# Immunization Coalition of Los Angeles County (ICLAC) Spring 2023 Vaccine Forum

# Routine Immunizations on Schedule for Everyone (RISE)

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## Disclosures

- Richard Dang
  - President, California Pharmacists Association
  - Vice Chair, Immunization Collation of Los Angeles County
  - Advisory Board, GSK (completed)

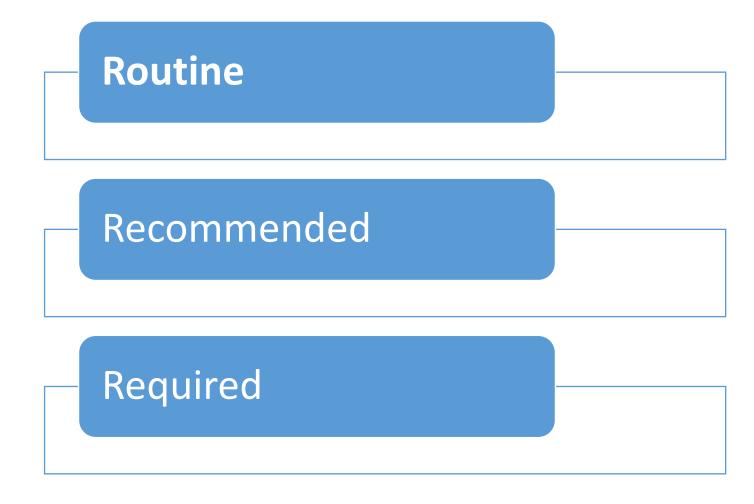


## Objectives

- 1) Review the CDC 2023 Immunization Schedules
- 2) Discuss helpful resources for clinicians and patients



## The Three "R's" of Vaccines





## Let's RISE



RSE

CDC is launching a new initiative Let's RISE - Routine Immunizations on-Schedule for Everyone



The COVID-19 pandemic caused disruptions in routine health services including vaccinations for adults and children.



Routine vaccine orders and administration are rebounding but unevenly. The rebound may be insufficient to catch up everyone that missed a vaccine dose in 2020 and 2021.



The full impact of the COVID-19 pandemic on routine vaccination coverage will take time to determine.



While we continue to investigate, CDC is prioritizing getting everyone caught-up on routine immunizations that were missed or delayed and continuing to strengthen adult vaccination rates above pre-pandemic levels.



Routine Immunizations on Schedule for Everyone | 2022 Data Review www.cdc.gov/vaccines/partners/routine-immunizations-lets-rise.html





After the COVID-19 pandemic started, we saw a concerning drop in childhood routine vaccinations. Provider vaccine orders for children through the Vaccines for Children (VFC) program decreased by 14% in 2020 after the national COVID-19 public health emergency was declared.

### SIGNS OF IMPROVEMENT

Childhood vaccine orders through the Vaccines for Children (VFC) program and administration have steadily rebounded. As of January 2023, childhood VFC vaccine orders are within 1% or exceed pre-pandemic levels.



Vaccination coverage has remained high and stable overall among young children, with coverage of more than **9 in 10** children for most recommended vaccines nationally

### **IMPROVEMENT NEEDED**

Vaccination coverage among certain groups declined since the start of the COVID-19 pandemic and may have not yet recovered.

Combined 7-vaccine series coverage by age 24 months fell 4-5% among children living below poverty or in rural areas.



### Kindergarten vaccination

coverage has steadily declined for all vaccines and in a majority of states over the past 2 school years to lowest levels in a decade.

### At least 1/4 Million

children that entered kindergarten during the pandemic are potentially susceptible to vaccine preventable disease.

Disparities in vaccination coverage by race/ethnicity, income, and insurance status that existed prior to the pandemic persisted in 2021 but did not worsen during the pandemic. However disparities worsened for young children living in rural areas.

### **IMMEDIATE ACTION NEEDED**

We must act now to catch up children who delayed or missed getting routine vaccinations during the COVID-19 pandemic.

Getting routine immunizations back on track is a goal that we can achieve by working together to reduce barriers, increase access, and strengthen vaccine confidence.







After the COVID-19 pandemic started, we saw a concerning drop in routine adolescent vaccinations. Provider orders for adolescent vaccines through the Vaccines for Children (VFC) program also decreased by approximately 25% for HPV, Men-ACWY, and Tdap year 2020 (October 2019-September 2020) as compared to the same time the previous year.

### SIGNS OF IMPROVEMENT

Adolescent vaccine orders through the Vaccines for Children (VFC) program and administration have steadily rebounded. Tdap and MenACWY VFC orders now exceed pre-pandemic levels.



Vaccination coverage has remained high and stable overall among children 13-17 years of age, with with coverage of more than 9 in 10 teenagers for recommended Tdap and Meningococcal vaccines nationally. Coverage for HPV also increased slightly with nearly 1 in 8 teens receiving a first dose of this vaccine.

### **IMPROVEMENT NEEDED**

Adolescent vaccination coverage for MenACWY and Tdap **fell 4-5%** among teens that became eligible for these vaccines during the pandemic. HPV vaccine orders through the Vaccines for Children (VFC) program have remained around **10% lower** than pre-pandemic levels.



### **IMMEDIATE ACTION NEEDED**

We must act now to catch up teens who delayed or missed getting routine vaccinations during the COVID-19 pandemic.

Getting routine immunizations back on track is a goal that we can achieve by working together to reduce barriers, increase access, and strengthen vaccine confidence.







Majority of U.S. Adults
Are Missing
Routine
Vaccinations

At least 3 out of every 4 adults are missing one or more routinely recommended vaccines



## **CDC Immunization Schedule**

- Updated annually, published February of each calendar year
- 2 schedules:
  - Child and Adolescent Schedule
  - Adult Schedule
- Routine vaccine recommendations based on:
  - Age
  - Medical Conditions/Special Conditions
- Contents:
  - Cover Page
  - Schedules
  - Catch-up Recommendations
  - Footnotes
  - Contraindications/Precautions



## What's New? Child/Adolescent Schedule

- Added COVID19, Priorix®, and PCV15 vaccines
- New abbreviations for COVID19 vaccines (1v, 2v, mRNA, aPS)
- Added guidance, revisions, or clarifications for: COVID-19, dengue, hepatitis B, influenza, MMR, meningococcal ACWY, menB, pneumococcal, poliovirus vaccines



## What's New? Adult Schedule

- Added COVID19, PreHevbrio®, and Priorix®
- New abbreviations for COVID19 vaccines (1v, 2v, mRNA, aPS)
- Added APhA has a new approving partner
- Added guidance, revisions, or clarifications for: COVID-19, hepatitis B, influenza, MMR, meningococcal, pneumococcal, poliovirus, Zoster vaccines



UNITED STATES

### Vaccines in the Child and Adolescent Immunization Schedule\*

Vaccine	Abbreviation(s)	Trade name(s)
COVID-19	1vCOV-mRNA	Comirnaty®/Pfizer- BioNTech COVID-19 Vaccine
		SPIKEVAX®/Moderna COVID-19 Vaccine
	2vCOV-mRNA	Pfizer-BioNTech COVID-19 Vaccine, Bivalent
		Moderna COVID-19 Vaccine, Bivalent
	1vCOV-aPS	Novavax COVID-19 Vaccine
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)	ActHIB® Hiberix®
	Hib (PRP-OMP)	PedvaxHIB®
Hepatitis A vaccine	НерА	Havrix <sup>®</sup> Vaqta <sup>®</sup>
Hepatitis B vaccine	НерВ	Engerix-B <sup>®</sup> Recombivax HB <sup>®</sup>
Human papillomavirus vaccine	HPV	Gardasil 9°
nfluenza vaccine (inactivated)	IIV4	Multiple
nfluenza vaccine (live, attenuated)	LAIV4	FluMist® Quadrivalent
Measles, mumps, and rubella vaccine	MMR	M-M-R II® Priorix®
Meningococcal serogroups A, C, W, Y vaccine	MenACWY-D	Menactra®
	MenACWY-CRM	Menveo®
	MenACWY-TT	MenQuadfi <sup>6</sup>
Meningococcal serogroup B vaccine	MenB-4C	Bexsero®
	MenB-FHbp	Trumenba®
Pneumococcal conjugate vaccine	PCV13 PCV15	Prevnar 13 <sup>®</sup> Vaxneuvance™
Pneumococcal polysaccharide vaccine	PPSV23	Pneumovax 23®
Poliovirus vaccine (inactivated)	IPV	IPOL®
Rotavirus vaccine	RV1 RV5	Rotarix <sup>®</sup> RotaTeq <sup>®</sup>
Tetanus, diphtheria, and acellular pertussis vaccine	Tdap	Adacel® Boostrix®
Tetanus and diphtheria vaccine	Td	Tenivac® Tdvax™
Varicella vaccine	VAR	Varivax <sup>®</sup>
Combination vaccines (use combination vaccines instead of separe		
OTaP, hepatitis B, and inactivated poliovirus vaccine	DTaP-HepB-IPV	Pediarix®
OTaP, inactivated poliovirus, and Haemophilus influenzae type b vaccine	DTaP-IPV/Hib	Pentacel®
DTaP and inactivated poliovirus vaccine	DTaP-IPV	Kinrix <sup>e</sup> Quadracel <sup>e</sup>
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 unkno extended intervals between doses. When a vaccine is not administered at the reco The use of trade names is for identification purposes only and does not imply endo	mmended age, administer	at a subsequent visit.

### How to use the child and adolescent immunization schedule

Determine recommended

vaccine by age (Table 1)

Determine recommended interval for catchup vaccination (Table 2)

Assess need for additional recommended vaccines by or other indication (Notes) (Table 3)

Review vaccine intervals, and medical condition special situations (Appendix)

Review types, frequencies, contraindications and precautions considerations for for vaccine types

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.aapa.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
- 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

Scan QR code for access to online schedule









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).

o determine minimum intervals betwe 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 vrs	11–12 yrs	13–15 vrs	16 yrs	17-18 )
Hepatitis B (HepB)	1 <sup>st</sup> dose		dose▶		4		3 <sup>rd</sup> dose										
Rotavirus (RV): RV1 (2-dose series), RV5 (3-dose series)			1 <sup>st</sup> dose	2 <sup>nd</sup> dose	See Notes												
Piphtheria, tetanus, acellular pertussis DTaP <7 yrs)			1st dose	2 <sup>nd</sup> dose	3 <sup>rd</sup> dose			<b>◄</b> 4 <sup>th</sup> c	lose▶			5 <sup>th</sup> dose					
aemophilus influenzae type b (Hib)			1st dose	2 <sup>nd</sup> dose	See Notes		<3 <sup>rd</sup> or 4 See N	th dose, Notes									
Pneumococcal conjugate PCV13, PCV15)			1 <sup>st</sup> dose	2 <sup>nd</sup> dose	3 <sup>rd</sup> dose		<b>◄</b> 4 <sup>th</sup> 0	lose>									
nactivated poliovirus IPV <18 yrs)			1 <sup>st</sup> dose	2 <sup>nd</sup> dose	<b>◄</b>		3 <sup>rd</sup> dose					4 <sup>th</sup> dose					S No
OVID-19 (1vCOV-mRNA, vCOV-mRNA, 1vCOV-aPS)									2- or 3-	dose primar	y series and	booster (Se	ee Notes)				
nfluenza (IIV4)								Annual vac	cination 1 c	r 2 doses				Annu	al vaccinatio	on 1 dose o	nly
nfluenza (LAIV4)												ual vaccinat or 2 doses	ion	Ann	ual vaccinati	on 1 dose o	only
Measles, mumps, rubella (MMR)					Seel	Notes	<b>◄</b> 1 <sup>st</sup> c	lose>				2 <sup>nd</sup> dose					
/aricella (VAR)							<b>◄</b> 1 <sup>st</sup> c	lose				2 <sup>nd</sup> dose					
lepatitis A (HepA)					See I	Votes		2-dose serie	es, See Note	s							
etanus, 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						1 <sup>st</sup> dose		2 <sup>nd</sup> dose	
Meningococcal B MenB-4C, MenB-FHbp)															See No	otes	
rneumococcal polysaccharide PPSV23)														See Notes			
engue (DEN4CYD; 9-16 yrs)															itive in ende areas (See N		
Range of recommended ages for all children		recommenc up vaccinati			nge of recor certain high				mended va			ecommende n shared clin				recomme ot applicab	





## Recommended Catch-up Immunization Schedule for Children and Adolescents Who Start Late or Who Are More than 1 Month Behind, United States, 2023

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
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 1st 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 (ActHib®, Pentacel®, Hiberix®), Vaxelis® 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 1st 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 1st 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* birthday 8 weeks (as final dose for healthy children) if 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) 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 is only necessary for children aged 12 through 59 months regardless of risk, or age 60 through 71 months with any risk, who received 3 doses before age 12 months.	
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	
	2,000		Children and adolescents age 7 through 18 years		
Meningococcal ACWY	Not applicable (N/A)	8 weeks	- Innaran and dassescents age 7 through 10 years		
Tetanus, diphtheria; tetanus, diphtheria, and acellular pertussis	7 years	4 weeks	4 weeks if first dose of DTaP/DT was administered before the 1st birthday 6 months (as final dose) if first dose of DTaP/DT or Tdap/Td was administered at or after the 1st birthday	6 months if first dose of DTaP/DT was administered before the 1st 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. 4 weeks if age 13 years or older			
Dengue	9 years	6 months	6 months		





### Recommended Child and Adolescent Immunization Schedule by Medical Indication, United States, 2023

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

		INDICATION									
VACCINE	Pregnancy	Immunocom- promised status (excluding HIV infection)	HIV infection <15% or total CD4 cell count of <200/mm <sup>3</sup>	n CD4+ count <sup>a</sup> ≥15% and total CD4 cell count of ≥200/mm <sup>3</sup>	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		SCID⁵									
Diphtheria, tetanus, and acellular pertussis (DTaP)											
Haemophilus influenzae type b											
Pneumococcal conjugate											
Inactivated poliovirus											
COVID-19		See Notes	See	Notes							
Influenza (IIV4)											
Influenza (LAIV4)			-4			Asthma, wheezing: 2–4yrs <sup>c</sup>					
Measles, mumps, rubella	*										
Varicella	*										
Hepatitis A											
Tetanus, diphtheria, and acellular pertussis (Tdap)											
Human papillomavirus	*										
Meningococcal ACWY											
Meningococcal B											
Pneumococcal polysaccharide											
Dengue											
Vaccination according t routine schedule recommended		Recommended for persons with an addition factor for which the vac would be indicated	onal risk 🛗 a ccine r	/accination is recomr and additional doses necessary based on n condition or vaccine.	may be nedical	recaution-vaccine night be indicated if benefit of protection outweighs risk of adverse reaction	be adminis	ded-vaccine should not	No recomme applicable	ndation/not	

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.

b. Severe Combined Immunodeficiency

c. LAIV4 contraindicated for children 2–4 years of age with asthma or wheezing during the preceding 12 months





Adult Immunization Schedule, 2023.

index.html.

Additional information

- Consult relevant ACIP statements for detailed recommendations at www.cdc.gov/vaccines/hcp/acip-recs/
- 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 3-2, 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.qov/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/immunocompetence.html, and Immunization in Special Clinical Circumstances (In: Kimberlin DW, Barnett ED, Lynfield Ruth, Sawyer MH, eds. *Red Book: 2021–2024 Report of the Committee on Infectious Diseases.* 32nd ed. Itasca, IL: American Academy of Pediatrics; 2021:72–86).
- For information about vaccination in the setting of a vaccine-preventable 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 vaccines included in the child and adolescent vaccine schedule are covered by VICP except dengue, PPSV23, and COVID-19 vaccines. COVID-19 vaccines that are authorized or approved by the FDA are covered by the Countermeasures Injury Compensation Program (CICP). For more information, see www.hrsa.gov/vaccinecompensation or www.hrsa.gov/cicp.

### **COVID-19 vaccination**

(minimum age: 6 months [Moderna and Pfizer-BioNTech COVID-19 vaccines], 12 years [Novavax COVID-19 Vaccine])

### **Routine vaccination**

- Primary series:
- Age 6 months-4 years: 2-dose series at 0, 4-8 weeks (Moderna) or 3-dose series at 0, 3-8, 11-16 weeks (Pfizer-BioNTech)
- Age 5-11 years: 2-dose series at 0, 4-8 weeks (Moderna) or 2-dose series at 0, 3-8 weeks (Pfizer-BioNTech)
- Age 12–18 years: 2-dose series at 0, 4-8 weeks (Moderna) or 2-dose series at 0, 3-8 weeks (Novavax, Pfizer-BioNTech)
- For booster dose recommendations see www.cdc. gov/vaccines/covid-19/clinical-considerations/interimconsiderations-us.html

### Special situations

Persons who are moderately or severely immunocompromised

- Primary series
- Age 6 months-4 years: 3-dose series at 0, 4, 8 weeks (Moderna) or 3-dose series at 0, 3, 11 weeks (Pfizer-BioNTech)
- Age 5–11 years: 3-dose series at 0, 4, 8 weeks (Moderna) or 3-dose series at 0, 3, 7 weeks (Pfizer-BioNTech)
- Age 12–18 years: 3-dose series at 0, 4, 8 weeks (Moderna) or 2-dose series at 0, 3 weeks (Novavax) or 3-dose series at 0, 3, 7 weeks (Pfizer-BioNTech)
- Booster dose: see www.cdc.gov/vaccines/covid-19/clinicalconsiderations/interim-considerations-us.html
- Pre-exposure prophylaxis (monoclonal antibodies) may be considered to complement COVID-19 vaccination. See www.cdc.gov/vaccines/covid-19/clinical-considerations/ interim-considerations-us.html#immunocompromised

For Janssen COVID-19 Vaccine recipients see COVID-19 schedule at www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html

**Note:** Administer an age-appropriate vaccine product for each dose. Current COVID-19 schedule and dosage formulation available at www.cdc.gov/vaccines/covid-19/downloads/COVID-19-immunization-schedule-ages-6months-older. pdf. For more information on Emergency Use Authorization (EUA) indications for COVID-19 vaccines, see www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/covid-19-vaccines.

### **Dengue vaccination** (minimum age: 9 years)

#### Routine vaccination

- Age 9–16 years living in areas with endemic dengue 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 <a href="https://www.cdc.gov/mmwr/volumes/70/rr/r7006a1.htm?s\_cid=rr7006a1\_w">www.cdc.gov/mmwr/volumes/70/rr/r7006a1.htm?s\_cid=rr7006a1\_w</a> and <a href="https://www.cdc.gov/dengue/vaccine/hcp/index.html">www.cdc.gov/dengue/vaccine/hcp/index.html</a>
- Dengue vaccine should not be administered to children traveling to or visiting endemic dengue areas.

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 4<sup>th</sup> 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 age12–15 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 age12–59 months and at least 8 weeks after dose 2.
- 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. Vaxelis® can be used for catch-up 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

- 1 dose

### Elective splenectomy:

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

### 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 no doses (age 15 months or older)

### **Hepatitis A vaccination** (minimum age: 12 months for routine vaccination)

#### Routine vaccination

 2-dose series (minimum interval: 6 months) at age 12–23 months

### **Catch-up vaccination**

- Unvaccinated persons through age 18 years should complete a 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.

 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).

### International travel

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

### **Hepatitis B vaccination** (minimum age: birth)

### Routine vaccination

- 3-dose series at age 0, 1–2, 6–18 months (use monovalent HepB vaccine for doses administered before age 6 weeks)
- Birth weight ≥2,000 grams: 1 dose within 24 hours of birth if medically stable
- Birth weight <2,000 grams: 1 dose at chronological age 1 month or hospital discharge (whichever is earlier and even if weight is still <2,000 grams).</li>
- Infants who did not receive a birth dose should begin the series as soon as possible (see Table 2 for minimum intervals).
- Administration of 4 doses is permitted when a combination vaccine containing HepB is used after the birth dose.
- Minimum intervals (see Table 2): when 4 doses are administered, substitute "dose 4" for "dose 3" in these calculations
- Final (3rd or 4th) dose: age 6–18 months (minimum age 24 weeks)
- Mother is HBsAq-positive
- Birth dose (monovalent HepB vaccine only): administer HepB vaccine and hepatitis B immune globulin (HBIG) (in separate limbs) within 12 hours of birth, regardless of birth weight.
- Birth weight <2000 grams: administer 3 additional doses of HepB vaccine beginning at age 1 month (total of 4 doses)
- Final (3rd or 4th) dose: administer at age 6 months (minimum age 24 weeks)
- Test for HBsAg and anti-HBs at age 9–12 months. If HepB series is delayed, test 1–2 months after final dose. Do not test before age 9 months.



#### Mother is HBsAg-unknown

If other evidence suggestive of maternal hepatitis B infection exists (e.g., presence of HBV DNA, HBeAg-positive, or mother known to have chronic hepatitis B infection), manage infant as if mother is HBsAg-positive

- Birth dose (monovalent HepB vaccine only):
- · Birth weight ≥2,000 grams: administer **HepB vaccine** within 12 hours of birth. Determine mother's HBsAg status as soon as possible. If mother is determined to be HBsAgpositive, administer **HBIG** as soon as possible (in separate limb), but no later than 7 days of age.
- · Birth weight <2,000 grams: administer **HepB vaccine** and **HBIG** (in separate limbs) within 12 hours of birth. Administer 3 additional doses of **HepB vaccine** beginning at age 1 month (total of 4 doses)
- Final (3rd or 4th) dose: administer at age 6 months (minimum age 24 weeks)
- If mother is determined to be HBsAg-positive or if status remains unknown, test for HBsAg and anti-HBs at age 9–12 months. If HepB series is delayed, test 1–2 months after final dose. Do not test before age 9 months.

### **Catch-up vaccination**

- Unvaccinated persons should complete a 3-dose series at 0, 1–2, 6 months. See Table 2 for minimum intervals
- 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:
- Heplisav-B<sup>®</sup>: 2-dose series at least 4 weeks apart
- PreHevbrio®: 3-dose series at 0, 1, and 6 months
- -Combined HepA and HepB vaccine, **Twinrix**\*: 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).

### **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 HBsAq-positive mothers
- Persons who are predialysis or on maintenance dialysis
- -Other immunocompromised persons
- For detailed revaccination recommendations, see www.cdc. gov/vaccines/hcp/acip-recs/vacc-specific/hepb.html.

**Note:** Heplisav-B and PreHevbrio are not recommended in pregnancy due to lack of safety data in pregnant persons

### Human papillomavirus vaccination (minimum age: 9 years)

### Routine and catch-up vaccination

- HPV vaccination routinely recommended at age 11–12 years (can start at age 9 years) and catch-up HPV vaccination recommended for all persons through age 18 years if not adequately vaccinated
- 2- or 3-dose series depending on age at initial vaccination:
- Age 9–14 years at initial vaccination: 2-dose series at 0, 6–12 months (minimum interval: 5 months; repeat dose if administered too soon)
- Age 15 years or older at initial vaccination: 3-dose series at 0, 1–2 months, 6 months (minimum intervals: dose 1 to dose 2: 4 weeks / dose 2 to dose 3: 12 weeks / dose 1 to dose 3: 5 months; repeat dose if administered too soon)
- Interrupted schedules: If vaccination schedule is interrupted, the series does not need to be restarted.
- No additional dose recommended when any HPV vaccine series has been completed using the recommended dosing intervals.

### Special situations

- Immunocompromising conditions, including HIV infection: 3-dose series, even for those who initiate vaccination at age 9 through 14 years.
- History of sexual abuse or assault: Start at age 9 years
- Pregnancy: Pregnancy testing not needed before vaccination; HPV vaccination not recommended until after pregnancy; no intervention needed if vaccinated while pregnant

#### Influenza vaccination

(minimum age: 6 months [IIV], 2 years [LAIV4], 18 years [recombinant influenza vaccine, RIV4])

### **Routine vaccination**

- Use any influenza vaccine appropriate for age and health status annually:
- 2 doses, separated by at least 4 weeks, for **children age 6 months-8 years** who have received fewer than 2 influenza vaccine doses before July 1, 2022, or whose influenza vaccination history is unknown (administer dose 2 even if the child turns 9 between receipt of dose 1 and dose 2)
- 1 dose for children age 6 months-8 years who have received at least 2 influenza vaccine doses before July 1, 2022
- 1 dose for all persons age 9 years or older

- For the 2022-2023 season, see www.cdc.gov/mmwr/volumes/71/rr/rr7101a1.htm.
- For the 2023–24 season, see the 2023–24 ACIP influenza vaccine recommendations.

### Special situations

- Egg allergy, hives only: Any influenza vaccine appropriate for age and health status annually
- 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 or LAIV4, administer
   in medical setting under supervision of health care provider
   who can recognize and manage severe allergic reactions.
- Severe allergic reaction (e.g., anaphylaxis) to a vaccine component or a previous dose of any influenza vaccine: see Appendix listing contraindications and precautions
- Close contacts (e.g., caregivers, healthcare personnel)
   of severely immunosuppressed persons who require a
   protected environment: these persons should not receive
   LAIV4. If LAIV4 is given, they should avoid contact with/
   caring for such immunosuppressed persons for 7 days after
   vaccination.

Measles, mumps, and rubella vaccination (minimum age: 12 months for routine vaccination)

#### Routine vaccination

- 2-dose series at age 12–15 months, age 4–6 years
- MMR or MMRV may be administered

**Note:** For dose 1 in children age 12–47 months, it is recommended to administer MMR and varicella vaccines separately. MMRV may be used if parents or caregivers express a preference.

### Catch-up vaccination

- Unvaccinated children and adolescents: 2-dose series at least 4 weeks apart
- The maximum age for use of MMRV is 12 years.
- Minimum interval between MMRV doses: 3 months



### **Special situations**

- International travel
- Infants age 6–11 months: 1 dose before departure; revaccinate with 2-dose series at age 12–15 months (12 months for children in high-risk areas) and dose 2 as early as 4 weeks later.
- Unvaccinated children age 12 months or older:
   2-dose series at least 4 weeks apart before departure
- In mumps outbreak settings, for information about additional doses of MMR (including 3rd dose of MMR), see www.cdc.gov/mmwr/volumes/67/wr/mm6701a7.htm

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 booster at age 16–18 years (minimum interval: 8 weeks)
- Age 16-18 years: 1 dose

### Special situations

Anatomic or functional asplenia (including sickle cell disease), HIV infection, persistent complement component deficiency, complement inhibitor (e.g., eculizumab, ravulizumab) use:

- Menveo®\*
- Dose 1 at age 2 months: 4-dose series (additional 3 doses at age 4, 6, and 12 months)
- Dose 1 at age 3–6 months: 3- or 4-dose series (dose 2 [and dose 3 if applicable] at least 8 weeks after previous dose until a dose is received at age 7 months or older, followed by an additional dose at least 12 weeks later and after age 12 months)
- Dose 1 at age 7–23 months: 2-dose series (dose 2 at least 12 weeks after dose 1 and after age 12 months)
- Dose 1 at age 24 months or older: 2-dose series at least 8 weeks apart

#### Menactra®

- Persistent complement component deficiency or complement inhibitor use:
- · Age 9–23 months: 2-dose series at least 12 weeks apart
- · Age 24 months or older: 2-dose series at least 8 weeks apart

- Anatomic or functional asplenia, sickle cell disease, or HIV infection:
- · Age 9-23 months: Not recommended
- Age 24 months or older: 2-dose series at least 8 weeks apart
- Menactra® must be administered at least 4 weeks after completion of PCV series.

### MenQuadfi®

 Dose 1 at age 24 months or older: 2-dose series at least 8 weeks apart

Travel to countries with hyperendemic or epidemic meningococcal disease, including countries in the African meningitis belt or during the Haji (www.cdc.gov/travel/):

- Children less than age 24 months:
- Menveo®\* (age 2-23 months)
- · Dose 1 at age 2 months: 4-dose series (additional 3 doses at age 4, 6, and 12 months)
- Dose 1 at age 3–6 months: 3- or 4-dose series (dose 2 [and dose 3 if applicable] at least 8 weeks after previous dose until a dose is received at age 7 months or older, followed by an additional dose at least 12 weeks later and after age 12 months)
- Dose 1 at age 7–23 months: 2-dose series (dose 2 at least 12 weeks after dose 1 and after age 12 months)
- Menactra® (age 9–23 months)
- 2-dose series (dose 2 at least 12 weeks after dose 1; dose 2 may be administered as early as 8 weeks after dose 1 in travelers)
- Children age 2 years or older: 1 dose Menveo<sup>®\*</sup>, Menactra<sup>®</sup>, or MenQuadfi<sup>®</sup>

First-year college students who live in residential housing (if not previously vaccinated at age 16 years or older) or military recruits:

1 dose Menveo®\*, Menactra®, or MenQuadfi®

Adolescent vaccination of children who received MenACWY prior to age 10 years:

- Children for whom boosters are recommended because of an ongoing increased risk of meningococcal disease (e.g., those with complement component deficiency, HIV, or asplenia): Follow the booster schedule for persons at increased risk.
- Children for whom boosters are not recommended (e.g., a healthy child who received a single dose for travel to a country where meningococcal disease is endemic): Administer MenACWY according to the recommended adolescent schedule with dose 1 at age 11–12 years and dose 2 at age 16 years.

\*Menveo has two formulations: lyophilized and liquid. The liquid formulation should not be used before age 10 years.

**Note: Menactra®** should be administered either before or at the same time as DTaP. MenACWY may be administered simultaneously with MenB vaccines if indicated, but at a different anatomic site, if feasible.

For MenACWY **booster dose recommendations** for groups listed under "Special situations" and in an outbreak setting and additional meningococcal vaccination information, see <a href="https://www.cdc.gov/mmwr/volumes/69/rr/rr6909a1.htm">www.cdc.gov/mmwr/volumes/69/rr/rr6909a1.htm</a>.

Meningococcal serogroup B vaccination (minimum age: 10 years [MenB-4C, Bexsero\*; MenB-FHbp, Trumenba\*])

### Shared clinical decision-making

- Adolescents not at increased risk age 16–23 years (preferred age 16–18 years) based on shared clinical decision-making:
- Bexsero®: 2-dose series at least 1 month apart
- Trumenba®: 2-dose series at least 6 months apart (if dose 2 is administered earlier than 6 months, administer a 3rd dose at least 4 months after dose 2)

### Special situations

Anatomic or functional asplenia (including sickle cell disease), persistent complement component deficiency, complement inhibitor (e.g., eculizumab, ravulizumab) use:

- Bexsero®: 2-dose series at least 1 month apart
- Trumenba®: 3-dose series at 0, 1–2, 6 months (if dose 2 was administered at least 6 months after dose 1, dose 3 not needed; if dose 3 is administered earlier than 4 months after dose 2, a 4<sup>th</sup> dose should be administered at least 4 months after dose 3)

**Note:** Bexsero® and Trumenba® are not interchangeable; the same product should be used for all doses in a series.

For MenB **booster dose recommendations** for groups listed under "Special situations" and in an outbreak setting and additional meningococcal vaccination information, see www.cdc.gov/mmwr/volumes/69/rr/rr6909a1.htm.



Pneumococcal vaccination (minimum age: 6 weeks [PCV13], [PCV15], 2 years [PPSV23])

### **Routine vaccination with PCV**

4-dose series at 2, 4, 6, 12–15 months

### Catch-up vaccination with PCV

- Healthy children age 24–59 months with any incomplete\* PCV series: 1 dose PCV
- For other catch-up guidance, see Table 2.

**Note:** PCV13 and PCV15 can be used interchangeably for children who are healthy or have underlying conditions. PCV15 is not indicated for children who have received 4 doses of PCV13 or another age appropriate complete PCV13 series.

### **Special situations**

Underlying conditions below: When both PCV and PPSV23 are indicated, administer PCV first. PCV and PPSV23 should not be administered during the same visit.

Chronic heart disease (particularly cyanotic congenital heart disease and cardiac failure); chronic lung disease (including asthma treated with high-dose, oral corticosteroids); diabetes mellitus:

#### Age 2-5 years

- Any incomplete\* series with:
- 3 PCV doses: 1 dose PCV (at least 8 weeks after any prior PCV dose)
- Less than 3 PCV doses: 2 doses PCV (8 weeks after the most recent dose and administered 8 weeks apart)
- No history of PPSV23: 1 dose PPSV23 (at least 8 weeks after completing all recommended PCV doses)

#### Age 6–18 years

- Any incomplete\* series with PCV: no further PCV doses needed
- No history of PPSV23: 1 dose PPSV23 (at least 8 weeks after completing all recommended PCV doses)

### Cerebrospinal fluid leak, cochlear implant:

#### Age 2-5 years

- Any incomplete\* series with:
- 3 PCV doses: 1 dose PCV (at least 8 weeks after any prior PCV dose)
- Less than 3 PCV doses: 2 doses PCV (8 weeks after the most recent dose and administered 8 weeks apart)
- No history of PPSV23: 1 dose PPSV23 (at least 8 weeks after completing all recommended PCV doses)

#### Age 6–18 years

- No history of either PCV or PPSV23: 1 dose PCV, 1 dose PPSV23 at least 8 weeks later
- Any PCV but no PPSV23: 1 dose PPSV23 at least 8 weeks after the most recent dose of PCV
- PPSV23 but no PCV: 1 dose PCV at least 8 weeks after the most recent dose of PPSV23

Sickle cell disease and other hemoglobinopathies; anatomic or functional asplenia; congenital or acquired immunodeficiency; HIV infection; chronic renal failure; nephrotic syndrome; malignant neoplasms, leukemias, lymphomas, Hodgkin disease, and other diseases associated with treatment with immunosuppressive drugs or radiation therapy; solid organ transplantation; multiple myeloma:

#### Age 2-5 years

- Any incomplete\* series with:
- 3 PCV doses: 1 dose PCV (at least 8 weeks after any prior PCV dose)
- Less than 3 PCV doses: 2 doses PCV (8 weeks after the most recent dose and administered 8 weeks apart)
- No history of PPSV23: 1 dose PPSV23 (at least 8 weeks after completing all recommended PCV doses) and a dose 2 of PPSV23 5 years later

#### Age 6-18 years

- No history of either PCV or PPSV23: 1 dose PCV, 2 doses PPSV23 (dose 1 of PPSV23 administered 8 weeks after PCV and dose 2 of PPSV23 administered at least 5 years after dose 1 of PPSV23)
- Any PCV but no PPSV23: 2 doses PPSV23
   (dose 1 of PPSV23 administered 8 weeks after the most recent dose of PCV and dose 2 of PPSV23 administered at least 5 years after dose 1 of PPSV23)
- PPSV23 but no PCV: 1 dose PCV at least 8 weeks after the most recent PPSV23 dose and a dose 2 of PPSV23 administered 5 years after dose 1 of PPSV23 and at least 8 weeks after a dose of PCV
- \*Incomplete series = Not having received all doses in either the recommended series or an age-appropriate catch-up series see Table 2 in ACIP pneumococcal recommendations at www.cdc.gov/mmwr/volumes/71/wr/mm7137a3.htm

For guidance on determining which pneumococcal vaccines a patient needs and when, please refer to the mobile app, which can be downloaded here: www.cdc.gov/vaccines/vpd/pneumo/hcp/pneumoapp.html

### 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 previous dose.

### 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\_%20 cid=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.

### Special situations

- Adolescents aged 18 years at increased risk of exposure to poliovirus with:
- No evidence of a complete polio vaccination series (i.e., at least 3 doses): administer remaining doses (1, 2, or 3 doses) to complete a 3-dose series
- Evidence of completed polio vaccination series (i.e., at least 3 doses): may administer one lifetime IPV booster

For detailed information, see: www.cdc.gov/vaccines/vpd/ polio/hcp/recommendations.html





### Rotavirus vaccination (minimum age: 6 weeks)

#### Routine vaccination

- Rotarix®: 2-dose series at age 2 and 4 months
- RotaTeq®: 3-dose series at age 2, 4, and 6 months
- If any dose in the series is either RotaTeq® or unknown, default to 3-dose series.

### Catch-up vaccination

- Do not start the series on or after age 15 weeks, 0 days.
- The maximum age for the final dose is 8 months, 0 days.
- For other catch-up 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 tetanus- and diphtheria-toxoid-containing vaccine.

### **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 (minimum age: 12 months)

#### Routine vaccination

- 2-dose series at age 12-15 months, 4-6 years
- VAR or MMRV may be administered\*
- Dose 2 may be administered as early as 3 months after dose 1 (a dose inadvertently administered after at least 4 weeks may be counted as valid)
- \*Note: For dose 1 in children age 12–47 months, it is recommended to administer MMR and varicella vaccines separately. MMRV may be used if parents or caregivers express a preference.

### **Catch-up vaccination**

- Ensure persons age 7–18 years without evidence of immunity (see MMWR at www.cdc.gov/mmwr/pdf/rr/rr5604.pdf) have a 2-dose series:
- Age 7-12 years: Routine interval: 3 months (a dose inadvertently administered after at least 4 weeks may be counted as valid)
- Age 13 years and older: Routine interval: 4–8 weeks (minimum interval: 4 weeks)
- The maximum age for use of MMRV is 12 years.





### **Guide to Contraindications and Precautions to Commonly Used Vaccines**

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

For COVID-19 vaccine contraindications and precautions see www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html#contraindications

Vaccine	Contraindicated or Not Recommended <sup>1</sup>	Precautions <sup>2</sup>
Influenza, egg-based, inactivated injectable (IIV4)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after previous dose of any influenza vaccine (i.e., any egg-based IIV, ccIIV, RIV, or LAIV of any valency)</li> <li>Severe allergic reaction (e.g., anaphylaxis) to any vaccine component<sup>3</sup> (excluding egg)</li> </ul>	<ul> <li>Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine</li> <li>Moderate or severe acute illness with or without fever</li> </ul>
Influenza, cell culture-based inactivated injectable [(ccllV4), Flucelvax <sup>®</sup> Quadrivalent]	• Severe allergic reaction (e.g., anaphylaxis) to any ccIIV of any valency, or to any component <sup>3</sup> of ccIIV4	<ul> <li>Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine</li> <li>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 setting under supervision of health care provider who can recognize and manage severe allergic reactions. May consult an allergist.</li> <li>Moderate or severe acute illness with or without fever</li> </ul>
Influenza, recombinant injectable [(RIV4), Flublok° Quadrivalent]	• Severe allergic reaction (e.g., anaphylaxis) to any RIV of any valency, or to any component <sup>3</sup> of RIV4	<ul> <li>Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine</li> <li>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.</li> <li>Moderate or severe acute illness with or without fever</li> </ul>
Influenza, live attenuated [LAIV4, Flumist <sup>©</sup> Quadrivalent]	Severe allergic reaction (e.g., anaphylaxis) after previous dose of any influenza vaccine (i.e., any egg-based IIV, ccIIV, RIV, or LAIV of any valency) Severe allergic reaction (e.g., anaphylaxis) to any vaccine component³ (excluding egg) Children age 2 – 4 years with a history of asthma or wheezing Anatomic or functional asplenia Immunocompromised due to any cause including, but not limited to, medications and HIV infection Close contacts or caregivers of severely immunosuppressed persons who require a protected environment Pregnancy Cochlear implant Active communication between the cerebrospinal fluid (CSF) and the oropharynx, nasopharynx, nose, ear or any other cranial CSF leak Children and adolescents receiving aspirin or salicylate-containing medications Received influenza antiviral medications oseltamivir or zanamivir within the previous 48 hours, peramivir within the previous 5 days, or baloxavir within the previous 17 days	<ul> <li>Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine</li> <li>Asthma in persons aged 5 years old or older</li> <li>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)]</li> <li>Moderate or severe acute illness with or without fever</li> </ul>

- 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/
- 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





Vaccine	Contraindicated or Not Recommended <sup>1</sup>	Precautions <sup>2</sup>
Dengue (DEN4CYD)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component <sup>3</sup> Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of chemotherapy, congenital immunodeficiency, long-term immunosuppressive therapy or patients with HIV infection who are severely immunocompromised)     Lack of laboratory confirmation of a previous Dengue infection	Pregnancy HIV infection without evidence of severe immunosuppression Moderate or severe acute illness with or without fever
Diphtheria, tetanus, pertussis (DTaP) Tetanus, diphtheria (DT)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup></li> <li>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</li> </ul>	<ul> <li>Guillain-Barré syndrome (GBS) within 6 weeks after previous dose of tetanus-toxoid—containing vaccine</li> <li>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</li> <li>For DTaP only: Progressive neurologic disorder, including infantile spasms, uncontrolled epilepsy, progressive encephalopathy; defer DTaP until neurologic status clarified and stabilized</li> <li>Moderate or severe acute illness with or without fever</li> </ul>
Haemophilus influenzae type b (Hib)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup></li> <li>For Hiberix, ActHib, and PedvaxHIB only: History of severe allergic reaction to dry natural latex</li> <li>Less than age 6 weeks</li> </ul>	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 <sup>3</sup> including neomycin	Moderate or severe acute illness with or without fever
Hepatitis B (HepB)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup> including yeast</li> <li>Pregnancy: Heplisav-B and PreHevbrio are not recommended due to lack of safety data in pregnant persons. Use other hepatitis B vaccines if HepB is indicated<sup>4</sup>.</li> </ul>	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 <sup>3</sup> including neomycin and yeast	Moderate or severe acute illness with or without fever
Human papillomavirus (HPV)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup></li> <li>Pregnancy: HPV vaccination not recommended.</li> </ul>	Moderate or severe acute illness with or without fever
Measles, mumps, rubella (MMR) Measles, mumps, rubella, and varicella (MMRV)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component <sup>a</sup> Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of chemotherapy, 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 (≤11 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     For MMRV only: Personal or family (i.e., sibling or parent) history of seizures of any etiology
Meningococcal ACWY (MenACWY) MenACWY-CRM (Menveo®); MenACWY-D (Menactra®); MenACWY-TT (MenQuadfi®)]	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup></li> <li>For MenACWY-D and Men ACWY-CRM only: severe allergic reaction to any diphtheria toxoidor CRM197-containing vaccine</li> <li>For MenACWY-TT only: severe allergic reaction to a tetanus toxoid-containing vaccine</li> </ul>	<ul> <li>For MenACWY-CRM only: Preterm birth if less than age 9 months</li> <li>Moderate or severe acute illness with or without fever</li> </ul>
Meningococcal B (MenB) MenB-4C (Bexsero®); MenB-FHbp (Trumenba®)]	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component <sup>3</sup>	Pregnancy     For MenB-4C only: Latex sensitivity     Moderate or severe acute illness with or without fever
Pneumococcal conjugate (PCV)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component <sup>3</sup> Severe allergic reaction (e.g., anaphylaxis) to any diphtheria-toxoid-containing vaccine or its component <sup>3</sup>	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 <sup>3</sup>	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 component <sup>3</sup>	Pregnancy     Moderate or severe acute illness with or without fever
Rotavirus (RV) [RV1 (Rotarix <sup>e</sup> ), RV5 (RotaTeq <sup>e</sup> )]	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup></li> <li>Severe combined immunodeficiency (SCID)</li> <li>History of intussusception</li> </ul>	Altered immunocompetence other than SCID     Chronic gastrointestinal disease     RV1 only: Spina bifica or bladder exstrophy     Moderate or severe acute illness with or without fever
Fetanus, diphtheria, and acellular oertussis (Tdap) Fetanus, diphtheria (Td)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup></li> <li>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</li> </ul>	<ul> <li>Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of tetanus-toxoid—containing vaccine</li> <li>History of Arthus-type hypersensitivity reactions after a previous dose of diphtheria-toxoid—containing or tetanus-toxoid—toxining vaccine; defer vaccination until at least 10 years have elapsed since the last tetanus-toxoid—containing vaccine</li> <li>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</li> <li>Moderate or severe acute illness with or without fever</li> </ul>
Varicella (VAR)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component <sup>3</sup> Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of chemotherapy, 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 (≤11 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 us 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     If using MMRV, see MMR/MMRV for additional precautions

- 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.
- 4. For information on the pregnancy exposure registries for persons who were inadvertently vaccinated with Heplisav-B or PreHevbrio while pregnant, please visit heplisavbpregnancyregistry.com/ or www.prehevbrio.com/#safety.



## Recommended Adult Immunization Schedule for ages 19 years or older

2023

### How to use the adult immunization schedule

Determine recommended vaccinations by age (Table 1)

Assess need for additional recommended vaccinations by medical condition or other indication (Table 2)

Review vaccine types, dosing frequencies and intervals, and considerations for special situations (Notes) 4 Review contraindications and precautions for vaccine types (Appendix)

### Vaccines in the Adult Immunization Schedule\*

Vaccine	Abbreviation(s)	Trade name(s)
COVID-19 vaccine	1vCOV-mRNA	Comirnaty®/Pfizer-BioNTech COVID-19 Vaccine
		SPIKEVAX®/Moderna COVID-19 Vaccine
	2vCOV-mRNA	Pfizer-BioNTech COVID-19 Vaccine, Bivalent
		Moderna COVID-19 Vaccine, Bivalent
	1vCOV-aPS	Novavax COVID-19 Vaccine
Haemophilus influenzae type b vaccine	Hib	ActHIB®
		Hiberix®
		PedvaxHIB®
Hepatitis A vaccine	НерА	Havrix®
		Vaqta®
Hepatitis A and hepatitis B vaccine	НерА-НерВ	Twinrix®
Hepatitis B vaccine	НерВ	Engerix-B®
		Heplisav-B®
		PreHevbrio®
		Recombivax HB®
Human papillomavirus vaccine	HPV	Gardasil 9°
Influenza vaccine (inactivated)	IIV4	Many brands
Influenza vaccine (live, attenuated)	LAIV4	FluMist® Quadrivalent
Influenza vaccine (recombinant)	RIV4	Flublok® Quadrivalent
Measles, mumps, and rubella vaccine	MMR	M-M-R II®
		Priorix®
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 conjugate vaccine	PCV15	Vaxneuvance™
D	PCV20	Prevnar 20™
Pneumococcal polysaccharide vaccine Poliovirus vaccine	PPSV23	Pneumovax 23° IPOL°
	IPV	
Tetanus and diphtheria toxoids	Td	Tenivac®
Takanan and dishkhada kanalda and a adhilan	T-1	Tdvax™
Tetanus and diphtheria toxoids and acellular	Tdap	Adacel®
pertussis vaccine Varicella vaccine	VAR	Boostrix® Varivax®
		1, 1, 2, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1,
Zoster vaccine, recombinant	RZV	Shingrix

<sup>\*</sup>Administer recommended vaccines if vaccination history is incomplete or unknown. Do not restart or add doses to vaccine series if there are extended intervals between doses. 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 College of Physicians (www.acponline.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.aapa.org), American Pharmacists Association (www.pharmacist.com), and Society for Healthcare Epidemiology of America (www.shea-online.org).

### Report

- Suspected cases of reportable vaccine-preventable diseases or outbreaks to the local or state health department
- Clinically significant postvaccination reactions to the Vaccine Adverse Event Reporting System at www.vaers.hhs.gov or 800-822-7967

### Injury claims

All vaccines included in the adult immunization schedule except PPSV23, RZV, and COVID-19 vaccines are covered by the National Vaccine Injury Compensation Program (VICP). COVID-19 vaccines that are authorized or approved by the FDA are covered by the Countermeasures Injury Compensation Program (CICP). For more information, see <a href="https://www.hrsa.gov/vaccinecompensation">www.hrsa.gov/cicp</a>.

#### **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
- Travel vaccine recommendations: www.cdc.gov/travel
- Recommended Child and Adolescent Immunization Schedule, United States, 2023: www.cdc.gov/vaccines/schedules/hcp/child-adolescent.html
- ACIP Shared Clinical Decision-Making Recommendations: www.cdc.gov/vaccines/acip/acip-scdm-fags.html



U.S. Department of Health and Human Services Centers for Disease Control and Prevention Scan QR code for access to online schedule







### Table 1 Recommended Adult Immunization Schedule by Age Group, United States, 2023

Vaccine	19–26 years	27-49 years		50-64 years	≥65 years	
COVID-19	2- or 3- dose primary series and booster (See Notes)					
Influenza inactivated (IIV4) or Influenza recombinant (RIV4)_		1 dose ann	ually			
Influenza live, attenuated (LAIV4)		1 dose anr	ually			
Tetanus, diphtheria, pertussis (Tdap or Td)	1 dose			p for wound management (see n	otes)	
Measles, mumps, rubella (MMR)		1 dose Tdap, then Td or Tdap booster every 10 years  1 or 2 doses depending on indication (if born in 1957 or later)				
<b>Varicella</b> (VAR)	2 doses (if born in 1980	or later)		2 doses		
Zoster recombinant (RZV)	2 doses for immunocompron	unocompromising conditions (see notes) 2 dose			oses	
Human papillomavirus (HPV)	2 or 3 doses depending on age at initial vaccination or condition	27 through 45 years				
Pneumococcal (PCV15, PCV20, PPSV23)		1 dose PCV15 follov OR 1 dose PCV20 (	•	and the second s	See Notes See Notes	
<b>Hepatitis A</b> (HepA)		2, 3, or 4 dos	es depen	ding on vaccine		
<b>Hepatitis B</b> (HepB)		2, 3, or 4 doses dep	ending o	on vaccine or condition		
Meningococcal A, C, W, Y (MenACWY)	1 or	2 doses depending on indica	tion, see	notes for booster recommendati	ons	
Meningococcal B (MenB)	2 or 3 dos 19 through 23 years	2 or 3 doses depending on vaccine and indication, see notes for booster recommendations  19 through 23 years				
<b>Haemophilus influenzae type b</b> (Hib)		1 or 3 doses depending on indication				
Recommended vaccination for adult lack documentation of vaccination, c	· · · · · · · · · · · · · · · · · · ·	ecommended vaccination for adults v dditional risk factor or another indica		Recommended vaccination based on clinical decision-making	shared No recommendation Not applicable	



### Table 2 Recommended Adult Immunization Schedule by Medical Condition or Other Indication, United States, 2023

Vaccine	Pregnancy			Asplenia, complement deficiencies	End-stage renal disease, or on hemodialysis	Heart or lung disease; alcoholism <sup>a</sup>	Chronic liver disease	Diabetes	Health care personnel <sup>b</sup>	Men who have sex with men	
COVID-19		See Notes									
IIV4 or RIV4				1	dose annually				or		
LAIV4		Contraindica	ted			Preca	ution		1 dose a	nnually	
Tdap or Td	1 dose Tdap each pregnancy			1 dose Tdap, t	hen Td or Tdap	booster every	10 years				
MMR	Contraindicated*	Contraindicated			1 or 2	doses depend	ling on indicati	on			
VAR	Contraindicated*	Contraindicated		2 doses							
RZV		2 doses at age ≥1		2 doses at age ≥50 years							
HPV	Not Recommended*	3 doses through age	26 years	2 or 3 do	ses through ag	e 26 years dep	ending on age	at initial vac	cination or co	ndition	
Pneumococcal (PCV15, PCV20, PPSV23)					1 dose PCV1	5 followed by	PPSV23 OR 1 d	ose PCV20 (s	ee notes)		
НерА						2, 3, or 4	d <mark>oses dependi</mark> r	ng on vaccine	•		
НерВ	3 doses (see notes)			2, 3, or 4 dos	es depending	on vaccine o	condition				
MenACWY		1 or 2 doses dependin	g on indication	, see notes for	booster recom	mendations					
MenB	Precaution	2 or	2 or 3 doses depending on vaccine and indication, see notes for booster recommendations								
Hib		3 doses HSCT <sup>c</sup> recipients only									
Recommended va for adults who me age requirement, documentation of vaccination, or lac evidence of past in	eet lack f k	Recommended vaccination for adults with an additional risk factor or another indication	commended vaccination adults with an additional decision-making benefit of protection benefit of protection adults with an additional decision-making benefit of protection Contraindicated or not recommended-vaccine should not be administered.								



### Notes

### Recommended Adult Immunization Schedule for ages 19 years or older, United States, 2023

For vaccine recommendations for persons 18 years of age or younger, see the Recommended Child and Adolescent Immunization Schedule.

### **COVID-19 vaccination**

### **Routine vaccination**

- Primary series: 2-dose series at 0, 4-8 weeks (Moderna) or 2-dose series at 0, 3-8 weeks (Novavax, Pfizer-BioNTech)
- Booster dose: see www.cdc.gov/vaccines/covid-19/ clinical-considerations/interim-considerations-us.html

### **Special situations**

### Persons who are moderately or severely immunocompromised

- Primary series
- 3-dose series at 0, 4, 8 weeks (Moderna) or 3-dose series at 0, 3, 7 weeks (Pfizer-BioNTech)
- 2-dose series at 0, 3 weeks (Novavax)
- Booster dose: see www.cdc.gov/vaccines/covid-19/ clinical-considerations/interim-considerations-us.html
- Pre-exposure prophylaxis (e.g., monoclonal antibodies) may be considered to complement COVID-19 vaccination. See www.cdc.gov/ vaccines/covid-19/clinical-considerations/interimconsiderations-us.html#immunocompromised

For Janssen COVID-19 Vaccine recipients see COVID-19 schedule at www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html.

Note: Current COVID-19 schedule available at www. cdc.gov/vaccines/covid-19/downloads/COVID-19-immunization-schedule-ages-6months-older.pdf. For more information on Emergency Use Authorization (EUA) indications for COVID-19 vaccines, please visit www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/covid-19-vaccines

### Haemophilus influenzae type b vaccination

### Special situations

- Anatomical or functional asplenia (including sickle cell disease): 1 dose if previously did not receive Hib; if elective splenectomy, 1 dose preferably at least 14 days before splenectomy
- Hematopoietic stem cell transplant (HSCT):
   3-dose series 4 weeks apart starting 6–12 months after successful transplant, regardless of Hib vaccination history

### **Hepatitis A vaccination**

### **Routine vaccination**

• Not at risk but want protection from hepatitis A (identification of risk factor not required): 2-dose series HepA (Havrix 6–12 months apart or Vaqta 6–18 months apart [minimum interval: 6 months]) or 3-dose series HepA-HepB (Twinrix at 0, 1, 6 months [minimum intervals: dose 1 to dose 2: 4 weeks / dose 2 to dose 3: 5 months])

### **Special situations**

- At risk for hepatitis A virus infection: 2-dose series HepA or 3-dose series HepA-HepB as above
- Chronic liver disease (e.g., persons with hepatitis B, hepatitis C, cirrhosis, fatty liver disease, alcoholic liver disease, autoimmune hepatitis, alanine aminotransferase [ALT] or aspartate aminotransferase [AST] level greater than twice the upper limit of normal)
- HIV infection
- Men who have sex with men
- Injection or noninjection drug use
- Persons experiencing homelessness
- -Work with hepatitis A virus in research laboratory or with nonhuman primates with hepatitis A virus infection

- -Travel in countries with high or intermediate endemic hepatitis A (HepA-HepB [Twinrix] may be administered on an accelerated schedule of 3 doses at 0, 7, and 21–30 days, followed by a booster dose at 12 months)
- Close, personal contact with international adoptee (e.g., household or regular babysitting) in first 60 days after arrival from country with high or intermediate endemic hepatitis A (administer dose 1 as soon as adoption is planned, at least 2 weeks before adoptee's arrival)
- **Pregnancy** if at risk for infection or severe outcome from infection during pregnancy
- Settings for exposure, including health care settings targeting services to injection or noninjection drug users or group homes and nonresidential day care facilities for developmentally disabled persons (individual risk factor screening not required)

### Hepatitis B vaccination

### **Routine vaccination**

- Age 19 through 59 years: complete a 2- or 3- or 4-dose series
- 2-dose series only applies when 2 doses of Heplisav-B\* are used at least 4 weeks apart
- 3-dose series Engerix-B, PreHevbrio\*, or Recombivax HB at 0, 1, 6 months [minimum intervals: dose 1 to dose 2: 4 weeks / dose 2 to dose 3: 8 weeks / dose 1 to dose 3: 16 weeks])
- 3-dose series HepA-HepB (Twinrix at 0, 1, 6 months [minimum intervals: dose 1 to dose 2: 4 weeks / dose 2 to dose 3: 5 months])
- -4-dose series HepA-HepB (Twinrix) accelerated schedule of 3 doses at 0, 7, and 21–30 days, followed by a booster dose at 12 months
- \*Note: Heplisav-B and PreHevbrio are not recommended in pregnancy due to lack of safety data in pregnant persons.



### Notes

### Recommended Adult Immunization Schedule, United States, 2023

- Age 60 years or older with known risk factors for hepatitis B virus infection should complete a HepB vaccine series.
- Age 60 years or older without known risk factors for hepatitis B virus infection may complete a HepB vaccine series.
- Risk factors for hepatitis B virus infection include:
- Chronic liver disease (e.g., persons with hepatitis C, cirrhosis, fatty liver disease, alcoholic liver disease, autoimmune hepatitis, alanine aminotransferase [ALT] or aspartate aminotransferase [AST] level greater than twice upper limit of normal)
- · HIV infection
- Sexual exposure risk (e.g., sex partners of hepatitis B surface antigen [HBsAg]-positive persons; sexually active persons not in mutually monogamous relationships; persons seeking evaluation or treatment for a sexually transmitted infection; men who have sex with men)
- Current or recent injection drug use
- Percutaneous or mucosal risk for exposure to blood (e.g., household contacts of HBsAgpositive persons; residents and staff of facilities for developmentally disabled persons; health care and public safety personnel with reasonably anticipated risk for exposure to blood or blood-contaminated body fluids; persons on maintenance dialysis, including in-center or home hemodialysis and peritoneal dialysis, and persons who are predialysis; patients with diabetes)
- Incarceration
- Travel in countries with high or intermediate endemic hepatitis B

### Special situations

- Patients on dialysis: complete a 3- or 4-dose series
- 3-dose series Recombivax HB at 0, 1, 6 months (note: use Dialysis Formulation 1 mL = 40 mcg)
- -4-dose series Engerix-B at 0, 1, 2, and 6 months (note: use 2 mL dose instead of the normal adult dose of 1 mL)

### **Human papillomavirus vaccination**

### **Routine vaccination**

- HPV vaccination recommended for all persons through age 26 years: 2- or 3-dose series depending on age at initial vaccination or condition:
- Age 15 years or older at initial vaccination:
   3-dose series at 0, 1-2 months, 6 months (minimum intervals: dose 1 to dose 2: 4 weeks / dose 2 to dose 3: 12 weeks / dose 1 to dose 3: 5 months; repeat dose if administered too soon)
- Age 9–14 years at initial vaccination and received
   1 dose or 2 doses less than 5 months apart:
   1 additional dose
- Age 9-14 years at initial vaccination and received 2 doses at least 5 months apart: HPV vaccination series complete, no additional dose needed
- Interrupted schedules: If vaccination schedule is interrupted, the series does not need to be restarted
- No additional dose recommended when any HPV vaccine series has been completed using the recommended dosing intervals.

### Shared clinical decision-making

 Some adults age 27-45 years: Based on shared clinical decision-making, 2- or 3-dose series as above

### **Special situations**

- Age ranges recommended above for routine and catch-up vaccination or shared clinical decisionmaking also apply in special situations
- Immunocompromising conditions, including HIV infection: 3-dose series, even for those who initiate vaccination at age 9 through 14 years.
- Pregnancy: Pregnancy testing is not needed before vaccination; HPV vaccination is not recommended until after pregnancy; no intervention needed if inadvertently vaccinated while pregnant

### Influenza vaccination

### Routine vaccination

- Age 19 years or older: 1 dose any influenza vaccine appropriate for age and health status annually.
- Age 65 years or older: Any one of quadrivalent high-dose inactivated influenza vaccine (HD-IIV4), quadrivalent recombinant influenza vaccine (RIV4), or quadrivalent adjuvanted inactivated influenza vaccine (alIV4) is preferred. If none of these three vaccines is available, then any other age-appropriate influenza vaccine should be used.
- For the 2022–2023 season, see www.cdc.gov/mmwr/ volumes/71/rr/rr7101a1.htm
- For the 2023–2024 season, see the 2023–2024 ACIP influenza vaccine recommendations.

### **Special situations**

- **Egg allergy, hives only:** any influenza vaccine appropriate for age and health status annually
- Egg allergy–any symptom 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 or LAIV4, administer in medical setting under supervision of health care provider who can recognize and manage severe allergic reactions.
- Close contacts (e.g., caregivers, healthcare workers) of severely immunosuppressed persons who require a protected environment: these persons should not receive LAIV4. If LAIV4 is given, they should avoid contact with/caring for such immunosuppressed persons for 7 days after vaccination.
- Severe allergic reaction (e.g., anaphylaxis) to a vaccine component or a previous dose of any influenza vaccine: see Appendix listing contraindications and precautions



### Notes

### Recommended Adult Immunization Schedule, United States, 2023

 History of Guillain-Barré syndrome within 6 weeks after previous dose of influenza vaccine: Generally, should not be vaccinated unless vaccination benefits outweigh risks for those at higher risk for severe complications from influenza

### Measles, mumps, and rubella vaccination

#### **Routine vaccination**

- No evidence of immunity to measles, mumps, or rubella: 1 dose
- Evidence of immunity: Born before 1957 (health care personnel, see below), documentation of receipt of MMR vaccine, laboratory evidence of immunity or disease (diagnosis of disease without laboratory confirmation is not evidence of immunity)

### **Special situations**

- Pregnancy with no evidence of immunity to rubella: MMR contraindicated during pregnancy; after pregnancy (before discharge from health care facility), 1 dose
- Nonpregnant persons of childbearing age with no evidence of immunity to rubella: 1 dose
- HIV infection with CD4 percentages ≥15% and CD4 count ≥200 cells/mm³ for at least 6 months and no evidence of immunity to measles, mumps, or rubella: 2-dose series at least 4 weeks apart; MMR contraindicated for HIV infection with CD4 percentage <15% or CD4 count <200 cells/mm³
- Severe immunocompromising conditions:
   MMR contraindicated
- Students in postsecondary educational institutions, international travelers, and household or close, personal contacts of immunocompromised persons with no evidence of immunity to measles, mumps, or rubella:
   2-dose series at least 4 weeks apart if previously did
- 2-dose series at least 4 weeks apart if previously did not receive any doses of MMR or 1 dose if previously received 1 dose MMR

- In mumps outbreak settings, for information about additional doses of MMR (including 3rd dose of MMR), see www.cdc.gov/mmwr/volumes/67/wr/mm6701a7. htm
- Health care personnel:
- Born before 1957 with no evidence of immunity to measles, mumps, or rubella:
   Consider 2-dose series at least 4 weeks apart for

protection against measles or mumps or 1 dose for protection against rubella

Born in 1957 or later with no evidence of immunity to measles, mumps, or rubella:
 2-dose series at least 4 weeks apart for protection against measles or mumps or at least 1 dose for protection against rubella

### **Meningococcal vaccination**

### Special situations for MenACWY

- Anatomical or functional asplenia (including sickle cell disease), HIV infection, persistent complement component deficiency, complement inhibitor (e.g., eculizumab, ravulizumab) use: 2-dose series MenACWY-D (Menactra, Menveo, or MenQuadfi) at least 8 weeks apart and revaccinate every 5 years if risk remains
- Travel in countries with hyperendemic or epidemic meningococcal disease, or microbiologists routinely exposed to Neisseria meningitidis: 1 dose MenACWY (Menactra, Menveo, or MenQuadfi) and revaccinate every 5 years if risk remains
- First-year college students who live in residential housing (if not previously vaccinated at age 16 years or older) or military recruits: 1 dose MenACWY (Menactra, Menveo, or MenQuadfi)
- For MenACWY booster dose recommendations for groups listed under "Special situations" and in an outbreak setting (e.g., in community or organizational settings and among men who have sex with men) and additional meningococcal vaccination information, see www.cdc.gov/mmwr/volumes/69/rr/rr6909a1.htm

### Shared clinical decision-making for MenB

• Adolescents and young adults age 16–23 years (age 16–18 years preferred) not at increased risk for meningococcal disease: Based on shared clinical decision-making, 2-dose series MenB-4C (Bexsero) at least 1 month apart or 2-dose series MenB-FHbp (Trumenba) at 0, 6 months (if dose 2 was administered less than 6 months after dose 1, administer dose 3 at least 4 months after dose 2); MenB-4C and MenB-FHbp are not interchangeable (use same product for all doses in series)

### Special situations for MenB

- Anatomical or functional asplenia (including sickle cell disease), persistent complement component deficiency, complement inhibitor (e.g., eculizumab, ravulizumab) use, or microbiologists routinely exposed to Neisseria meningitidis:
- 2-dose primary series MenB-4C (Bexsero) at least 1 month apart or 3-dose primary series MenB-FHbp (Trumenba) at 0, 1–2, 6 months (if dose 2 was administered at least 6 months after dose 1, dose 3 not needed; if dose 3 is administered earlier than 4 months after dose 2, a fourth dose should be administered at least 4 months after dose 3); MenB-4C and MenB-FHbp are not interchangeable (use same product for all doses in series); 1 dose MenB booster 1 year after primary series and revaccinate every 2–3 years if risk remains
- Pregnancy: Delay MenB until after pregnancy unless at increased risk and vaccination benefits outweigh potential risks
- For MenB booster dose recommendations for groups listed under "Special situations" and in an outbreak setting (e.g., in community or organizational settings and among men who have sex with men) and additional meningococcal vaccination information, see www.cdc.qov/mmwr/volumes/69/rr/rr6909a1.htm

**Note:** MenB vaccines may be administered simultaneously with MenACWY vaccines if indicated, but at a different anatomic site, if feasible.



### Recommended Adult Immunization Schedule, United States, 2023

### Pneumococcal vaccination

### Routine vaccination

- Age 65 years or older who have:
- Not previously received a dose of PCV13, PCV15, or PCV20 or whose previous vaccination history is unknown: 1 dose PCV15 OR 1 dose PCV20. If PCV15 is used, this should be followed by a dose of PPSV23 given at least 1 year after the PCV15 dose. A minimum interval of 8 weeks between PCV15 and PPSV23 can be considered for adults with an immunocompromising condition,\* cochlear implant, or cerebrospinal fluid leak to minimize the risk of invasive pneumococcal disease caused by serotypes unique to PPSV23 in these vulnerable groups.
- Previously received only PCV7: follow the recommendation above.
- Previously received only PCV13: 1 dose PCV20 at least 1 year after the PCV13 dose OR complete the recommended PPSV23 series as described here www.cdc.gov/vaccines/vpd/pneumo/downloads/pneumo-vaccine-timing.pdf.
- Previously received only PPSV23: 1 dose PCV15 OR 1 dose PCV20 at least 1 year after the PPSV23 dose.
   If PCV15 is used, it need not be followed by another dose of PPSV23.
- Previously received both PCV13 and PPSV23 but NO PPSV23 was received at age 65 years or older: 1 dose PCV20 at least 5 years after their last pneumococcal vaccine dose OR complete the recommended PPSV23 series as described here www.cdc.gov/vaccines/vpd/pneumo/downloads/pneumo-vaccine-timing.pdf.
- Previously received both PCV13 and PPSV23, AND PPSV23 was received at age 65 years or older:
   Based on shared clinical decision-making, 1 dose of PCV20 at least 5 years after the last pneumococcal vaccine dose.

 For guidance on determining which pneumococcal vaccines a patient needs and when, please refer to the mobile app which can be downloaded here: www.cdc. gov/vaccines/vpd/pneumo/hcp/pneumoapp.html

### Special situations

- Age 19–64 years with certain underlying medical conditions or other risk factors\*\* who have
- Not previously received a PCV13, PCV15, or PCV20 or whose previous vaccination history is unknown: 1 dose PCV15 OR 1 dose PCV20. If PCV15 is used, this should be followed by a dose of PPSV23 given at least 1 year after the PCV15 dose. A minimum interval of 8 weeks between PCV15 and PPSV23 can be considered for adults with an immunocompromising condition,\* cochlear implant, or cerebrospinal fluid leak
- **Previously received only PCV7:** follow the recommendation above.
- Previously received only PCV13: 1 dose PCV20 at least 1 year after the PCV13 dose OR complete the recommended PPSV23 series as described here www.cdc.gov/vaccines/vpd/pneumo/downloads/ pneumo-vaccine-timing.pdf.
- Previously received only PPSV23: 1 dose PCV15 OR 1 dose PCV20 at least 1 year after the PPSV23 dose.
   If PCV15 is used, it need not be followed by another dose of PPSV23.
- Previously received both PCV13 and PPSV23 but have not completed the recommended series: 1 dose PCV20 at least 5 years after their last pneumococcal vaccine dose OR complete the recommended PPSV23 series as described here www.cdc.gov/vaccines/vpd/pneumo/downloads/pneumo-vaccine-timing.pdf.
- For guidance on determining which pneumococcal vaccines a patient needs and when, please refer to the mobile app which can be downloaded here: www.cdc. gov/vaccines/vpd/pneumo/hcp/pneumoapp.html

- \*Note: Immunocompromising conditions include chronic renal failure, nephrotic syndrome, immunodeficiency, iatrogenic immunosuppression, generalized malignancy, human immunodeficiency virus, Hodgkin disease, leukemia, lymphoma, multiple myeloma, solid organ transplants, congenital or acquired asplenia, sickle cell disease, or other hemoglobinopathies.
- \*\*Note: Underlying medical conditions or other risk factors include alcoholism, chronic heart/liver/ lung disease, chronic renal failure, cigarette smoking, cochlear implant, congenital or acquired asplenia, CSF leak, diabetes mellitus, generalized malignancy, HIV, Hodgkin disease, immunodeficiency, iatrogenic immunosuppression, leukemia, lymphoma, multiple myeloma, nephrotic syndrome, solid organ transplants, or sickle cell disease or other hemoglobinopathies.

### Polio vaccination

#### Routine vaccination

Routine poliovirus vaccination of adults residing in the United States is not necessary.

### **Special situations**

- Adults at increased risk of exposure to poliovirus with:
- No evidence of a complete polio vaccination series (i.e., at least 3 doses): administer remaining doses (1, 2, or 3 doses) to complete a 3-dose series
- Evidence of completed polio vaccination series (i.e., at least 3 doses): may administer one lifetime IPV booster

For detailed information, see: www.cdc.gov/vaccines/vpd/polio/hcp/recommendations.html



### Tetanus, diphtheria, and pertussis vaccination

### Routine vaccination

• Previously did not receive Tdap at or after age 11 years: 1 dose Tdap, then Td or Tdap every 10 years

### **Special situations**

- Previously did not receive primary vaccination series for tetanus, diphtheria, or pertussis: 1 dose Tdap followed by 1 dose Td or Tdap at least 4 weeks later, and a third dose of Td or Tdap 6–12 months later (Tdap can be substituted for any Td dose, but preferred as first dose), Td or Tdap every 10 years thereafter
- Pregnancy: 1 dose Tdap during each pregnancy, preferably in early part of gestational weeks 27–36
- Wound management: Persons with 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 who have not previously received Tdap or whose Tdap history is unknown. If a tetanus-toxoid-containing vaccine is indicated for a pregnant woman, use Tdap. For detailed information, see www.cdc.gov/mmwr/volumes/69/wr/mm6903a5.htm

### Varicella vaccination

### Routine vaccination

- No evidence of immunity to varicella: 2-dose series 4-8 weeks apart if previously did not receive varicellacontaining vaccine (VAR or MMRV [measles-mumpsrubella-varicella vaccine] for children); if previously received 1 dose varicella-containing vaccine, 1 dose at least 4 weeks after first dose
- Evidence of immunity: U.S.-born before 1980 (except for pregnant persons and health care personnel [see below]), documentation of 2 doses varicella-containing vaccine at least 4 weeks apart, diagnosis or verification of history of varicella or herpes zoster by a health care provider, laboratory evidence of immunity or disease

### Special situations

- Pregnancy with no evidence of immunity to varicella: VAR contraindicated during pregnancy; after pregnancy (before discharge from health care facility), 1 dose if previously received 1 dose varicellacontaining vaccine or dose 1 of 2-dose series (dose 2: 4–8 weeks later) if previously did not receive any varicella-containing vaccine, regardless of whether U.S.-born before 1980
- Health care personnel with no evidence of immunity to varicella: 1 dose if previously received 1 dose varicella-containing vaccine; 2-dose series 4-8 weeks apart if previously did not receive any varicella-containing vaccine, regardless of whether U.S.-born before 1980
- HIV infection with CD4 percentages ≥15% and CD4 count ≥200 cells/mm³ with no evidence of immunity: Vaccination may be considered (2 doses 3 months apart); VAR contraindicated for HIV infection with CD4 percentage <15% or CD4 count <200 cells/mm³</li>
- Severe immunocompromising conditions:
   VAR contraindicated

### Zoster vaccination

### Routine vaccination

- Age 50 years or older\*: 2-dose series recombinant zoster vaccine (RZV, Shingrix) 2–6 months apart (minimum interval: 4 weeks; repeat dose if administered too soon), regardless of previous herpes zoster or history of zoster vaccine live (ZVL, Zostavax) vaccination.
- \*Note: Serologic evidence of prior varicella is not necessary for zoster vaccination. However, if serologic evidence of varicella susceptibility becomes available, providers should follow ACIP guidelines for varicella vaccination first. RZV is not indicated for the prevention of varicella, and there are limited data on the use of RZV in persons without a history of varicella or varicella vaccination.

### **Special situations**

- Pregnancy: There is currently no ACIP recommendation for RZV use in pregnancy. Consider delaying RZV until after pregnancy.
- Immunocompromising conditions (including persons with HIV regardless of CD4 count)\*\*:
   2-dose series recombinant zoster vaccine (RZV, Shingrix) 2–6 months apart (minimum interval: 4 weeks; repeat dose if administered too soon).
   For detailed information, see www.cdc.gov/shingles/vaccination/immunocompromised-adults.html
- \*\*Note: If there is no documented history of varicella, varicella vaccination, or herpes zoster, providers should refer to the clinical considerations for use of RZV in immunocompromised adults aged ≥19 years and the ACIP varicella vaccine recommendations for further guidance: www.cdc.gov/mmwr/volumes/71/wr/mm7103a2.htm



### **Appendix**

### Recommended Adult Immunization Schedule, United States, 2023

### **Guide to Contraindications and Precautions to Commonly Used Vaccines**

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

### For COVID-19 vaccine contraindications and precautions see

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

Vaccine	Contraindicated or Not Recommended <sup>1</sup>	Precautions <sup>2</sup>
Influenza, egg-based, inactivated injectable (IIV4)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after previous dose of any influenza vaccine (i.e., any egg-based IIV, ccIIV, RIV, or LAIV of any valency)</li> <li>Severe allergic reaction (e.g., anaphylaxis) to any vaccine component<sup>3</sup> (excluding egg)</li> </ul>	<ul> <li>Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine</li> <li>Moderate or severe acute illness with or without fever</li> </ul>
Influenza, cell culture-based inactivated injectable [(ccIIV4), Flucelvax® Quadrivalent]	Severe allergic reaction (e.g., anaphylaxis) to any ccIIV of any valency, or to any component <sup>3</sup> of ccIIV4	<ul> <li>Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine</li> <li>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 setting under supervision of health care provider who can recognize and manage severe allergic reactions. May consult an allergist.</li> <li>Moderate or severe acute illness with or without fever</li> </ul>
Influenza, recombinant injectable [(RIV4), Flublok° Quadrivalent]	Severe allergic reaction (e.g., anaphylaxis) to any RIV of any valency, or to any component <sup>3</sup> of RIV4	<ul> <li>Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine</li> <li>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.</li> <li>Moderate or severe acute illness with or without fever</li> </ul>
Influenza, live attenuated [LAIV4, Flumist* Quadrivalent]	Severe allergic reaction (e.g., anaphylaxis) after previous dose of any influenza vaccine (i.e., any egg-based IIV, ccIIV, RIV, or LAIV of any valency) Severe allergic reaction (e.g., anaphylaxis) to any vaccine component³ (excluding egg) Anatomic or functional asplenia Immunocompromised due to any cause including, but not limited to, medications and HIV infection Close contacts or caregivers of severely immunosuppressed persons who require a protected environment Pregnancy Cochlear implant Active communication between the cerebrospinal fluid (CSF) and the oropharynx, nasopharynx, nose, ear, or any other cranial CSF leak Received influenza antiviral medications oseltamivir or zanamivir within the previous 48 hours, peramivir within the previous 5 days, or baloxavir within the previous 17 days.	<ul> <li>Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine</li> <li>Asthma in persons aged 5 years old or older</li> <li>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)]</li> <li>Moderate or severe acute illness with or without fever</li> </ul>

- 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.



### **Appendix**

### Recommended Adult Immunization Schedule, United States, 2023

Vaccine	Contraindicated or Not Recommended <sup>1</sup>	Precautions <sup>2</sup>
Haemophilus influenzae type b (Hib)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup></li> <li>For Hiberix, ActHib, and PedvaxHIB only: History of severe allergic reaction to dry natural latex</li> </ul>	Moderate or severe acute illness with or without fever
Hepatitis A (HepA)	$\bullet$ Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component $^3$ including neomycin	Moderate or severe acute illness with or without fever
Hepatitis B (HepB)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup> including yeast</li> <li>Pregnancy: Heplisav-B and PreHevbrio are not recommended due to lack of safety data in pregnant persons.</li> <li>Use other hepatitis B vaccines if HepB is indicated<sup>4</sup></li> </ul>	Moderate or severe acute illness with or without fever
Hepatitis A- Hepatitis B vaccine [HepA-HepB, (Twinrix®)]	$\bullet$ Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component $^3$ including neomycin and yeast	Moderate or severe acute illness with or without fever
Human papillomavirus (HPV)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup></li> <li>Pregnancy: HPV vaccination not recommended</li> </ul>	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 <sup>3</sup> Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of chemotherapy, 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	<ul> <li>Recent (≤11 months) receipt of antibody-containing blood product (specific interval depends on product)</li> <li>History of thrombocytopenia or thrombocytopenic purpura</li> <li>Need for tuberculin skin testing or interferon-gamma release assay (IGRA) testing</li> <li>Moderate or severe acute illness with or without fever</li> </ul>
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 component <sup>3</sup> For MenACWY-D and MenACWY-CRM only: severe allergic reaction to any diphtheria toxoid–or CRM197–containing vaccine     For MenACWY-TT only: severe allergic reaction to a tetanus toxoid-containing vaccine	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 <sup>3</sup>	Pregnancy For MenB-4C only: Latex sensitivity Moderate or severe acute illness with or without fever
Pneumococcal conjugate (PCV15, PCV20)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup></li> <li>Severe allergic reaction (e.g., anaphylaxis) to any diphtheria-toxoid—containing vaccine or to its vaccine component<sup>3</sup></li> </ul>	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 <sup>3</sup>	Moderate or severe acute illness with or without fever
Tetanus, diphtheria, and acellular pertussis (Tdap) Tetanus, diphtheria (Td)	<ul> <li>Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component<sup>3</sup></li> <li>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</li> </ul>	<ul> <li>Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of tetanus-toxoid-containing vaccine</li> <li>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</li> <li>Moderate or severe acute illness with or without fever</li> <li>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</li> </ul>
Varicella (VAR)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component <sup>3</sup> Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of chemotherapy, 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	<ul> <li>Recent (≤11 months) receipt of antibody-containing blood product (specific interval depends on product)</li> <li>Receipt of specific antiviral drugs (acyclovir, famciclovir, or valacyclovir) 24 hours before vaccination (avoid use of these antiviral drugs for 14 days after vaccination)</li> <li>Use of aspirin or aspirin-containing products</li> <li>Moderate or severe acute illness with or without fever</li> </ul>
Zoster recombinant vaccine (RZV)	• Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component <sup>3</sup>	Moderate or severe acute illness with or without fever     Current herpes zoster infection

- 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.
- 4. For information on the pregnancy exposure registries for persons who were inadvertently vaccinated with Heplisav-B or PreHevbrio while pregnant, please visit heplisavbpregnancyregistry.com/ or www.prehevbrio.com/#safety.

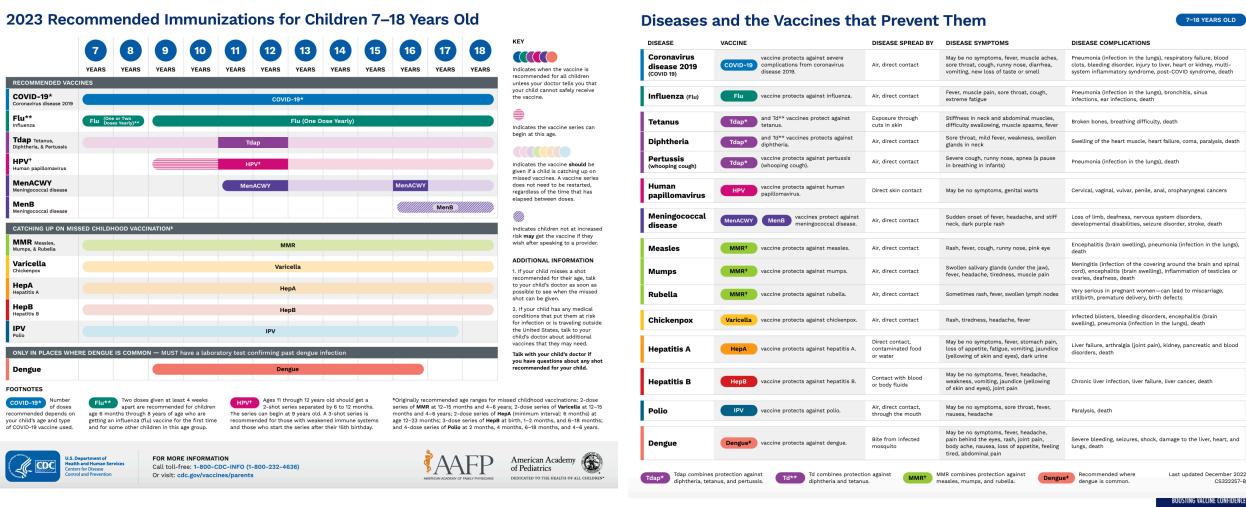


## Resources: "Parent Friendly Schedules"

#### 2023 Recommended Immunizations for Children from Birth Through 6 Years Old 4-6 VACCINE HepB RV\* DTaP Diphtheria, Hib\* Haemophilus influenzae type b PCV13, PCV15 COVID-19\*\* Flu† Flu (One or Two Doses Yearly) MMR Measles Varicella HepA<sup>‡</sup> FOOTNOTES ADDITIONAL INFORMATION Two doses of Hep A vaccine 1. If your child misses 2. If your child has any medical of doses least 4 weeks apart are needed for lasting conditions that put them at child's doctor if a shot recommended Administering a third dose are recommended for children protection. The 2 doses should be given risk for infection (e.g., sickle cell, recommended depends on for their age, talk to you have question: your child's age and type at age 6 months depends age 6 months through 8 years of between age 12 and 23 months. Both on the brand of Hib or of COVID-19 vaccine used. age who are getting an influenza doses should be separated by at least soon as possible to or is traveling outside the United rotavirus vaccine used for (flu) vaccine for the first time 6 months. Children 2 years and older see when the missed States, talk to your child's doctor previous dose. and for some other children in who have not received 2 doses of Hep A shot can be given. about additional vaccines that American Academy Call toll-free: 1-800-CDC-INFO (1-800-232-4636) of Pediatrics Or visit: cdc.gov/vaccines/parents

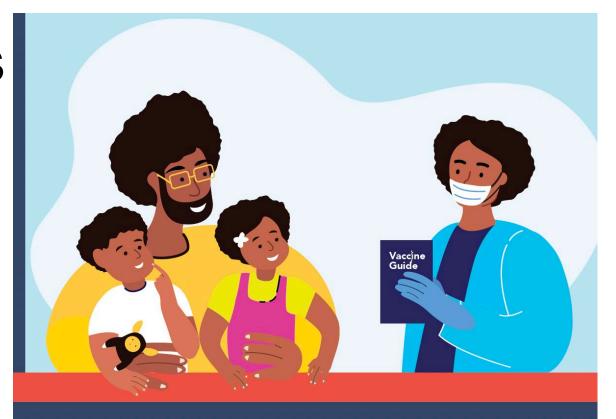
DISEASE	VACCINE	DISEASE SPREAD BY	DISEASE SYMPTOMS	DISEASE COMPLICATIONS
Hepatitis B	HepB vaccine protects against hepatitis B.	Contact with blood or body fluids	May be no symptoms, fever, headache, weakness, vomiting, jaundice (yellowing of skin and eyes), joint pain	Chronic liver infection, liver failure, liver cancer, death
Rotavirus	RV vaccine protects against rotavirus.	Through the mouth	Diarrhea, fever, vomiting	Severe diarrhea, dehydration, death
Diphtheria	DTaP* vaccine protects against diphtheria.	Air, direct contact	Sore throat, mild fever, weakness, swollen glands in neck	Swelling of the heart muscle, heart failure, coma, paralysis, death
Pertussis (whooping cough)	DTaP* vaccine protects against pertussis (whooping cough).	Air, direct contact	Severe cough, runny nose, apnea (a pause in breathing in infants)	Pneumonia (infection in the lungs), death
Tetanus	DTaP* vaccine protects against tetanus.	Exposure through cuts in skin	Stiffness in neck and abdominal muscles, difficulty swallowing, muscle spasms, fever	Broken bones, breathing difficulty, death
Haemophilus influenzae type b (Hib)	Hib vaccine protects against Haemophilus influenzae type b.	Air, direct contact	May be no symptoms unless bacteria enter the blood	Meningitis (infection of the covering around the brain and spinal cord), intellectual disability, epiglottitis (life-threatening infection that can block the windpipe and lead to serious breathing problems), pneumonia (infection in the lungs), death
Pneumococcal disease (PCV13, PCV15)	PCV vaccine protects against pneumococcal disease.	Air, direct contact	May be no symptoms, pneumonia (infection in the lungs)	Bacteremia (blood infection), meningitis (infection of the covering around the brain and spinal cord), death
Polio	IPV vaccine protects against polio.	Air, direct contact, through the mouth	May be no symptoms, sore throat, fever, nausea, headache	Paralysis, death
Coronavirus disease 2019 (COVID-19)	vaccine protects against severe complications from coronavirus disease 2019.	Air, direct contact	May be no symptoms, fever, muscle aches, sore throat, cough, runny nose, diarrhea, vomiting, new loss of taste or smell	Pneumonia (infection in the lungs), respiratory failure, blood clots, bleeding disorder, injury to liver, heart or kidney, multisystem inflammatory syndrome, post-COVID syndrome, death
Influenza (Flu)	Flu vaccine protects against influenza.	Air, direct contact	Fever, muscle pain, sore throat, cough, extreme fatigue	Pneumonia (infection in the lungs), bronchitis, sinus infections ear infections, death
Measles	MMR** vaccine protects against measles.	Air, direct contact	Rash, fever, cough, runny nose, pink eye	Encephalitis (brain swelling), pneumonia (infection in the lungs), death
Mumps	MMR** vaccine protects against mumps.	Air, direct contact	Swollen salivary glands (under the jaw), fever, headache, tiredness, muscle pain	Meningitis (infection of the covering around the brain and spinal cord), encephalitis (brain swelling), inflammation of testicles or ovaries, deafness, death
Rubella	MMR** vaccine protects against rubella.	Air, direct contact	Sometimes rash, fever, swollen lymph nodes	Very serious in pregnant women—can lead to miscarriage, stillbirth, premature delivery, birth defects
Chickenpox	Varicella vaccine protects against chickenpox.	Air, direct contact	Rash, tiredness, headache, fever	Infected blisters, bleeding disorders, encephalitis (brain swelling), pneumonia (infection in the lungs), death
Hepatitis A	HepA vaccine protects against hepatitis A.	Direct contact, contaminated food	May be no symptoms, fever, stomach pain, loss of appetite, fatigue, vomiting, jaundice (yellowing of skin and eyes), dark urine	Liver failure, arthralgia (joint pain), kidney, pancreatic and blood disorders, death

## Resources: "Parent Friendly Schedules"



## **★ICLAC** Resources

- Parents Guide to Recommended & Required Vaccinations
- Child Vaccination Booklet
- Available in multiple languages: English, Spanish, Chinese



## PARENTS GUIDE to Recommended and Required Vaccinations



## Additional Resources

CDC Vaccine Schedules App for Healthcare Providers



Download "CDC Vaccine Schedules" free for iOS and Android devices.

Download the app



Materials for You and Your Practice

Help strengthen communication between you and parents.



Materials to Share With Parents

Help parents understand vaccine benefits and risks.



## **QUESTIONS?**

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