them with their healthcare provider.
- CDC offers a prevaccination checklist, where people can check if they are considered moderately or severely immunocompromised.

Prevaccination Checklis for COVID-19 Vaccinatio	st on
For vaccine recipients: The following questions will help us determine if there is any reason you shin to get the COVID-19 vaccine today. If you answer "yes" to any question, t does not necessarily mean you should not be vaccinated. It just means additional questions may be asked. If a question is not clear, please ask your validation of the totage of the totaget of	Name
	Yes No know
Are you feeling sick today?	
Have you ever received a dose of COVID-19 vaccine? If yes, which vaccine product(s) did you receive? Pfizer-BioNTech Moderna	Another Product Johnson)
How many doses of COVID-19 vaccine have you received?	
Did you bring your vaccination record card or other documenta	tion?
 Do you have a health condition or are you undergoing treatment or severely immunocompromised? (This would include treatment for cance immunosuppressive therapy or high-dose corticasteroids, CAR-T-cell therapy, hematopo or Wisket-Main's syndrome). 	that makes you moderately
 Have you received hematopoietic cell transplant (HCT) or CAR-T-c COVID-19 vaccine? 	ell therapies since receiving
 Have you ever had an allergic reaction to: (This would include a severe allergic reaction (eg., anaphylaxis) that required treatment to ao to the hospital. It would also include an alleraic reaction that caused hives. swelline A component of a COVID-19 vaccine, including either of the following: o Polyethylene glycol (<i>PEG</i>), which is found in some medications, such colonoscopy procedures 	with epinephrine or EpiPen* or that caused you 2. or resoliratory distress. includina wheezina.) as laxatives and preparations for
 Polysorbate, which is found in some vaccines, film coated tablets, and 	d intravenous steroids
A previous dose of COVID-19 vaccine	
 Have you ever had an allergic reaction to another vaccine (other th or an injectable medication? (This would include a severe allergic reaction (leg., anaphylaxii) that required treatment to go to the hospital. It would also include an allergic reaction that caused hives, swelling 	han COVID-19 vaccine) with epinephrine or EpiPen* or that caused you a or respiratory distress, including wheezing.)
Check all that apply to you:	
Am a female between ages 18 and 49 years old	Have a bleeding disorder
Am a male between ages 12 and 29 years old	Take a blood thinner
Have a history of myocarditis or pericarditis	Have a history of heparin-induced thrombocytopenia (HIT)
Have been treated with monoclonal antibodies or convalescent serum to prevent or treat COVID-19	Am currently pregnant or breastfeeding
Diagnosed with Multisystem Inflammatory Syndrome (MIS-C or	Have received dermal fillers Have a history of Guillain-Barré Syndrome (GRS)

https://www.cdc.gov/vaccines/covid-19/downloads/pre-vaccination-screening-form.pdf

Pediatric COVID-19 Vaccination Recommendations



Ages 4 years and younger



Ages 5–11 years



Ages 12–17 years

Recommendations for Children Ages 4 Years and Younger

- Currently no FDA-approved or authorized COVID-19 vaccine for children ages 4 years and younger
- Should not receive any COVID-19 vaccine doses at this time



Recommendations for Children Ages 5–11 Years

- Children in this age group
 - Should receive a total of 2
 or 3 doses, based on their
 degree of immune
 suppression
 - Should only receive Pfizer-BioNTech COVID-19 Vaccine



Recommendations for Children Ages 5–11 Years

Children who are NOT moderately or severely immunocompromised should receive a total of 2 doses.



Recommendations for Children Ages 5–11 Years Who Are Moderately or Severely Immunocompromised

Children who ARE moderately or severely immunocompromised should receive a total of 3 doses.



Recommendations for Adolescents Ages 12–17 Years

- Adolescents in this age group
 - Should receive a total of 3
 or 4 doses based on their
 degree of immune
 suppression
 - Should only receive Pfizer-BioNTech COVID-19 Vaccine



Recommendations for Adolescents Ages 12–17 Years

 Adolescents who are NOT moderately or severely immunocompromised should receive a total of 3 doses.



Primary Series Interval of 3-8 Weeks



Primary Series Interval

• High COVID-19

disease

community levels



- Reduced myocarditis risk
- **Optimize vaccine** ulleteffectiveness

Recommendations for Adolescents Ages 12–17 Years Who are Moderately or Severely Immunocompromised

68

- Adolescents who are moderately or severely immunocompromised should receive a total of 4 doses.
- They may receive a second booster dose for a total of 5 doses.



Second Booster Dose

- New recommendation
- Only three groups of people MAY receive a second booster

People 12 years and older who are moderately or severely immunocompromised

People 50 years and older

People 18 years and older who received both a primary and booster dose of Janssen COVID-19 vaccine

Summary of Recommendations by Age and Immune Status



Knowledge Check

How should vaccinators verify that a patient is moderately or severely immunocompromised?

Knowledge Check

A 15-year-old patient with severe immune compromise has the following vaccination history:

Dose 1 (primary): 6/01/21 Dose 2 (primary): 6/22/21 Dose 3 (primary): 8/14/21 Dose 4 (booster): 1/14/22 **True/false**: They **may** get a 2nd booster today?


Staying Up to Date

- CDC recommends everyone get up to date with their COVID-19 vaccinations.
- Being up to date means a person has received all recommended doses in their primary vaccine series, and a booster dose, when eligible. Receipt of a second booster dose is not necessary to be considered up to date at this time.



Age indications	5 through 11 years	12 years and older	12 years and older
Doses per vial	10	6	6
Dilution required	Yes—1.3 mL	Yes—1.8 mL	No
Dose	10 mcg	30 mcg	30 mcg
Dose volume	0.2 mL	0.3 mL	0.3 mL

	Orange cap	Purple cap	Gray cap
Age indications	5 through 11 years	12 years and older	12 years and older
Doses per vial	10	6	6
Dilution required	Yes—1.3 mL	Yes—1.8 mL	No
Dose	10 mcg	30 mcg	30 mcg
Dose volume	0.2 mL	0.3 mL	0.3 mL

	Orange cap	Purple cap	Gray cap
Age indications	5 through 11 years	12 years and older	12 years and older
Doses per vial	10	6	6
Dilution required	Yes—1.3 mL	Yes—1.8 mL	No
Dose	10 mcg	30 mcg	30 mcg
Dose volume	0.2 mL	0.3 mL	0.3 mL

	Orange cap	Purple cap	Gray cap
Age indications	5 through 11 years	12 years and older	12 years and older
Doses per vial	10	6	6
Dilution required	Yes—1.3 mL	Yes—1.8 mL	No
Dose	10 mcg	30 mcg	30 mcg
Dose volume	0.2 mL	0.3 mL	0.3 mL

Children should receive the ageappropriate vaccine formulation and follow the schedule based on their age on the day of vaccination, regardless of their size or weight.

Dosing and Formulation

- Children ages 5–11 years should receive the 10 µg Pfizer-BioNTech COVID-19 Vaccine (orange cap vial) formulation.
- Adolescents ages 12 years and older should receive the 30 μg Pfizer-BioNTech COVID-19 Vaccine (purple or gray cap vial) formulation.
- If a child turns 12 years old between their first and second dose, they should receive the age-appropriate 30 µg Pfizer-BioNTech COVID-19 Vaccine (purple or gray cap vial) formulation for their second dose.

Vaccine Dosage and Schedule

- Vaccine dosage is based on age and not size or weight
- Work differently than other medications
- Clinical trials determine the best dosage and schedule

Common Errors for COVID-19 Pediatric Vaccination

Error	Recommended action
Unauthorized age group (recipients ages 4 years and younger)	Do not give another dose at this time.
Unauthorized age group (recipients ages 5–11 years)	If Moderna vaccine administered : Do not repeat the dose (Moderna dose "counts"). Give Pfizer-BioNTech for the next dose in the schedule, if applicable.
If ages 5–11 years and Pfizer-BioNTech purple or gray cap inadvertently administered	Do not repeat dose.
If ages 12–17 years and Pfizer-BioNTech orange cap inadvertently administered	Do not repeat dose. However, based on clinical judgement, a repeat dose of Pfizer-BioNTech vaccine ≥12 years formulation (30 µg, purple or gray cap) may be administered at an interval 3-8 weeks after the dose given in error.

https://www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html#appendix-c

Coadministration

- COVID-19 vaccines may be administered without regard to timing of other vaccines.
- Providers are encouraged to offer all vaccines at the same visit.
- Best practices for multiple injections include:
 - Label each syringe
 - Separate injection sites by 1 inch or more, if possible.
 - Administer the COVID-19 vaccine and vaccines that may be more likely to cause a local reaction in different limbs, if possible.

Knowledge Check

A 12-year-old patient (DOB 03/01/2010) received 2 primary doses at age 11 years. Dose 1: 11/03/2021 (Pfizer orange cap) Dose 2: 11/24/2021 (Moderna, error) She is due for her booster dose 4/24/2021. What should she receive?

https://www.gettyimages.com/

Key Takeaways

- COVID-19 vaccines are safe and effective.
- COVID-19 vaccination recommended for everyone ages 5 years and older in the United States for the prevention of COVID-19.
 - Between 2 to 5 doses depending on the age and immune status
- Children should receive the formulation based on their age on the day of vaccination.
- Children and adolescents should get up to date as soon as possible.

Resources

- Interim Clinical Considerations for Use of COVID-19 Vaccines Currently Approved or Authorized in the United States <u>https://www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html</u>
- U.S. COVID-19 Vaccine Product Information <u>https://www.cdc.gov/vaccines/covid-19/info-by-product/index.html</u>
- Stay Up to Date with Your COVID-19 Vaccines <u>https://www.cdc.gov/coronavirus/2019-ncov/vaccines/stay-up-to-date.html</u>
- Comfort and Restraint Techniques <u>https://www.youtube.com/watch?v=r1dGpTCgerE</u>
- Before, During, and After Shots: <u>www.cdc.gov/vaccines/parents/visit/before-during-after-shots.html</u>
- Fainting (Syncope) after Vaccination: <u>www.cdc.gov/vaccinesafety/concerns/fainting.html</u>
- Vaccine Administration: Needle Gauge and Length: www.cdc.gov/vaccines/hcp/admin/downloads/vaccine-administration-needle-length.pdf
- Vaccine Administration: Intramuscular (IM) Injection Children 3 through 6 years of age: <u>https://www.cdc.gov/vaccines/hcp/admin/downloads/IM-Injection-3-6-Years.pdf</u>
- Vaccine Administration: Intramuscular (IM) Injection Children 7 through 18 years of age: <u>www.cdc.gov/vaccines/hcp/admin/downloads/IM-Injection-children.pdf</u>
- Vaccinate with Confidence communication materials
 - Routinely recommended vaccines: <u>www.cdc.gov/vaccines/partners/vaccinate-with-confidence.html</u>
 - COVID-19 vaccine: <u>www.cdc.gov/vaccines/covid-19/vaccinate-with-confidence.html</u>
- Epidemiology and Prevention of Vaccine-Preventable Diseases, Vaccine Administration Chapter: <u>www.cdc.gov/vaccines/pubs/pinkbook/vac-admin.html</u>
- You Call the Shots Vaccine Administration: <u>www2.cdc.gov/vaccines/ed/vaxadmin/va/ce.asp</u>
- General Best Practice Guidelines for Immunization: <u>www.cdc.gov/vaccines/hcp/acip-recs/general-recs/index.html</u>



Centers for Disease Control and Prevention. (2021). COVID-19 vaccine. www.cdc.gov/vaccines/covid-19/vaccinate-with-confidence.html

Centers for Disease Control and Prevention. (2021). Epidemiology and Prevention of Vaccine-Preventable Diseases, Vaccine Administration Chapter.

www.cdc.gov/vaccines/pubs/pinkbook/vac-admin.html

Centers for Disease Control and Prevention. (2021). Routinely recommended vaccines.

https://www.cdc.gov/vaccines/partners/vaccinate-with-confidence.html

Centers for Disease Control and Prevention. (2021). U.S. COVID-19 Vaccine Product Information.

https://www.cdc.gov/vaccines/covid-19/info-by-product/index.html

Centers for Disease Control and Prevention. (2021). Vaccine Administration: Intramuscular (IM) Injection Children 3 through 6 years of age.

https://www.cdc.gov/vaccines/hcp/admin/downloads/IM-Injection-3-6-Years.pdf

Centers for Disease Control and Prevention. (2021). Vaccine Administration: Intramuscular (IM) Injection Children 7 through 18 years of age.

www.cdc.gov/vaccines/hcp/admin/downloads/IM-Injection-children.pdf



For more information, contact CDC 1-800-CDC-INFO (232-4636) TTY: 1-888-232-6348 www.cdc.gov



The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the Centers for Disease Control and Prevention.



DHA Childhood Catchup Immunization Efforts

Navy CAPT Gregory H. Gorman, M.D., M.H.S. Executive Director, Defense Health Board Chair, DHA Complex Pediatrics Clinical Community Falls Church, Va.





Defense Health Agency Memo on Childhood Catch-Up Immunizations



DEFENSE HEALTH AGENCY 7700 ARLINGTON BOULEVARD, SUITE 5101 FALLS CHURCH, VIRGINIA 22042-5101

30 JUL 2021

MEMORANDUM FOR DIRECTOR, COASTAL MISSISSIPPI MARKET DIRECTOR, JACKSONVILLE MARKET DIRECTOR, NATIONAL CAPITAL REGION MARKET DIRECTOR, CENTRAL NORTH CAROLINA MARKET DIRECTOR, TIDEWATER MARKET DIRECTOR, COLORADO MARKET DIRECTOR, SAN ANTONIO MARKET DIRECTOR, PUGET SOUND MARKET DIRECTOR, SAN DIEGO MARKET DIRECTOR, HAWAII MARKET DIRECTOR, AUGUSTA MARKET DIRECTOR, CENTRAL TEXAS MARKET DIRECTOR. COASTAL NORTH CAROLINA MARKET DIRECTOR, LOW COUNTRY MARKET DIRECTOR, SACRAMENTO MARKET DIRECTOR, SOUTHWEST GEORGIA MARKET DIRECTOR, SOUTHWEST KENTUCKY MARKET DIRECTOR, SMALL MARKET AND STAND-ALONE MILITARY MEDICAL TREATMENT FACILITY ORGANIZATION LEAD, DIRECT SUPPORT ORGANIZATION, ARMY LEAD, DIRECT SUPPORT ORGANIZATION, NAVY LEAD, DIRECT SUPPORT ORGANIZATION, AIR FORCE

SUBJECT: Interim Guidance for Childhood Immunization Catch-Up

This memorandum establishes interim guidance for improving childhood immunization rates. Military Health System (MHS) childhood immunization rates have fallen from 85-95% to 70-80% in the setting of the COVID-19 pandemic. For measles, the population vaccine rate rate 500% - 1

Champions for each Market

Expanded use of Immunization Registry to identify and contact children for catch-up immunizations

Scripts incorporating best practices to communicate with parents and guardians

Training for population health managers and champions

Suggested innovative means to deliver immunizations

Central tracking and reporting by Market





MMR # 1 Immunization Rates of 16-18 month old Children Enrolled in Military Treatment Facilities, Aug 2021 – April 2022



The enrolled population ranged from ~5800-6900 during the time period.



Medically Ready Force... Ready Medical Force





Best Practices to Catch-up Child Immunizations:

Examples from the 2021 Winners of the Health Resources & Services Administration's Promoting Pediatric Primary Prevention (P4) Challenge

Text & Chat-bots (Chicago, IL; Alexandria, VA)	Partnerships w schools (Urbana, IL; Jefferson Cor	vith unty, AL)	Partnerships with Dentistry (Denver, CO; Vista, CA)		Partnerships with local departments of health (Manassas, VA)
Registries to betterTargetidentify patientsp(Bronx, NY; Denver, CO; Chicago, IL)(Virgit)		Targetir popu (Virginia; Ba	ng at-risk lations altimore, MD)	Mob (Manassas, Peoria, IL; Logan	ile clinics VA; Columbus, OH; Bremerton, WA; County, WV)
Resources to overcome barriers (Baltimore, MD; Nashville, TN; Alexandria, VA)	Off-hours Clir (Vista, CA)	nics	Partnersh churc (Memphi	ips with hes ^{s, TN)}	Offering incentives (San Fernando Valley, CA)

Full list and complete description of funded programs at

https://mchb.hrsa.gov/funding/challenge-competitions/p4challenge/our-20-final-winners/winners-showcase







Department of Defense Memorandum. (2021). Interim Guidance for Childhood Immunization Catch-up.

Department of Defense Memorandum, Deputy Assistant of Medical Affairs, Defense Health Agency.

Health Resources and Services Administration.(2022). P4 challenge winners showcase.

https://mchb.hrsa.gov/funding/challenge-competitions/p4challenge/our-20-final-winners/winners-

showcase

Morbidity and Mortality Weekly Report. (2022). # 1 Immunization Rates of 16-18 month old Children

Enrolled in Military Treatment Facilities. https://www.cdc.gov/mmwr/volumes/71/wr/mm7116a1.htm





Questions





To receive CE/CME credit, you must register by 0800 on 29 APR 2022 to qualify for the receipt of CE/CME credit or a certificate of attendance. You must complete the program posttest and evaluation before collecting your certificate. The posttest and evaluation will be available through 12 MAY 2022 at 2359 ET. Please complete the following steps to obtain CE/CME credit:

- 1. Go to URL: <u>https://www.dhaj7-cepo.com/content/apr-2022-ccss-military-children-and-youth</u>
- 2. Search for your course using the Catalog, Calendar, or Find a course search tool.
- 3. Click on the REGISTER/TAKE COURSE tab.
 - a. If you have previously used the CEPO CMS, click login.
 - b. If you have not previously used the CEPO CMS click register to create a new account.
- 4. Follow the onscreen prompts to complete the post-activity assessments:
 - a. Read the Accreditation Statement
 - b. Complete the Evaluation
 - c. Take the Posttest
- 5. After completing the posttest at 80% or above, your certificate will be available for print or download.
- 6. You can return to the site at any time in the future to print your certificate and transcripts at: <u>https://www.dhaj7-cepo.com/</u>
- 7. If you require further support, please contact us at: <u>dha.ncr.j7.mbx.cepo-cms-support@mail.mil</u>



