Michigan Immunization Update

November 2023



Agenda



- Winter is coming (*winter is here*)
 - COVID Commercialization/Bridge Access Program
 - RSV
- State Immunization Coverage Rates
- Challenges
- Strategies to improve rates
- Vaccine Champions



Division of Immunization Priorities



- Ensure equitable access to all vaccines
 - Vaccine deserts
 - Additional data
- Improve routine immunization coverage
 - Best and promising practices

Evaluation and Impact Assessment

- Vaccine champions
- Retain VFC Providers
- Flu, COVID, RSV
 - Education/Training • <u>https://www.michigan.gov/mdhhs/keep-mi-</u> <u>healthy/chronicdiseases/seasonal-respiratory-viruses</u>







Influenza (flu)

Respiratory Syncytial Virus (RSV)

COVID-19



COVID Commercialization and Bridge Access Program



Key Points



- Commercialization transition from vaccine supplied by the government to being supplied by manufacturers for profit.
- MDHHS **ONLY** has access to public vaccine stock through:
 - Bridge Access Program
 - VFC
- MI-AVP and VFC providers can order vaccine through MDHHS.
 - Normal process through MCIR e-ordering.
- COVID-19 vaccines are covered by private/public insurers, and provided at no cost to uninsured/underinsured adults through the Bridge Access Program.
- Pharmacies have a separate contract with the federal government to provide vaccines to uninsured.

Ordering Controls



- COVID-19 vaccines are becoming available by manufacturer over time rather than all at once.
- Vaccines will continue to vary in distribution:
 - Moderna and Novavax will be centrally distributed.
 - Pfizer will ship directly from the manufacturer.
- Controlled ordering (allocations) helps ensure that COVID-19 vaccines are available to all awardees to support their programmatic plans.
- MDHHS and CDC will regularly monitor allocations and orders in VTrckS.
- As additional COVID-19 vaccines become available, CDC will make updates and adjustments to allocations to align with the ordering activity of awardees.

What is the Role of MDHHS?



Provider Enrollment

- Leverage existing providers enrolled in the awardee's Adult/317 Program and/or enroll new providers.
- Enroll additional providers in areas with low access to vaccination sites, low COVID-19 vaccination coverage, and/or high uninsured rates or providers serving persons in carceral settings.
- Work to ensure that approximately 25% of Bridge Access Program COVID-19 vaccines are distributed through HRSA-supported health centers.
 - All HRSA-supported health centers are eligible to enroll in the awardee's existing Adult/317 Program.
- Coordinate community outreach.

Bridge Program



MI-AVP Providers

- LHDs
- FQHCs
- Migrant, tribal, and rural health centers

Pharmacies

- Walgreens
- CVS
- eTrueNorth

To sign up, please visit: www.joinetruenorth.com





COVID Vaccine Provider Deserts



Vaccine desert More than a 15 minute drive to the closest active COVID-19 vaccination site.

OO Potential vaccination sites within deserts To view, see "Find potential new sites" on the menu.



- Work with eTrueNorth to identify independent pharmacies and encourage them to enroll.
- Identify opportunities for community outreach.
- Work with vaccine champions.

RSV Vaccines



RSV Protection this Season



New Immunizations to Protect Against Severe RSV

| | Who Does It Protect? | Type of Product | Is It for Everyone in Group? | |
|---------------|-------------------------|---------------------------------------|--|--|
| B | Adults 60 and over | RSV vaccine | Talk to your doctor first | |
| | Babies | RSV antibody given to baby | All infants entering or born during RSV season. Small group of older babies for second season. | |
| E Contraction | Babies | RSV vaccine given during pregnancy | Can get if you are 32–36 weeks pregnant during September–January | |
| www | v.cdc.gov/rsv | | | |

RSV Monoclonal Antibody



- August ACIP met to discuss nirsevimab and voted unanimously on the inclusion of the new monoclonal on the VFC contract; it is recommended for:
 - Infants younger than 8 months in their first RSV season.
 - Children aged 8 months to 19 months who are at higher risk for contracting RSV.
- Challenges:
 - Limited supply
 - Cost
 - Timing
 - Administration considerations
 - Birthing facility vs pediatric office
 - Recording inventory and administrations in MCIR

MDHHS Prioritization



- Limited allocations of nirsevimab.
- MDHHS implements CDC recommendations for healthcare providers on use of nirsevimab during limited allocations:
 - For infants weighing <5 kg, ACIP recommendations are unchanged.
 - For infants weighing ≥5 kg, prioritize using 100 mg nirsevimab doses in infants at highest risk of severe RSV disease:
 - Young infants aged <6 months.
 - American Indian and Alaska Native infants aged <8 months.
 - Infants aged 6 through 7 months with conditions that place them at high risk of severe RSV disease.
 - In palivizumab-eligible children aged 8–19 months, suspend using nirsevimab for the 2023–2024 RSV season. These children should receive palivizumab per <u>AAP recommendations</u>.
 - Continue offering nirsevimab to American Indian and Alaska Native children aged 8–19 months who are not palivizumab-eligible and who live in remote regions.
 - Follow <u>AAP recommendations</u> for palivizumab-eligible infants aged <8 months when the appropriate dose of nirsevimab is not available.

MDHHS Prioritization cont....



- Avoid using two 50 mg doses for infants weighing ≥5 kilograms (≥11 pounds), because 50 mg doses should be reserved only for infants weighing <5 kilograms (<11 pounds).
- Providers should encourage pregnant people to receive RSVpreF vaccine (Abrysvo, Pfizer) during 32 weeks' gestation through 36 weeks.
- VFC supply stock should remain prioritized for VFC eligible children.
 - While there are supply constraints for nirsevimab, bi-directional borrowing will not be allowed for VFC providers.
- MDHHS will continue to review nirsevimab supply and send further messages regarding its usage and prioritization.
- Noontime knowledge slides cover prioritization guidance in detail:
 - <u>https://www.michigan.gov/mdhhs/adult-child-serv/childrenfamilies/immunization/mdhhs-covid-19-vaccine-webinars</u>

RSV Pregnant Persons



- September 22 ACIP met to discuss RSV vaccines for pregnant persons to protect infants:
 - Recommended administration 32-36 weeks.
 - VFC doses for pregnant persons < 18 years.
- Challenges:
 - Provider awareness of birth parent immunization status at pediatric visits for the infant.
 - MCIR forecasting.

RSV Older Adults



- Shared clinical decision making
 - High risk due to comorbidities
 - Frailty
 - Long term care residents
- Challenges
 - Billing insurance
 - Coadministration

Michigan's Immunization Rates



Childhood Coverage is Alarmingly Low





HHS

Series - 4 DTaP, 3 Polio, 1 MMR, 3 Hib, 3 HepB, 1 Varicella, 4 PCV

All Regions Impacted





Adolescent Coverage is Not Affected as Much





Statewide 3-month moving average



Series - 1 Tdap, 3 Polio, 2 MMR, 3 HepB, 2 Varicella, 1 MenACWY

Similar Pattern in All Regions





Number of Counties with Childhood Coverage of 70% or Lower





Childhood Series Coverage by County







Counties shaded green have coverage < 70%

2023



More 7th Graders Attend Schools with Low Completion of Required Series





Middle schools with vaccine coverage <70%

A Bright Spot





Trends in Childhood Immunizations 2013 - 2023



Challenges



- Increasing vaccine hesitancy
- Distrust in government
- Data quality
- New and more expensive immunization products
- Staff turnover/burnout

Strategies



- Better data, better use of the data
 - Data quality focus
 - Available, complete, timely, valid
 - Deduplication
- Trusted messengers
- Access
 - Provider deserts
 - Retain VFC providers
 - Alternative providers/community outreach



AA1 Project



Target population:

• Children 19 through 35 months of age who live in Oscoda, Gladwin, or Houghton counties.

Goals:

- Identify and establish collaborations with influential community members and organizations
- Conduct listening sessions/focus groups
- Conduct vaccine clinics at non-traditional venues
- Reduce provider level-vaccine hesitancy.
- Increase vaccination coverage and timeliness in these jurisdictions.
- Reduce school and childcare waivers.
- Established a model that can be used in other jurisdictions or scaled up to regional or statewide interventions.

| Socioeconomic indicators | Oscoda | Gladwin | Houghton | State Average |
|---|--------|---------|----------|---------------|
| Percent ≤ 150% poverty level | 24.7 | 29.0 | 30.5 | 23.2 |
| Percent Uninsured | 13.6 | 9.0 | 7.0 | 6.3 |
| Medicaid Eligibility among children 19- | 53.2 | 52.8 | 47.9 | 47.5 |
| 36 months | | | | |
| Vaccination | Oscoda | Gladwin | Houghton | State Average |
| Primary childhood series (4313314) | 31.8% | 58.9% | 55.5% | 65.8% |

Vaccine Champions



- Convene identified vaccine champions from a variety of health organizations to discuss immunization priorities in Michigan
- First step in aligning and advancing shared goals, developing relationships, and reestablishing a statewide Immunization Coalition.
- Commit to prioritizing vaccine uptake across the state and identify best practices to move this important work forward.
- "Vaccine Champions" will meet leaders from other sectors (health insurance payers, medical organizations, health systems, and local health departments) resulting in opportunities for cross-collaboration.





Recap



- COVID Vaccines are commercialized.
 - MDHHS has a role in Bridge and VFC and in coordinating community outreach.
- RSV immunizations (vaccines for pregnancy and monoclonal antibodies) are being rolled out.
- Immunization coverage continues to decline statewide and retention of VFC Providers is a key part of reversing that trend.
- Rebuilding trust, ensuring equitable access, improving the way we gather and use data will also be keys to improving our coverage.



Back to Basics: Vaccine Errors and Ways to Avoid Them-Pediatric and Adult Case Studies

FALL IMMUNIZATION CONFERENCE 2023 PRESENTER: DIANNE ANKLEY BSN, RN



Use the Vaccine Administration Protocols:

Review Immunization History

Reviewing and assessing a patient's immunization history should be done at every health care visit to help determine which vaccines may be needed

Assess for Needed Immunizations

Use the current Advisory Committee on Immunization Practices (ACIP) immunization schedule to determine what recommended vaccines are needed based on the patient's immunization history

Screen for Contraindications and Precautions

Screening for contraindications and precautions can prevent adverse events following vaccination. All patients should be screened for contraindications and precautions prior to administering any vaccine, even if the patient has previously received that vaccine

Vaccine Administration Protocols | CDC



Vaccine Administration Protocols Cont:

Educate the Patient

Health care professionals should be prepared to provide comprehensive vaccine information

Prepare the Vaccine(s)

Proper preparation is critical for maintaining the integrity of the vaccine during transfer from the vial to the syringe

Administer the Vaccine(s)

Each vaccine has a recommended administration route and site, which are based on clinical trials, practical experience, and theoretical considerations

Document the Vaccination(s)

Health care providers are required by law to record certain information in a patient's medical record

Vaccine Administration Protocols | CDC

Before You Vaccinate Consider the Following:

During your assessment, respond to these four questions:

- 1. What vaccine(s) does the person need today?
- 2. Based on the persons current medical history are there any contraindications or precautions to receiving the recommended vaccines today?
- 3. Are there any medical conditions or other indications that would suggest the need for additional vaccines?
- 4. What vaccine(s) will you give today?
- 5. When should this person return and what vaccines will be recommended at that visit?

Vaccine Administration Protocols | CDC



Know Your Pediatric Resources and Where to Find Them

2023 Recommended Immunization Schedule for Children and Adolescents



2023 Centers for Disease Control (CDC) Child and adolescent Immunization Schedule
| Table 1 Recomm | 0-19 ner | vacci nded (| nation Child a | recomn ind Ad | nendati olesce | ions ha nt Imr | ve chai nuniza | nged. Fin ation Sc | nd the la hedule | e for ages | mmend i 18 ye | ations a ars or y | t www.cdc ounger, l | gov/c United | ovidscher States, | dule 2023 | |
|---|-------------|---------------------------|--------------------------|----------------------|-------------------------|-----------------------------|---------------------------|-----------------------------|------------------------------|--------------------------------------|------------------|-----------------------------|---|----------------------|-------------------------------------|--------------------------|-----------------------------|
| These recommendations must be To determine minimum intervals be | e rea | d with the | e notes th see the ca | at follow. | For those edule (Tab | who fall be ple 2). | hind or st | art late, prov | vide catch-u | up vaccination | at the earli | iest opportu | nity as indicate | ed by the | green bars. | | |
| 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 yr | s 7–10 yrs 1 | 11–12 yrs | 13–15 yrs 16 | 5 yrs 17–18 yr: | |
| Hepatitis B (HepB) | | 1 [≠] dose | < 2 nd | dose> | | 4 | | 3 rd dose | | | | | | | | | |
| Rotavirus (RV): RV1 (2-dose series), RV5 (3-dose series) | | | | 1 st dose | 2 nd dose | See Notes | | | | | | | | | | = Ran | ige of recommended ages |
| Diphtheria, tetanus, acellular pertus (DTaP <7 yrs) | sis | | | 1ª dose | 2 nd dose | 3 rd dose | | | ∢ 4 th dos | se> | | 5 th dos | e | | | for | all children |
| Haemophilus influenzae type b (Hib) | | | | 1¤ dose | 2 nd dose | See Notes | | | dose, otes | | | | | | | = Ran | ge of recommended ages for |
| Pneumococcal conjugate (PCV13, PCV15) | | | | 1ª dose | 2 nd dose | 3 rd dose | | ∢ 4 th de | ose ——• | | - | | . | | | cate | ch-up |
| Inactivated poliovirus (IPV <18 yrs) | | | | 1ª dose | 2 nd dose | • | | — 3 rd dose — | | | | 4 th dos | e | | | Dav | |
| COVID-19 (1vCOV-mRNA, 2vCOV-mRNA, 1vCOV-aPS) | | | | | | | | | | 2- or 3- dose pr | imary series | s and booster | (See Notes) | | | = Ran | ige of recommended ages for |
| Influenza (IIV4) | | | | | | | | P | Annual vaccin | nation 1 or 2 dos | es | | 07 - | A | | cer | tain nign-risk groups |
| Influenza (LAIV4) | | | | | | | | | | | | Annual vaccir 1 or 2 dos | nation les | | E 23 | = Rec | ommended vaccination can |
| Measles, mumps, rubella (MMR) | | | | | | Seel | Notes | ∢ 1 [#] do | ose> | G. | | 2 nd dos | e | | | beg | in in this age group |
| Varicella (VAR) | | | | | | _ | | ∢ 1 [#] do | ose> | | | 2 nd dos | e | | | - Rec | ommended based on shared |
| Hepatitis A (HepA) | | | | | | Seel | Notes | 2 | -dose series, | See Notes | | | _ | | | clin | ical decision making |
| Tetanus, diphtheria, acellular pertus (Tdap ≥7 yrs) | sis | | | | | | | | | | | | | do | | Ciiii | |
| Human papillomavirus (HPV) | | | | | | | | | | | | | | See Note | | = No | recommendation/not |
| Meningococcal (MenACWY-D ≥9 mo MenACWY-CRM ≥2 mos, MenACWY ≥2years) | ŤT | | _ | | | | | | See Notes | | | | | 1ª do | | арр | licable |
| Meningococcal B (MenB-4C, MenB-FHbp) | | | | | | | | | | | | | | | Secnoies | | |
| Pneumococcal polysaccharide (PPSV23) | | | | | | | | | | | | | S | See Notes | | | |
| Dengue (DEN4CYD; 9-16 yrs) | | | | | | | | | | | | | | Seroposi dengue a | tive in endemic reas (See Notes) | | |
| Range of recommended ages for all children | | Range of r for catch-u | ecommend up vaccinati | led ages ion | Rar for | nge of recor certain hig | nmended a n-risk group | ages os | Recomme can begin | ended vaccinatio in this age grou | p | Recommer on shared o | nded vaccination clinical decision-n | n based making | No reco not ap | ommendation/ plicable | |
| | | | | | _ | | | | | | | | | | | | |

Birth-18 Years Immunization Schedule – Healthcare Providers | CDC

| Table 3 | Recomm United St | ended Child tates, 2023 | and Adolescent In | munization S | Schedule by M | Aedical India | cation, | | | | | |
|---|---|--|---|--|--|---|--|------------------------------------|-----------|--------------|-----------------|---------|
| Always use this table in a | onjunction wit | th Table 1 and the N | otes that follow. | | | | | | | | | |
| | | | HIV infection CD4+ count* | | | | 1 | | | | | |
| | | Immunocom- promised status | <15% or total ≥15% and to | tal end-stage renal | | CSFlea | ak Asplenia or | Chronic | | | | |
| VACCINE | Pregnancy | (excluding Hiv infection) | of <200/mm ³ of ≥200/mr | ht disease, or on hemodialysis | chronic lung dis | ease implan | ear persistent complement nt component deficiencies | disease Diabete | es | | | |
| Hepatitis B | | | | | | | | | | | | |
| Rotavirus | | SCID ^b | | | | | = Vaccination a | ccording to | the rou | itine recor | nmended | |
| Diphtheria, tetanus, and acellular pertussis (DTaP) | | | | | | | | | | | michaea | |
| Haemophilus influenzae type b | | | | | | | schedule | | | | | |
| Pneumococcal conjugate | | | | | | | = Recommende | ed for perso | ons with | n an additio | onal risk facto | or for |
| Inactivated poliovirus | | | | | | | which the vac | cine would | be indi | cated | | |
| COVID-19 | | See Notes | See Notes | | | | | | | | | |
| Influenza (IIV4) | | | | | | 12 23 | = Vaccination is | s recommen | nded, ai | nd additior | hal doses ma | y be |
| Influenza (LAIV4) | | | | | Asthma, wheezin | | necessary bas | sed on medi | ical con | dition. See | e Notes | |
| Measles, mumps, rubella | * | | | | | | = Precaution-va | accine migh | t he inc | licated if b | enefit of pro | tection |
| Varicella | * | | | | | | outweighs ris | k of adverse | e reactio | nouted in 5 | cheffe of pro | |
| Hepatitis A | | | | | | | out reigns his | | | 011 | | |
| Tetanus, diphtheria, and acellular pertussis (Tdap) | | | | | | | = Contraindicat | ed or not re | ecomm | ended-vac | cine should r | not be |
| Human papillomavirus | * | | | | | | administered | (*Vaccinate | e after p | oregnancy) | | |
| Meningococcal ACWY | | | | | | | | | | | | |
| Meningococcal B | | | | | | | = No recomme | ndation/not | t applic | able | | |
| Pneumococcal polysaccharide | | | | | | | | | | | | |
| Dengue | | | | | | | | | _ | | | |
| Vaccination according routine schedule recommended | to the | Recommended for persons with an additio factor for which the vac would be indicated | conal risk Vaccination is re and additional d necessary based condition or vac | commended, oses may be on medical ine. See Notes. | Precaution-vaccine might be indicated if be of protection outweight of adverse reaction | enefit recom s risk be adr *Vacci | aindicated or not nmended-vaccine should not ministered inate after pregnancy | No recommendation/ne applicable | ot | | | |
| a. For additional informat www.cdc.gov/vaccines b. Severe Combined Imm c. LAIV4 contraindicated f | ion regarding HIV /hcp/acip-recs/ge unodeficiency or children 2–4 ye | laboratory parameters a neral-recs/immunocom | and use of live vaccines, see the Ge petence.html and Table 4-1 (footn or wheezing during the preceding | neral Best Practice Guideli ote J) at www.cdc.gov/va 12 months | nes for Immunization, "Alt accines/hcp/acip-recs/ger | ered Immunocompete neral-recs/contraindica | ence," at ations.html. | | | | | |

Birth-18 Years Immunization Schedule – Healthcare Providers | CDC

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

Routine vaccination

- ActHIB[®], Hiberix[®], Pentacel[®], or Vaxelis[®]: 4-dose series (3dose 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 -3-dose series 4 weeks ap successful transplant, re

 Anatomic or functional sickle cell disease): Age 12-59 months - Unvaccinated or only 1 2 doses, 8 weeks apart

- 2 or more doses before
- 1 dose at least 8 weeks a Unvaccinated* persons age

- 1 dose

• Elective splenectomy:

Unvaccinated* persons age - 1 dose (preferably at leas

• HIV infection: Age 12-59 months Unvaccinated or only 1 2 doses, 8 weeks apart - 2 or more doses before a 1 dose at least 8 weeks a

- Unvaccinated* persons age - 1 dose
- Immunoglobulin deficie complement deficiency: Age 12-59 months Unvaccinated or only 1
- 2 doses, 8 weeks apart - 2 or more doses before a
- 1 dose at least 8 weeks a *Unvaccinated = Less than

14 months) OR no doses (**Hepatitis A vaccinatio** (minimum age: 12 mor

Routine vaccination

- 2-dose series (minimum i age 12-23 months
- **Catch-up vaccination** Unvaccinated persons threads and the second s
- a 2-dose series (minimum Persons who previously re
- older should receive dose

For vaccination recommendations for persons ages 19 years or older, see the Recommended Adult Immunization Schedule, 2023.

Additional information

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

For calculating intervals between doses, 4 weeks = 28 days. Intervals of \geq 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.gov/travel/.

For vaccination of persons with immunodeficiencies, see Table 8-1, Vaccination of persons with primary and secondary immunodeficiencies, in General Best Practice Guidelines for Immunization at www.cdc.gov/vaccines/hcp/acip-recs/ general-recs/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.

See Addendum for new or updated ACIP vaccine recommendations

Notes Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2023

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/clinicalconsiderations/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 www.cdc.gov/mmwr/volumes/70/rr/ rr7006a1.htm?s_cid=rr7006a1_w and www.cdc.gov/dengue/ vaccine/hcp/index.html
- 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 4th dose that was inadvertently administered as early as age 12 months may be counted if at least 4 months have elapsed since dose 3.

Catch-up vaccination

 Dose 5 is not necessary if dose 4 was administered at age 4 years or older and at least 6 months after dose 3.

For other catch-up guidance, see Table 2.

Special situations

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

Recommended Child and Adolescent Immunization Schedule, United States, 2023 (cdc.gov)

| Appendix | Recommended Child and Adolescent Im | munization Schedule | for ages 18 years or younger, United States, 2023 | | | |
|--|--|--|---|--|--|--|
| Vaccine | Contraindicated or Not Recommended ¹ | | | | | |
| Dengue (DEN4CYD) | Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of cher immunodeficiency, long-term immunosuppressive therapy or patients with H immunocompromised) Lack of laboratory confirmation of a previous Dengue infection | Appendix Guide to Contraindication | Recommended Child and Adolescent Immunization Sched | ule for ages 18 years or younger, United States, 2023 | | |
| Diphtheria, tetanus, pertussis (DTaP) Tetanus, diphtheria (DT) | Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine For DTaP only: Encephalopathy (e.g., coma, decreased level of consciousness, to another identifiable cause within 7 days of administration of previous dose | Adapted from Table 4-1 in Adv recs/general-recs/contraindic | sory Committee on Immunization Practices (ACIP) General Best Practice Guidelines for Immuniz ations.html and ACIP's Recommendations for the Prevention and Control of 2022-23 seasonal i | ation: Contraindication and Precautions available at www.cdc.gov/vaccines/hcp/acip- nfluenza with Vaccines available at www.cdc.gov/mmwr/volumes/71/rr/rr7101a1.htm. | | |
| Haemophilus influenzae type b (Hib) | Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine For Hiberix, ActHib, and PedvaxHIB only: History of severe allergic reaction to Less than age 6 weeks | For COVID-19 vaccine | contraindications and precautions see www.cdc.gov/vaccines/covid-19/clinical- | considerations/interim-considerations-us.html#contraindications | | |
| Hepatitis A (HepA) | Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine | | | | | |
| Hepatitis B (HepB) | Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine Prenancy: Heolisay-B and PreHeybrio are not recommended due to lack of safet | Vaccine | Contraindicated or Not Recommended ¹ | Precautions ² | | |
| Hepatitis A-Hepatitis B vaccine [HepA- HepB, (Twinrix*)] | hepatitis B vaccines if HepB is indicated ⁴ . Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component ¹ including neomycin and yeast | Influenza, egg-based, inactivated injectable (IIV4) | Severe allergic reaction (e.g., anaphylaxis) after previous dose of any influenza vaccine (i.e., any egg-based IIV, ccIIV, RIV, or LAIV of any valency) Severe allergic reaction (e.g., anaphylaxis) to any vaccine component³ (excluding egg) | Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine Moderate or severe acute illness with or without fever | | |
| Human papillomavirus (HPV) | Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine Pregnancy: HPV vaccination not recommended | Influenza, cell culture-based | Severe allergic reaction (e.g., anaphylaxis) to any cclIV of any valency, or to any component³ | Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of | | |
| Measles, mumps, rubella (MMR) Measles, mumps, rubella, and varicella (MMRV) | Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of cher congenital immunodeficiency, long-term immunosuppressive therapy or pat HIV infection who are severely immunocompromised) Pregnancy Framily history of altered immunocompetence, unless verified clinically or by laboratory testing as immunocompetent | inactivated injectable [(ccllV4), Flucelvax® Quadrivalent] | of ccIIV4 | influenza vaccine Persons with a history of severe allergic reaction (e.g., anaphylaxis) after a previous dose of any egg-based IIV, RIV, or LAIV of any valency. If using cclV4, administer in medical setting under supervision of health care provider who can recognize and manage severe allergic reactions. May consult an allergist. Moderate or severe acute illness with or without fever | | |
| 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 For MenACWY-D and Men ACWY-CRM only: severe allergic reaction to any dip or CRM197-containing vaccine For MenACWY-TT only: severe allergic reaction to a tetanus toxoid-containing | Influenza, recombinant injectable [(RIV4), Flublok® | \bullet Severe allergic reaction (e.g., an aphylaxis) to any RIV of any valency, or to any component 3 of RIV4 | Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine Persons with a history of severe allergic reaction (e.g., anaphylaxis) after a previous dose | | |
| Meningococcal B (MenB) [MenB-4C (Bexsero®); MenB-FHbp (Trumenba®)] | Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine | Quadrivalent] | | 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 | | |
| Pneumococcal conjugate (PCV) | Severe allergic reaction (e.g. anaphylaxis) after a previous dose or to a vaccine Severe allergic reaction (e.g. anaphylaxis) to any diphtheria-toxoid-containing | | | 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 | Influenza, live attenuated | Severe allergic reaction (e.g., anaphylaxis) after previous dose of any influenza vaccine (i.e., | Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of | | |
| Poliovirus vaccine, inactivated (IPV) Potaviaus (PV) (PV/1 (Potavix [®]) | Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine Severe allergic reaction (e.g., anaphylaxic) after a previous dose or to a vaccine | [LAIV4, Flumist® Quadrivalent] | any egg-based IIV, ccIIV, RIV, or LAIV of any valency) • Severe allergic reaction (e.g., anaphylaxis) to any vaccine component ³ (excluding egg) | influenza vaccine • Asthma in persons aged 5 years old or older | | |
| RV5 (RotaTeq*)] | Severe conditional immunodeficiency (SCID) History of intussusception | | 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 | 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, | | |
| Tetanus, diphtheria, and acellular pertussis (Tdap) Tetanus, diphtheria (Td) | Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine For Tdap only: Encephalopathy (e.g., coma, decreased level of consciousness, to another identifiable cause within 7 days of administration of previous dose | | infection • Close contacts or caregivers of severely immunosuppressed persons who require a protected environment • Pregnancy • Cochlear implant | Moderate or severe acute illness with or without fever | | |
| Varicella (VAR) | Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine Severe immunodeficiency (e.g., hematologic and solid tumors, receipt of cher congenital immunodeficiency, long-term immunosuppressive therapy or pat HIV infection who are severely immunocompromised) Pregnancy Family history of altered immunocompetence, unless verified clinically or by laboratory testing as immunocompetent | | Active communication between the cereorospinal 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 | | | |
| When a contraindication is present When a precaution is present, vacc Guidelines for Immunization. www Vaccination providers should chec www.fda.gov/vaccines-blood-blok For information on the pregnancy | t, a vaccine should NOT be administered. Kroger A, Bahta L, Hunter P. ACIF cination should generally be deferred but might be indicated if the benef w.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html & FDA-approved prescribing information for the most complete and upda logics/approved-products/vaccines-licensed-use-united-states. exposure registries for persons who were inadvertently vaccinated with H | When a contraindication is p contraindications.html When a precaution is preser General Best Practice Guide Vaccination providers shoul vaccines are available at www. | present, a vaccine should NOT be administered. Kroger A, Bahta L, Hunter P. ACIP General Best Protection for the should generally be deferred but might be indicated if the benefit of protection fr lines for Immunization. www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.htm d check FDA-approved prescribing information for the most complete and updated information ww.fda.gov/vaccines-blood-biologics/approved-products/vaccines-licensed-use-united-states | actice Guidelines for Immunization. www.cdc.gov/vaccines/hcp/acip-recs/general-recs/ om the vaccine outweighs the risk for an adverse reaction. Kroger A, Bahta L, Hunter P. ACIP nl , including contraindications, warnings, and precautions. Package inserts for U.Slicensed | | |

Recommended Child and Adolescent Immunization Schedule, United States, 2023 (cdc.gov)

| Child and Adolesce | ent Immunization Sc | hedule by Age | | | |
|---|---|--|---|---|---|
| Recommendations for Ages 18 Ye | ars or Younger, United States, 2023 | | | | |
| Print | Addendum – Cł Schedule for ag | nild and Ado ges 18 years | olescent Recor s or younger, L | mmended Immunization Jnited States, 2023 | |
| See Addendum for new or update | Vaccines and Other Immunizing Agents | Re | commendations | Effective Date of Recommendation* | |
| | Respiratory syncytial virus | Addendum In addition to the recommen recommendations have been Vaccines and Other | Recommended Child and ndations presented in the previous sections in adopted by the CDC Director and are now | d Adolescent Immunization Schedule for ages 18 years or younger, Uni of this Immunization Schedule, ACIP has approved the following recommendations by majority vote since it official. Links are provided if these recommendations have been published in <i>Morbidity and Mortality Week</i> | ted States, 2023 October 20, 2022. The following <i>ly Report (MMWR)</i> . Effective Date of |
| Using the schedule | | Immunizing Agents COVID-19 (Moderna, Pfizer-BioNTech) | All persons ≥6 months of age should rec For detailed information, see: www.cdc.g | eive 2023–2024 (monovalent, XBB containing) COVID-19 vaccines as authorized under EUA or approved by BLA jov/covidschedule | Recommendation* September 12, 2023 |
| To make vaccination recommendations, 1. Determine needed vaccines based o 2. Determine appropriate intervals for | | Respiratory syncytial virus [RSV-mAb (Nirsevimab)] | All infants younger than 8 months and b 1 week of birth either in hospital or outp Infants younger than age 8 months not b 1 dose of nirsevimab shortly before the s Infants aged 8-19 months with chronic I supplemental oxygen) any time during t for length <10th percentile; or with man abnormalities on chest imaging that per Infants 8-19 months who are American I Infants who are age-eligible and underg For detailed information, see: www.cdc.c | orn shortly before or during the RSV season should receive 1 dose of nirsevimab within atient setting born during RSV season and now entering their first RSV season should receive tart of RSV season ung disease of prematurity requiring medical support (e.g., chronic corticosteroid therapy, diuretic therapy, or he 6-month period before start of the second RSV season; severe immunocompromise; cystic fibrosis with weig festation of severe lung disease (e.g., previous hospitalization for pulmonary exacerbation in the first year of life sist when stable) should receive 1 dose of nirsevimab shortly before start of second RSV season ndian or Alaska Native should receive 1 dose of nirsevimab before start of second RSV season oing cardiac surgery with cardiopulmonary bypass should receive 1 additional dose of nirsevimab after surgery gov/mmwr/volumes/72/wr/mm7234a4.htm?s_cid=mm7234a4_w | ht August 3, 2023 e or |
| 3. Assess for medical conditions and ot | COVID-19 (Moderna Pfize | Poliovirus (IPV) | Adolescents age 18 years who are known primary vaccination series with inactivat Adolescents age 18 years who have receipoliovirus exposure may receive another | n or suspected to be unvaccinated or incompletely vaccinated against polio should complete a ed polio vaccine (IPV). ived a primary series of trivalent oral polio vaccine (tOPV) or IPV in any combination and who are at increased ri dose of IPV. Available data do not indicate the need for more than a single lifetime booster dose with IPV for ad | sk of June 27, 2023 Jults. |
| Review special situations (<u>Vaccination</u> Review contraindications and precautions | COVID-15 (Moderna, 1120 | Influenza (IIV4, cclV4, RIV4, LAIV4) | All persons ages ≥6 months with egg all- that is otherwise appropriate for the reci Affirm the updated MMWR Recommend. Committee on Immunization Practices— | ergy should receive influenza vaccine. Any influenza vaccine (egg based or non-egg based) pient's age and health status can be used. ations and Reports, "Prevention and Control of Seasonal Influenza with Vaccines: Recommendations of the Advis United States, 2023-24 Influenza Season" www.cdc.gov/mmwr/volumes/72/rr/rr7202a1.htm | sory June 27, 2023 |
| 6. See Addendum for new or updated A | | Pneumococcal (PCV15, PCV20) | Use of either pneumococcal conjugate v currently recommended PCV dosing and For children with an incomplete PCV vac PCV dosing and schedules is recommend- Healthy children aged 24–59 months - Children with specified health conditio For children aged 2–18 years with any ris Using 21 dose(s) of PCV20: No addition updated as additional data become av - Using PCV13 or PCV15 (no PCV20): A d For children aged 6–18 years with any ris PCV20 is recommended. When PCV15 is | accines (PCV) PCV15 or PCV20 is recommended for all children aged 2-23 months according to i schedules. cination status, use of either PCV15 or PCV20 according to currently recommended ded for: ins(2) aged 24 through 71 months is condition who have received all recommended doses of PCV before age 6 years hal doses of any pneumococcal vaccine are indicated. This recommendation may be ailable. ose of PCV20 or PPSV23 using previously recommended dosing and schedules is recommended. is condition who have not received any dose of PCV13, PCV15, or PCV20, a single dose of PCV15 or used, it should be followed by a dose of PPSV23 at least 8 weeks later if not previously given. | June 27, 2023 |

Birth-18 Years Immunization Schedule – Healthcare Providers | CDC

General Best Practice Guidelines for Immunization

Updated August 1, 2023

Best Practices Guidance Kroger A, Bahta L, Long S, Sanchez P

Introduction Purpose and topics covered in this report...

History

History of development of: Timing and Spacing, Contraindications and Precautions, Preventing and Managing Adverse Reactions...

Timing and Spacing of Immunobiologics

Vaccine scheduling, supply and lapsed schedule, spacing of doses, simultaneous and nonsimultaneous administration, licensed combination vaccines, interchangeability of formulations, extra doses, conjugate vaccines...

Contraindications and Precautions

General principles, standards of valid contraindications and precautions, and conditions incorrectly perceived as contraindications...

Preventing and Managing Adverse Reactions

Benefit and risk communication, reporting adverse reactions, National Vaccine Injury Compensation Program...

Vaccine Administration

Infection control and sterile technique, route of administration, multiple and jet injections. alleviating discomfort and pain, clinical implications of nonstandard practices...

TABLE 3-2. Recommended and minimum ages and intervals between vaccine doses

| Vaccine and dose number | Recommended age for this dose | Minimum age for this dose | Recommended interval to next dose | Minimum interval to next dose |
|----------------------------|-------------------------------|---------------------------------|---|-------------------------------------|
| DTaP-1 ^(e) | 2 months | 6 weeks | 8 weeks | 4 weeks |
| DTaP-2 | 4 months | 10 weeks | 8 weeks | 4 weeks |
| DTaP-3 | 6 months | 14 weeks | 6-12 months ^(f) | 6 months ^(f) |
| DTaP-4 | 15-18 months | 15 months ^(f) | 3 years | 6 months |
| DTaP-5 ^(g) | 4-6 years | 4 years | - | _ |
| HepA-1 ^(e) | 12-23 months | 12 months | 6-18 months | 6 months |
| НерА-2 | ≥18 months | 18 months | - | |
| HepB-1 ^(h) | Birth | Birth | 4 weeks-4 months | 4 weeks |
| НерВ-2 | 1-2 months | 4 weeks | 8 weeks-17 months | 8 weeks |
| HepB-3 ⁽ⁱ⁾ | 6-18 months | 24 weeks | - | _ |
| Hib-1 ⁽⁾⁾ | 2 months | 6 weeks | 8 weeks | 4 weeks |
| Hib-2 | 4 months | 10 weeks | 8 weeks | 4 weeks |

ACIP General Best Practice Guidelines for Immunization | CDC

Recommended Catch-up Immunization Schedule for Children and Adolescents Who Start Late or Who Are More Table 2 Table 2 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 the section with rable r and many and many and many set the section appropriate for the child's age. Always use the section with rable r and many set to be restarted.

| | | | | Children age 4 months through 6 years | | |
|---|--|--|---|---|--|------------------|
| Vaccine | Minimum Age for | | | Minimum Interva Between Doses | | |
| | Dose 1 | Dose 1 to Dose 2 | Dose 2 | o Dose s | Dose 3 to Dose 4 | Dose 4 to Dose 5 |
| Hepatitis B | Birth | 4 weeks | 8 weeks | and at least 16 weeks after first dose mage for the final dose is 24 weeks | | |
| Rotavirus | 6 weeks Maximum age for first dose is 14 weeks, 6 days. | 4 weeks | 4 weeks maximu | m age for final dose is 8 months, 0 days | | |
| Diphtheria, tetanus, and acellular pertussis | 6 weeks | 4 weeks | 4 weeks | | 6 months | 6 months |
| Haemophilus influenzae type b | 6 weeks | No further doses needed if first dose was administered at age 15 months or older. 4 weeks if first dose was administered before the 1 st birthday. 8 weeks (as final dose) if first dose was administered at age 12 through 14 months. | No furth if previou 4 weeks if curren 1 previou 8 weeks if curren OR if curren administ OR if both d | her doses needed us dose was administered at age 15 months or older t age is younger than 12 months and first dose was administered at younger than age 7 months and at least us dose was PRP-T (ActHib [®] , Pentacel [®] , Hiberix [®]), Vaxelis [®] or unknown and age 12 through 59 months (as final dose) t age is younger than 12 months and first dose was administered at age 7 through 11 months; t age is 12 through 59 months and first dose was administered before the 1 st birthday and second dose was tered at younger than 15 months; oses were PedvaxHIB [®] and were administered before the 1st birthday | 8 weeks (as final dose) This dose only necessary for children age 12 through 59 months who received 3 doses before the 1 st birthday. | |
| Pneumococcal conjugate | 6 weeks | No further doses needed for healthy children if first dose was administered at age 24 months or older 4 weeks if first dose was administered before the 1 st birthday 8 weeks (as final dose for healthy children) if first dose was administered at the 1 st birthday or after | No furth for healt 4 weeks if curren 8 weeks if previor OR if curren | her doses needed hy children if previous dose was administered at age 24 months or older t age is younger than 12 months and previous dose was administered at <7 months old (as final dose for healthy children) us dose was administered between 7–11 months (wait until at least 12 months old); t 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 6 month if current | t age is <4 years Is (as final dose) t 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 Note | | See Notes | |
| | | | Chil | dren and adolescents age 7 through 18 years | 4); 4); | |
| Meningococcal ACWY | Not applicable (N/A) | 8 weeks | | | | |
| Tetanus, diphtheria; tetanus, diphtheria, and acellular pertussis | 7 years | 4 weeks | 4 weeks if first do 6 month if first do | ise of DTaP/DT was administered before the 1 st birthday is (as final dose) ise of DTaP/DT or Tdap/Td was administered at or after the 1 st birthday | 6 months if first dose of DTaP/DT was administered before the 1 st birthday | |
| Human papillomavirus | 9 years | Routine dosing intervals are recommended. | | | | |
| Hepatitis A | N/A | 6 months | | | | |
| Hepatitis B | N/A | 4 weeks | 8 weeks | and at least 16 weeks after first dose | | |
| Inactivated poliovirus | N/A | 4 weeks | 6 month A fourth the prev | is dose is not necessary if the third dose was administered at age 4 years or older and at least 6 months after ious 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 month | s | | |
| | | | | | | |

Recommended Child and Adolescent Immunization Schedule, United States, 2023 (cdc.gov)

Other Catch-Up Resources



www.cdc.gov/vaccines/schedules/hcp/imz/catchup.html#guidance

COVID-19 Vaccine Clinical Considerations

| Interim Clinical Considerations for Use | of COVID-19 | | | | | | |
|--|--|---|---|--|-------------------|--|-------------------------------------|
| Vaccines in the United States | Ages 6 months-4 ye | ears | | | | | |
| Print | COVID-19 vaccination I prior to updated (2023 | Ages 5–11 years [‡] | er of updated | Vaccine vial | | | |
| Summary of recent changes (last updated October 24, 2023): Age transitions: Updated guidance for children who transition during the initial CC 4 years to age 5 years and children who are moderately or severely immunocomp years to age 12 years to receive the age-appropriate dosage based on their age or Interchangeability of COVID 10 upgeinger Clarification of circumstances in which account of the coving severely immunocomp of the coving severely immunocoving severely immunocovi | Formula) mRNA vacc Unvaccinated | COVID-19 vaccination history prior to updated (2023–2024 Formula) mRNA vaccine* | Updated (2023– 2024 Formula) mRNA vaccine | Number of updated (2023–2024 Formula) mRNA doses indicated | Dosage (mL/ug) | Vaccine vial cap and label colors | Interval between doses |
| Interchangeability of COVID-19 vaccines. Clarinication of circumstances in which ac doses from different manufacturers may be considered when doses from the sam recommended. | 1 dose any Moderna 2 or more doses any Mo | Unvaccinated | Moderna | 1 | 0.25 mL/25 ug | Dark blue cap; green label | - |
| Reference Materials | 1 dose any Pfizer-BioNT | | Pfizer-BioNTech | 1 | 0.3 mL/10 ug | Blue cap; blue label | - |
| <u>COVID-19 Vaccination Recommendations Infographic</u> (Updated 10/13/2023) COVID-19 Vaccination Recommendations Infographic | 2 doses any Pfizer-BioN | 1 or more doses any mRNA | Moderna OR | 1 | 0.25 mL/25 ug | Dark blue cap; green label | At least 8 weeks after last dose |
| (Immunocompromised) (Updated 10/13/2023) | 3 or more doses any Pfi BioNTech | | Pfizer-BioNTech | 1 | 0.3 mL/10 ug | Blue cap; blue label | At least 8 weeks after last dose |

Interim Clinical Considerations for Use of COVID-19 Vaccines | CDC



Pediatric Case Studies



Case Study 1: Noah DOB: 6-2-23 Age: 5 months

- Healthy 5-month-old
- Weight today: 10 lbs. 10 oz.
- Received first HepB in the hospital on 6-2-23
- Has not received any other vaccines
- No contraindications or precautions to any vaccines
- Does not have any high-risk indications

NOAH'S MCIR RECORD:

| | | Immunizat | | | Oth | |
|---------------------------|-----------------------------------|-----------|--------|--------|--------|---------|
| Series | Dose 1 | Dose 2 | Dose 3 | Dose 4 | Dose 5 | Dose 6+ |
| DTP/DTaP/ DT/Td/Tdap | | | | | | |
| Polio | | | | | | |
| MMR | | | | | | |
| Hib | | | | | | |
| Hepatitis B | 06/02/2023 Hep B (ped/adol) | | | | | |
| Varicella | | | | | | |
| Pneumococcal Conjugate | | | | | | |
| Hepatitis A | | | | | | |
| Seasonal Influenza | | | | | | |
| SARS-CoV-2 | | | | | | |
| RSV | | | | | | |

Case Study #1: Noah Question #1:

Which vaccines should Noah receive today?

- A. HepB, Rotavirus (RV), DTaP, Hib, Polio
- B. HepB, DTaP, Hib, Pneumococcal, Polio, Nirsevimab (RSV)
- C. HepB, RV, Pneumococcal, Nirsevimab (RSV)
- D. HepB, RV, DTaP, Hib, Pneumococcal, Polio, Nirsevimab(RSV)
- E. None of the above

Question #1: Answer: B

- Hep B
- DTaP
- Hib
- Pneumococcal
- Polio
- Nirsevimab (RSV)
- *Using combination vaccines equals fewer shots:
 - Pediarix (DTaP/IPV/HepB) OR
 - Pentacel (DTaP/IPV/Hib) OR
 - Vaxelis (DTaP/IPV/Hib/HepB)

ACIP Timing and Spacing Guidelines for Immunization | CDC



| Child and Adolescent Immur | vization Schedule | by Age | | | | | |
|--|---|---|--|--|--|--|--|
| Recommendations for Ages 18 Years or Younger, U | Inited States, 202 Heal | thcare Providers: RSV Prevention | | | | | |
| <u>Print</u> | Infor | Information | | | | | |
| See Addendum for new or updated ACIP vaccine recom | RSV Imn | nunization for Infants and Young Children | | | | | |
| Addendum - Child and Add | Diescent R <u>Print</u> | | | | | | |
| Schedule for ages 18 years Vaccines and Other Rec Immunizing Agents Respiratory syncytial virus [RSV-mAb (Nirsevimab)] | • All infant and born the RSV s dose of r of birth e outpatier | On October 23, 2023, CDC released a health advisory notice to communicate interim recommendations regarding the limited supply of nirsevimab, the new preventive antibody to protect infants against severe RSV. Read more: <u>Limited Availability of Nirsevimab in the United States—Interim CDC Recommendations</u> | | | | | |
| | Infants younger than not born during RSV s now entering their fir should receive 1 dose nirsevimab shortly be of RSV season | age 8 months season and rst RSV season e of efore the start | | | | | |
| | Birth-18 Years Immuni | ization Schedule – Healthcare Providers CDC | | | | | |

Case Study: Noah

| Vaccine | Birth | 1 mo | 2 mos | 4 mos | 6 mos |
|--|----------------------------|---------------------------|----------------------|---|------------------------------|
| Hepatitis B (HepB) | 1 [≠] dose | ◄ 2 nd c | dose► | \overleftrightarrow | ۹ |
| Rotavirus (RV): RV1 (2-dose series), RV5 (3-dose series) | | | 1 st dose | 2" Se | See Notes |
| Diphtheria, tetanus, acellular pertussis (DTaP <7 yrs) | | | 1 [#] dose | 2" Se | 3 rd dose |
| Haemophilus influenzae type b (Hib) | | | 1 [≠] dose | 2" Se | See Notes |
| Pneumococcal conjugate (PCV13, PCV15) | | | 1 st dose | 2" ~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~ | 3 rd dose |
| Inactivated poliovirus (IPV <18 yrs) | | | 1 [≠] dose | 2"2" Se | ۹ |
| COVID-19 (1vCOV-mRNA, 2vCOV-mRNA, 1vCOV-aPS) | | | | | |
| Influenza (IIV4) | | | | | |
| Influenza (LAIV4) | | $\mathbf{X} = \mathbf{v}$ | vaccine | es that | were |
| Measles, mumps, rubella (MMR) | | r | ecom | mende | ed for |
| Varicella (VAR) | | 2 months of a | | | age |
| Hepatitis A (HepA) | | | | | See |
| Tetanus, diphtheria, acellular pertussis (Tdap ≥7 yrs) | | | | | |
| Human papillomavirus (HPV) | | | | | |
| Meningococcal (MenACWY-D ≥9 mos, MenACWY-CRM ≥2 mos, MenACWY-TT ≥2years) | | | | | |
| Meningococcal B (MenB-4C, MenB-FHbp) | | | | | |
| Pneumococcal polysaccharide (PPSV23) | | | | | |
| Dengue (DEN4CYD; 9-16 yrs) | | | | | |
| Range of recommended ages for all children | Range of re for catch-u | ecommend p vaccinatio | ed ages on | Rai | nge of recor certain higł |

| | | - | 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 | | | | | |
| Hepatit <mark>is</mark> B | Birth | ⁴ Rotavirus | | 6 weeks | | | | |
| Rotavirus | 6 weeks Maximum age for first dose is 14 weeks, 6 days | | | Maximum age for first | | | | |
| Diphtheria, tetanus, and acellular pertussis | 6 weeks | 4 | | dose is 14 weeks, 6 days. | | | | |
| Haemophilus influenzae type b | 6 weeks | No further doses needed if first dose was administered at age 15 months or older. 4 weeks if first dose was administered before the 1 st birthday. 8 weeks (as final dose) if first dose was administered at age 12 through 14 months. | No further doses a if previous dose wa 4 weeks if current age is yo 1 previous dose wa 8 weeks and age if current age is yo OR if current age is 12 administered at yo OR if both doses were | Use catch-up schedule to determine: • Maximum/minimum | | | | |
| Pneumococcal conjugate | 6 weeks | No further doses needed for healthy children if first dose was administered at age 24 months or older 4 weeks if first dose was administered before the 1 st birthday 8 weeks (as final dose for healthy children) if first dose was administered at the 1 st birthday or after | No further doses for healthy childrei 4 weeks if current age is yo 8 weeks (as final o if previous dose wa OR if current age is 12 | age for the 2-month vaccines to be given today Minimum interval | | | | |
| Inactivated poliovirus | 6 weeks | 4 weeks | 4 weeks if current age is <4 6 months (as fina if current age is 4 y | between 2-month doses and next 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 | | | | | |

Recommended Child and Adolescent Immunization Schedule, United States, 2023 (cdc.gov)

Case Study: Noah Question #2:

When does Noah need to return and what vaccines will he need at that time?

- A. 2-3 months: Hep B, DTaP, IPV, Flu
- B. 1 month: DTaP, Hib, IPV, Pneumococcal, Nirsevimab (RSV)
- C. 2 months: DTaP, Hib, Pneumococcal, IPV, COVID-19
- D. 1 month: DTaP, Hib, Pneumococcal, IPV, Flu, COVID-19
- E. None of the above

Question #2: Answer: D-1 month

- DTaP
- Hib
- Pneumococcal
- IPV
- Flu
- COVID-19

*Using combination vaccines equals fewer shots

- Pediarix (DTaP, IPV, Hep B) OR
- Pentacel (DTaP, IPV, Hib) **OR**
- Vaxelis (DTaP/IPV/Hib/Hep B)

Ped-Flu-Dose-Volume-23-24_FINAL.pdf (michigan.gov)

NOAH'S MCIR RECORD:

| | Immuni | Immunizations | | | | | | | |
|---------------------------|---|---|--------|--------|--------|---------|--|--|--|
| Series | Dose 1 | Dose 2 | Dose 3 | Dose 4 | Dose 5 | Dose 6+ | | | |
| DTP/DTaP/ DT/Td/Tdap | 11/03/2023 DTaP-Hep B-IPV (Pediarix) 5mos 1dy | | | | | | | | |
| Polio | 11/03/2023 DTaP-Hep B-IPV (Pediarix) 5mos 1dy | | | | | | | | |
| MMR | | | | | | | | | |
| Hib | 11/03/2023 Hib (historical) 5mos 1dy | | | | | | | | |
| Hepatitis B | 06/02/2023 Hep B (ped/adol) | 11/03/2023 DTaP-Hep B- IPV (Pediarix) 5mos 1dy | | | | | | | |
| Varicella | | | | | | | | | |
| Pneumococcal Conjugate | 11/03/2023 PCV15 (VAXNEUVANCE) 5mos 1dy | | | | | | | | |
| Hepatitis A | | | | | | | | | |
| Seasonal Influenza | | | | | | | | | |
| SARS-CoV-2 | | | | | | | | | |
| RSV | 11/03/2023 RSV Nirsevimab(Beyfortus)0.5ml 5mos 1dy | | | | | | | | |
| | | | | | | | | | |

| Vaccine | Birth | 1 mo | 2 mos | 4 mos | 6 mos |
|--|---------------------------|--------------------------|---------------------|----------------------|------------------------------|
| Hepatitis B (HepB) | 1 [≠] dose | ◄ 2 nd c | dose> | | ۹ |
| Rotavirus (RV): RV1 (2-dose series), RV5 (3-dose series) | | | 1ª dose | 2 nd dose | See Notes |
| Diphtheria, tetanus, acellular pertussis (DTaP <7 yrs) | | | 1 [#] dose | 2 nd dose | |
| Haemophilus influenzae type b (Hib) | | | 1¤ dose | 2 nd dose | Sector |
| Pneumococcal conjugate (PCV13, PCV15) | | | 1 [⊄] dose | 2 nd dose | 3 Are |
| Inactivated poliovirus (IPV <18 yrs) | | | 1≝ dose | 2 nd dose | \bigstar |
| COVID-19 (1vCOV-mRNA, 2vCOV-mRNA, 1vCOV-aPS) | | | | | |
| Influenza (IIV4) | | | | | |
| Influenza (LAIV4) | | | | | |
| Measles, mumps, rubella (MMR) | | | | | See N |
| Varicella (VAR) | | | | | |
| Hepatitis A (HepA) | | | | | See N |
| Tetanus, diphtheria, acellular pertussis (Tdap ≥7 yrs) | | | | | |
| Human papillomavirus (HPV) | | | | | |
| Meningococcal (MenACWY-D ≥9 mos, MenACWY-CRM ≥2 mos, MenACWY-TT ≥2years) | | | | | |
| Meningococcal B (MenB-4C, MenB-FHbp) | | | | | |
| Pneumococcal polysaccharide (PPSV23) | | | | | |
| Dengue (DEN4CYD; 9-16 yrs) | | | | | |
| Range of recommended ages for all children | Range of r for catch-u | ecommend ip vaccinati | ed ages on | Rar for | nge of recon certain high |

| | | | | Children age 4 months through 6 years |
|--|---|---|--|--|
| Vaccine | Minimum Age for | | | Minimum Interval Retween Doces |
| | Dose 1 | Dose 1 to Dose 2 | | Dose 2 to Dose 3 |
| Hepatitis B | | | Dose 2 | 2 to Dose 3 |
| Rotavirus | 6 weeks Maximum age for first dose is 14 weeks, 6 days. | 4 weeks | 8 wee minim | ks and at least 16 weeks after first dose oum age for the final dose is 24 weeks |
| Diphtheria, tetanus, and acellular pertussis | 6 weeks | 4 weeks | | 4 weeks |
| Haemophilus influenzae type b | 6 weeks | No further doses nee if first dose was admir months or older. 4 weeks if first dose was admir 1 st birthday. 8 weeks (as final dos if first dose was admir 12 through 14 monthe | eded nistered at age 15 nistered before the e) nistered at age s. | 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 <i>and</i> first dose was administered at younger than age 7 months <i>and</i> at least 1 previous dose was PRP-T (ActHib®, Pentacel®, Hiberix®), Vaxelis® or unknown 8 weeks <i>and</i> age 12 through 59 months (as final dose) if current age is younger than 12 months <i>and</i> first dose was administered at age 7 through 11 months; OR if current age is 12 through 59 months <i>and</i> first dose was administered before the 1 st birthday <i>and</i> second dose was administered at younger than 15 months; OR if both doses were PedvaxHIB® and were administered before the 1st birthday |
| Pneumococcal conjugate | 6 weeks | No further doses new children if first dose w age 24 months or olde 4 weeks if first dose was admir 1 st birthday 8 weeks (as final dos children) if first dose was admir 1 st birthday or after | eded for healthy ras administered at er histered before the e for healthy histered at the | 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 |
| 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 |

| Influenza (minimum 18 years [re Routine vac • Use any influ status annua - 2 doses, sep 6 months- 2 influenza whose influ (administer receipt of d | vaccination age: 6 mont combinant combinant cination enza vaccine lly: barated by at 8 years who vaccine dose enza vaccina dose 2 even ose 1 and dose | ths [IIV], 2 year influenza vaco appropriate for least 4 weeks, fo have received fe s before July 1, 2 tion history is ur if the child turns se 2) 6 months-8 yea | rs [LAIV cine, RIV age and r childre wer than 022, or known 9 betwe | 4], /4]) health en age | 2 Ever Flu v vacc were COV form ensu For a adm child Fluze ageo | 023-24 Seasonal Influenz yone aged 6 months and older should rec vaccine dose volume is based on the perso ine errors reported between June 2020 at e listed as numbers 1 and 3 of the most fr ID-19 vaccines. Wrong age and associated nulations of influenza vaccines (31%). ² It is the children are adequately protected aga children aged 6 through 35 months, flu va inistered. There are multiple licensed ina fren aged 6 through 35 months: Afluria [®] C one [®] Quadrivalent. The cell cultured-based d 6 months and older (Flucelvax [®] Quadrival You're Using This Vaccine (IIV4) ¹ | ceive flu vaccine every year. on's age and the flu vaccine product that is us and December 2021, wrong vaccine (24%) and is equent types of vaccine events other than tho d wrong dose errors occurred frequently betwis important to prevent flu vaccine administration inst flu. accine dose volume is dependent on the productivated influenza vaccines, quadrivalent (IIV4) Quadrivalent, FluLaval [®] Quadrivalent, Fluarix [®] C d inactivated influenza vaccine (ccIIV4) is availablent). Dose Volume for Ages 6-35 M 0.25 ml aperdose | br Children ed. ¹ Among wrong age (13%) use related to een age-related ion errors to uct that is 1) available for Quadrivalent, and able for persons | Volumes for Children for only 1 dose. If 0.25 mL is used 25 mL remaining in the single-dose vial tration errors ^{4, 5} : al/Fluarix rather than the 0.25 mL dose of the product that was t discovered until the following day, 0.5 mL of FluLaval, Fluarix, or Revaccinating the day after the initial ble. |
|---|---|--|---|--|---|--|--|--|---|
| have receiv | ed at least 2 i | nfluenza vaccine | doses | | | Afluria (Seqirus) | 0.25 mL per dose | | ii-dose vial rather than the |
| - 1 dose for all persons age 9 years or older | | | | | Fluzone (Sanofi Pasteur) | 0.25 mL OR 0.5 mL per dose *No preference is expressed for either of | e dose volume. | d as valid. Notify the parent/guardian accine administration errors. If the ter an age-appropriate IIV dose (use | |
| Ages 6 months Avears | | | | | | ou're Using This Vaccine (ccIIV4)* | Dose Volume for Ages 6-35 M | onths | |
| | | | | | | Flucelvax (Seqirus) | 0.5mL per dose | | ther than the recommended 0.5 mL |
| COVID-19 vaccination history prior to updated (2023–2024 Formula) mRNA vaccine* | Number of updatedVaccine vialcination historyUpdated (2023- (2023-2024 Formula)Capred (2023-20242024 Formula)mRNA vaccine dosesDosageRNA vaccine*mRNA vaccineindicated(mL/ug)colors | | Interval between doses | Refer to the Flu Vaccine Presentation Chart ¹ for available presentations of each of these vaccines. ildren aged 3 years and older, dose volume for standard-dose IIV is 0.5 mL regardless of the flue product being administered. ses of 2023-24 flu vaccine are needed ³ , the same vaccine product does not need to be used for both Use any age-appropriate flu vaccine that is available that day, ensuring you use the correct dose e for the product you are administering. miss an opportunity to vaccinate! Dose volume is based on the child's age on the day of vaccine istration. For example: | | e vaccines. ss of the flu he used for both correct dose of vaccine he dose volume | 0.25 mL dose of Afluria to correct this i should revaccinate the 5-year-old lay after the initial substandard dose healthy children aged 2 years and ichigan.gov/flu/resources/resources-for- sources/ismp-national-vaccine-errors- w.michigan.gov/flu/resources/resources- | | |
| Unvaccinated | Moderna | 2 | 0.25 mL/25 ug | Dark blue cap; green label | Dose 1 and Dose 2: 4–8 weeks ⁺ | based on the IIV/ccIIV product used . When the child returns 4 weeks later regardless of the IIV/ccIIV product use 7, the needed volume for a child aged 6 ning the appropriate volume (as supplie | for dose 2 and is now aged 3 years , the dose ed. through 35 months may be administered from ed by the manufacturer), a single-dose vial, or | volume is 0.5 mL n a prefilled syringe a multi-dose vial. | ing System (VAERS): <u>https://vaers.hhs.gov</u> . so the status of the error can be <u>endations of the Advisory Committee on</u> <u>ndations and Reports / August 25, 2023 /</u> html |
| | Pfizer-BioNTech | 3 | 0.3 ml /3 | Yellow can | Dose 1 and | er Guidance on Fluzone Quadrivalent: | , | | J, <u>www.cdc.gov/vaccines</u> , or |
| | - HZGE DIONECCI | 5 | ug | yellow label | Dose 2: 3–8 weeks ⁺ Dose 2 and Dose 3: | NOTE: Fluzone Quadrivalent is approve 0.5 mL per dose. The 0.25-mL prefilled syringe of Fluzon Fluzone Quadrivalent is used for a child | ed for children aged 6 through 35 months at e ne Quadrivalent is no longer available. If a pref d in this age group, the dose volume will be 0. | ither 0.25 mL or filled syringe of 5 mL per dose. | n errors, visit Immunize.org's Ask the |
| | | | | | At least 8 weeks | Michigan Department of Health ar | nd Human Services – Division of Immunization | Rev. August 30, 2023 Page 1 of 2 | of Immunization Rev. August 30, 2023 Page 2 of 2 |

Interim Clinical Considerations for Use of COVID-19 Vaccines | CDC

Resources for Health Professionals (michigan.gov)

Case Study 2: Winnie DOB: 5-9-12

Age: 11 years

- Healthy 11-year-old
- Has egg allergy:
 - \circ Hives
 - Nasal congestion
 - \circ Vomiting
- No contraindications or precautions to any vaccines
- Does not have any high-risk indications

WINNIE'S MCIR RECORD:

| | | Immunizatio | ons | | | Other |
|----------------------------|---|--|--|--|--|---------|
| Series | Dose 1 | Dose 2 | Dose 3 | Dose 4 | Dose 5 | Dose 6+ |
| DTP/DTaP/ DT/Td/Tdap | 07/10/2012 DTaP-Hep B- IPV (Pediarix) 2mos 1dy | 09/11/2012 DTaP-Hep B- IPV (Pediarix) 4mos 2dys | 11/12/2012 DTaP-Hep B- IPV (Pediarix) 6mos 3dys | 09/09/2013 DTaP (pediatric) 1yr 4mos | 08/05/2019 Tdap (adol/adult) 7yrs 2mos | |
| Polio | 07/10/2012 DTaP-Hep B- IPV (Pediarix) 2mos 1dy | 09/11/2012 DTaP-Hep B- IPV (Pediarix) 4mos 2dys | 11/12/2012 DTaP-Hep B- IPV (Pediarix) 6mos 3dys | 08/05/2019 IPV (polio) 7yrs 2mos | | |
| MMR | 05/13/2013 MMR 1yr | 08/05/2019 MMRV (ProQuad) 7yrs 2mos | | | | |
| Hepatitis B | 05/10/2012 Hep B (ped/adol) 1dy | 07/10/2012 DTaP-Hep B- IPV (Pediarix) 2mos 1dy | 11/12/2012 DTaP-Hep B- IPV (Pediarix) 6mos 3dys | | | |
| Varicella | 05/13/2013 Varicella (Varivax) 1yr | 08/05/2019 MMRV (ProQuad) 7yrs 2mos | | | | |
| HPV | | | | | | |
| Hepatitis A | 05/13/2013 Hep A (ped/adol) 1yr | 12/09/2013 Hep A (ped/adol) 1yr 7mos | | | | |
| Seasonal Influenza | 11/12/2012 Influenza (Historical) 6mos 3dys | 12/17/2012 Influenza (Historical) 7mos 8dys | 09/13/2013 Influenza (Historical) 1yr 4mos | 10/10/2014 Influenza (Historical) 2yrs 5mos | 10/13/2015 Influenza (Historical) 3yrs 5mos | |
| Meningococcal Conjugate | | | | | | |
| SARS-CoV-2 | | | | | | |

Case Study #2: Winnie Question #1: Which vaccines should Winnie receive today?

- A. Tdap, HPV, MenACWY, Flu, and COVID-19
- B. HPV, MenB, MenACWY, and Tdap
- C. MenB, COVID-19, and HPV
- D. Td, MenACWY, Flu, and COVID-19
- E. None of the above



Question #1: Answer: A

- Tdap
- HPV
- MenACWY
- Flu
- COVID-19



See Addendum for new or updated ACIP vaccine recommendations Table 1 Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2023

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

| Vaccine | Birth | 1 mo | 2 mos | 4 mos | 6 mos | 9 mos | 12 mos | 15 mos | 18 mos | 19–23 mos | 2-3 yrs | 4-6 yrs | 7-10 yrs | 11–12 yrs | 3–15 yrs | 16 yrs | 17-18 yrs |
|--|----------------------------|--------------------------|----------------------|----------------------|------------------------------|---------------------------|--|---|------------------------------|---------------------|--------------|------------------------------|--------------|------------------------------|---------------------------|-------------------------|---------------|
| Hepatitis B (HepB) | 1 st dose | < 2 nd c | dose> | | | | 3 rd dose | | | | | | | | | | |
| Rotavirus (RV): RV1 (2-dose series), RV5 (3-dose series) | | | 1 st dose | 2 nd dose | See Notes | | | | | | | | | | | | |
| Diphtheria, tetanus, acellular pertussis (DTaP <7 yrs) | | | 1ª dose | 2 nd dose | 3 [™] dose | | | ∢ 4 th d | loseÞ | | | 5 th dose | | | | | |
| Haemophilus influenzae type b (Hib) | | | 1 [≈] dose | 2 nd dose | See Notes | | ▲ <u>3</u> rd or 4 See 1 | th dose. Notes | | | | | | | | | |
| Pneumococcal conjugate (PCV13, PCV15) | | | 1≝ dose | 2 nd dose | 3 ^{el} dose | | ∢ 4 th ¢ | dose• | | | | | | | | | |
| Inactivated poliovirus (IPV <18 yrs) | | | 1 st dose | 2 nd dose | • | | 3 rd dose | | | | | 4 th dose | | | | | See Notes |
| COVID-19 (1vCOV-mRNA, 2vCOV-mRNA, 1vCOV-aPS) | | | | | | | | | 2- or 3- | dose primar | y series and | booster (Se | e Notes) | $\overleftarrow{\mathbf{x}}$ | | | |
| Influenza (IIV4) | | | | | | | | Annual vac | cination 1 o | r 2 doses | | | | Annua | vaccinatio | in 1 dose oi | nly |
| Influenza (LAIV4) | | | | | | | | | | | Ann | ual vaccinat I or 2 doses | ion | Annu | il vaccinati | on 1 dose c | only |
| Measles, mumps, rubella (MMR) | | | | | See I | Notes | 4 1* c | dose> | | | | 2 nd dose | | | | | |
| Varicella (VAR) | | | | | | | 4 1* c | lose• | | | | 2 nd dose | | | | | |
| Hepatitis A (HepA) | | | | | See I | Notes | | 2-dose serie | es, See Note | 5 | | | | | | | |
| Tetanus, diphtheria, acellular pertussis (Tdap ≥7 yrs) | | | | | | | | | | | | | | | | | |
| Human papillomavirus (HPV) | | | | | | | | | | | | | 55 | NA N | | | |
| Meningococcal (MenACWY-D ≥9 mos, MenACWY-CRM ≥2 mos, MenACWY-TT ≥2years) | | | | | | · · · | | See Notes | | | | | IT TO THE | יאלי | | 2 nd dose | |
| Meningococcal B (MenB-4C, MenB-FHbp) | | | | | | | | | | | | | | 1 | See No | tes | |
| Pneumococcal polysaccharide (PPSV23) | | | | | | | | | | | | | | See Notes | | | |
| Dengue (DEN4CYD; 9-16 yrs) | | | | | | | | | | | | | | Seroposit dengue a | ve in ende eas (See No | mic otes) | |
| Range of recommended ages for all children | Range of re for catch-u | ecommend ip vaccinati | ed ages on | Ran | nge of recor certain high | mmended a h-risk group | iges is | Recommended Recommende Recommended Recommended | mended vao jin in this ag | cination e group | R | ecommende n shared clin | d vaccinatio | on based making | No | recommer t applicabl | ndation/ e |
| | | | | | | | | | 10.7 | | | | | | | | |

<u>0-18yrs-child-combined-schedule.pdf (cdc.gov)</u>

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

Notes

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.

<u>0-18yrs-child-combined-schedule.pdf (cdc.gov)</u>

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:

| • P a | and an all a second second second a second beauting and and and a second s | - | | | | | | | |
|--------------|--|----------------|--|--|--|--|--|--|--|
| • Tr | MenB is shared clinical decision- | e 2 | | | | | | | |
| Wa | making for adolescents aged 16 | e 3 | | | | | | | |
| nc aft | through 23 years that are not at | ionths st | | | | | | | |
| 41 | increased risk. For High-Risk | | | | | | | | |
| Not | patients see Immunization | ble; | | | | | | | |
| the For | Schedule footnotes. | es. clictod | | | | | | | |
| und | For and in an outbreak setting and | | | | | | | | |

additional meningococcal vaccination information, see www.cdc.gov/mmwr/volumes/69/rr/rr6909a1.htm.

| Vaccine Type | Brand Names | Routine Recommendation for People Who Are Not in a Risk Group ^{1,2,3} | General Guidelines |
|----------------------|---|--|---|
| MenACWY ² | Menactra® MenQuadfi® Menveo® | One dose at age 11-12 years; booster dose at age 16 years | 1 dose required⁴ at 11 years of age or older upon entry into 7th grade or higher Products are interchangeable if age appropriate, but same vaccine is recommended for complete series Intramuscular injection |
| MenB ³ | Bexsero [®] Trumenba [®] | Shared clinical decision making for persons 16 years through 23 years without high-risk conditions: Bexsero: 2 dose series at least 1 month apart OR Trumenba: 2 dose series at least 6 months apart | Products are not inter- changeable Intramuscular injection MenB vaccine is not routinely recommended at 11-12 years of age |

¹ For Child and Adolescent Immunization Schedule by Age at: <u>www.cdc.gov/vaccines/schedules/hcp/imz/child-adolescent.html</u> ² For more information regarding Meningococcal Conjugate Vaccine (MenACWY), including guidance for catch-up and for persons who are at high-

risk, refer to the Meningococcal Conjugate Quick Look at: <u>www.michigan.gov/vaccinequicklooks</u> and Immunize.org's handout titled, "Meningococcal ACWY vaccine recommendations by Age and Risk Factor" at: <u>www.immunize.org/catq.d/p2018.pdf</u>

³ For more information regarding Meningococcal Serogroup B (MenB), including guidance for catch-up and for persons who are at high-risk, refer to the Meningococcal Serogroup B Quick Look at: <u>www.michigan.gov/vaccinequicklooks</u> and Immunize.org's handout titled, "Meningococcal B Vaccine Recommendations by Age and Risk Factor" at: <u>www.immunize.org/catg.d/p2035.pdf</u>

⁴ For vaccines required for school entry in Michigan, refer to "Vaccines Required for School Entry in Michigan" at: <u>https://mcir.org/wp-</u> content/uploads/2021/08/SchoolEntryRegVaccinesParentsMI 5.3.2021approvedfinalpublish.pdf

Routine-Recommendations-for-Meningococcal-Vaccines_5-25-23_FINAL.pdf (michigan.gov)

| Ages 5–11 years⁺ | | | Points to consid • For a complet the Quick Loc • Egg-ba • Q | der for the 2023-24 Influenza Season te list of contraindications and precautions for influenza vaccines review oks for: |
|---|---|--|---|---|
| COVID-19 vaccination history prior to updated (2023–2024 Formula) mRNA vaccine* | Updated (2023– 2024 Formula) mRNA vaccine | Number of updated (2023–2024 Formula mRNA doses indicate | cclIV4: clIV4: ⊆ Influenza va Inactiva Recom Live Att Cell Cu Severe aller, component With the exagrowing viru | 2023-24 INTIGENZA VACCINATION FOR Persons Who Report Egg Allergy For the 2023-24 influenza season, the Advisory Committee on Immunization Practices (ACIP) recommends the following: 1. All persons aged 6 months and older with egg allergy should receive influenza vaccine |
| Unvaccinated | Moderna | 1 | Tolerance tc For clinics th for maintain how to use i Emerge | Any influenza vaccine (egg based or nonegg based) that is otherwise appropriate for the recipient's age and health status can be used (i.e., any IIV4, RIV4, or LAIV4) Egg allergy in and of itself necessitates no additional safety measures for influenza vaccination beyond those recommended for any recipient of any vaccine, regardless of severity of previous reaction to egg |
| | Pfizer-BioNTech | 1 | airway Refer t at <u>http</u> For persons doses of infl flu product | Severe and life-threatening reactions to vaccines can rarely occur with any vaccine and in any vaccine recipient, regardless of allergy history. All vaccines should be administered in settings in which personnel and equipment needed for rapid recognition and treatment of acute hypersensitivity reactions are available |
| 1 or more doses any mRNA | Moderna OR | 1 | No post-vac persons; ho minutes afte syncope occ | All vaccination providers should be familiar with their office emergency plan and be certified in cardiopulmonary resuscitation (CPR) Remember: It is important to screen and review the contraindications and precautions for any vaccine. With flu vaccine it is important to know the type of flu vaccine being administered to assess for vaccine specific contraindications and precautions. |
| | Pfizer-BioNTech | 1 | Prevention and Contr Immunization Practic | For further information on contraindications and precautions review the Quick Looks for Influenza Vaccines (IIV4, LAIV4, ccIIV4, and RIV4) at: <u>www.Michigan.gov/vaccinequicklooks</u> |
| Interim Clinical Considerati | ons for Use of COVII | D-19 Vaccines CDC | 25, 2023 / 72(2);1–25 regarding flu vaccinat Mich | Prevention and Control of Seasonal Influenza with Vaccines: Recommendations of the Advisory Committee on Immunization Practices — United States, 2023–24 Influenza Season Recommendations and Reports / August 25, 2023 / 72(2);1–25, located at www.cdc.gov/season Recommendations and Reports / August 25, 2023 / 72(2);1–25, located at www.cdc.gov/vaccines/hcp/acip-recs/index.html. For further information regarding flu vaccination, refer to www.ddc.gov/vaccines, or www.cdc.gov/vaccines, or <a h<="" td=""> |

Case Study #2: Winnie Question #2:

When does Winnie need to return and what vaccines will she need at that time?

- A. 1 month: MenACWY, HPV, and Flu
- B. 1 month: MenB, and HPV
- C. 6 months: HPV
- D. 4-8 weeks: HPV, and COVID-19
- E. None of the above

Answer: C

• 6 months: HPV



WINNIE'S MCIR RECORD:

| | | Immunizatio | | | Other | |
|----------------------------|---|--|--|--|--|--|
| Series | Dose 1 | Dose 2 | Dose 3 | Dose 4 | Dose 5 | Dose 6+ |
| DTP/DTaP/ DT/Td/Tdap | 07/10/2012 DTaP-Hep B- IPV (Pediarix) 2mos 1dy | 09/11/2012 DTaP-Hep B- IPV (Pediarix) 4mos 2dys | 11/12/2012 DTaP-Hep B- IPV (Pediarix) 6mos 3dys | 09/09/2013 DTaP (pediatric) 1yr 4mos | 08/05/2019 Tdap (adol/adult) 7yrs 2mos | 09/26/2023 Tdap (adol/adult) 11yrs 4mos |
| Polio | 07/10/2012 DTaP-Hep B- IPV (Pediarix) 2mos 1dy | 09/11/2012 DTaP-Hep B- IPV (Pediarix) 4mos 2dys | 11/12/2012 DTaP-Hep B- IPV (Pediarix) 6mos 3dys | 08/05/2019 IPV (polio) 7yrs 2mos | | |
| MMR | 05/13/2013 MMR 1yr | 08/05/2019 MMRV (ProQuad) 7yrs 2mos | | | | |
| Hepatitis B | 05/10/2012 Hep B (ped/adol) 1dy | 07/10/2012 DTaP-Hep B- IPV (Pediarix) 2mos 1dy | 11/12/2012 DTaP-Hep B- IPV (Pediarix) 6mos 3dys | | | |
| Varicella | 05/13/2013 Varicella (Varivax) 1yr | 08/05/2019 MMRV (ProQuad) 7yrs 2mos | | | | |
| HPV | 09/26/2023 HPV9 11yrs 4mos | | | | | |
| Hepatitis A | 05/13/2013 Hep A (ped/adol) 1yr | 12/09/2013 Hep A (ped/adol) 1yr 7mos | | | | |
| Seasonal Influenza | 11/12/2012 Influenza (Historical) 6mos 3dys | 12/17/2012 Influenza (Historical) 7mos 8dys | 09/13/2013 Influenza (Historical) 1yr 4mos | 10/10/2014 Influenza (Historical) 2yrs 5mos | 10/13/2015 Influenza (Historical) 3yrs 5mos | 09/26/2023 Influenza IIV4 (P-Free Inj) 11yrs 4mos |
| Meningococcal Conjugate | 09/26/2023 MCV4 (Menactra or Menveo) 11yrs 4mos | | | | | |
| SARS-CoV-2 | 09/26/2023 COVID-19 PFR-BNT (23- 24) 10mcg 11yrs 4mos | | | | | |

WINNIE'S MCIR RECORD:

| | | Immunizatio | JIIS | | | other | |
|----------------------------|---|--|--|--|--|--|--------------------------------------|
| Series | Dose 1 | Dose 2 | Dose 3 | Dose 4 | Dose 5 | Dose 6+ | Status |
| DTP/DTaP/ DT/Td/Tdap | 07/10/2012 DTaP-Hep B- IPV (Pediarix) 2mos 1dy | 09/11/2012 DTaP-Hep B- IPV (Pediarix) 4mos 2dys | 11/12/2012 DTaP-Hep B- IPV (Pediarix) 6mos 3dys | 09/09/2013 DTaP (pediatric) 1yr 4mos | 08/05/2019 Tdap (adol/adult) 7yrs 2mos | 09/26/2023 Tdap (adol/adult) 11yrs 4mos | Up-To-Date Next Due 09/26/2033 |
| Polio | 07/10/2012 DTaP-Hep B- IPV (Pediarix) 2mos 1dy | 09/11/2012 DTaP-Hep B- IPV (Pediarix) 4mos 2dys | 11/12/2012 DTaP-Hep B- IPV (Pediarix) 6mos 3dys | 08/05/2019 IPV (polio) 7yrs 2mos | | | Series Complete |
| MMR | 05/13/2013 MMR 1yr | 08/05/2019 MMRV (ProQuad) 7yrs 2mos | | | | | Series Complete |
| Hepatitis B | 05/10/2012 Hep B (ped/adol) 1dy | 07/10/2012 DTaP-Hep B- IPV (Pediarix) 2mos 1dy | 11/12/2012 DTaP-Hep B- IPV (Pediarix) 6mos 3dys | | | | Series Complete |
| Varicella | 05/13/2013 Varicella (Varivax) 1yr | 08/05/2019 MMRV (ProQuad) 7yrs 2mos | | | | | Series Complete |
| HPV | 09/26/2023 HPV9 11yrs 4mos | | | | | | Up-To-Date Next Due 03/26/2024 |
| Hepatitis A | 05/13/2013 Hep A (ped/adol) 1yr | 12/09/2013 Hep A (ped/adol) 1yr 7mos | | | | | Series Complete |
| Seasonal Influenza | 11/12/2012 Influenza (Historical) 6mos 3dys | 12/17/2012 Influenza (Historical) 7mos 8dys | 09/13/2013 Influenza (Historical) 1yr 4mos | 10/10/2014 Influenza (Historical) 2yrs 5mos | 10/13/2015 Influenza (Historical) 3yrs 5mos | 09/26/2023 Influenza IIV4 (P-Free Inj) 11yrs 4mos | Season Complete |
| Meningococcal Conjugate | 09/26/2023 MCV4 (Menactra or Menveo) 11yrs 4mos | | | | | | Up-To-Date Next Due 05/09/2028 |
| SARS-CoV-2 | 09/26/2023 COVID-19 PFR-BNT (23- 24) 10mcg 11yrs 4mos | | | | | | Up-To-Date |
| | | | | | | | |



Know Your Adult Resources and Where to Find Them

2023 Recommended Immunization Schedule For Adults

| addition to the recomm ctober 20, 2022. The foll orbidity and Mortality W | n dations oring reco | presented in the prev mmendations have be t (MMWR). | ous sectio en adopte | ns of this Immunization Schedule, ACII d by the CDC Director and are now off | P has approved the following recomme ficial. Links are provided if these recom | endations by majority vote since mendations have been published in | | | | | |
|---|---|---|-------------------------|---|---|--|--|------------------------|--|--|--|
| /accines | | Table 2 | Recomr | nended Adult Immunizati | on Schedule by Medical C | ondition or Other Indicat | ion, United States, 2023 | | | | |
| lespiratory syncytial virus RSV) | laternal asonal | Vaccine | Pregna | ncy compromised percentage (excluding HIV infection) <15% or | tion CD4 Asplenia, and count complement ≥15% and deficiencies hemodia | age Heart or al lung disease; disease alvois alcoholism ⁴ | Diabetes Health care personnel ⁴ Men v personnel ⁴ with n | rho sex sen | | | |
| | | COVID-19 | | Table 1 Recommen | See Addendum Ided Adult Immunization | n for new or updated ACIP va Schedule for ages 19 year | ccine recommendations s or older, United States, | 2023 | | | |
| OVID-19 (Moderna, fizer-BioNTech) | Il persor valent n | IIV4 or RIV4 | | Vaccine | 19-26 years | 27-49 years | 50-64 years | ≿65 years | | | |
| | • or detail | Tdap or Td | 1 dose T preg | Influenza inactivated (IIV4) or | | 2- or 3- dose prima | ary series and booster (see notes) | | | | |
| tespiratory syncytial | duits 60 | MMR | Contrain | Influenza recombinant (RIV4) | 1 dose annually | | | | | | |
| 100 (101) | - or detain | VAR | Contrain | (LAIV4) | 1 dose annually 1 dose Td/Tdap for wound management (see notes) | | | | | | |
| | dults where the second se | RZ V | | (Tdap or Td) | | 1 dose Tdap, then Td or Td | ap booster every 10 years | iona) | | | |
| oliovirus (IPV) | dults wh cposure | HPV Pneumococcal | Recomm | Measles, mumps, rubella (MMR) | | ing on indication 57 or later) | For healthcare personne see notes | | | | |
| | . I persor | PPSV23) HepA | | Varicella (VAR) | 2 doses (if born in 1980 o | or later) | 2 doses | | | | |
| nfluenza (IIV4, ccIV4, | at is oth | НерВ | 3 di (see r | Zoster recombinant (RZV) | 2 doses for immunocomprom | ising conditions (see notes) | 2 di | oses | | | |
| .,, | ecomme ww.cdc | MenACWY | | Human papillomavirus (HPV) | 2 or 3 doses depending on age at initial vaccination or condition | 27 through 45 years | | | | | |
| e effective date is the date | nen the C | MenB | Preca | Pneumococcal (PCV15, PCV20, PPSV23) | | 1 dose PCV15 followed by OR 1 dose PCV20 (see no | PPSV23 | See Notes See Notes | | | |
| | _ | Hib Recommended v | ccination | Hepatitis A (HepA) | | 2, 3, or 4 doses dep | ending on vaccine | | | | |
| | | for adults who m age requirement, documentation o vaccination, or la | wt fack k | Hepatitis B (HepB) | | 2, 3, or 4 doses dependin | g on vaccine or condition | | | | |
| | | evidence of past | nfection en not epp | Meningococcal A, C, W, Y (MenACWY) | 1 or 2 | doses depending on indication, s | ee notes for booster recommendat | tions | | | |
| | | | | Meningococcal B (MenB) | 2 or 3 dose 19 through 23 years | es depending on vaccine and indice | ation, see notes for booster recom | mendations | | | |
| | | | | Haemophilus influenzae type b | | 1 or 3 doses depen | ding on indication | | | | |

- Adult vaccination is based primarily on risk conditions
- Schedule notes include information on risk groups, minimum and recommended intervals
- Addendum includes new or updated ACIP vaccine recommendations

2023 (CDC) Adult Immunization 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 prim | ary series and booster (See Notes) | |
| Influenza inactivated (IIV4) or Influenza recombinant (RIV4) | | | | |
| Influenza live, attenuated (LAIV4) | | ecommended vaccination for aduling requirement, lack documentation | ts who meet tion of | |
| Tetanus, diphtheria, pertussis (Tdap or Td) | v | accination, or lack evidence of pa | st infection | ee notes) |
| Measles, mumps, rubella (MMR) | = Re | ecommended vaccination for adul | ts with an | For healthcare personnel, see notes |
| Varicella (VAR) | a | dditional risk factor or another in | dication 2 do | ses |
| Zoster recombinant (RZV) | 2 dos = Re | commended vaccination based o | n shared clinical | 2 doses |
| Human papillomavirus (HPV) | 2 or 3 dose dose dose dose dose dose dose dose | lecision-making | | |
| Pneumococcal (PCV15, PCV20, PPSV23) | = Nc | o recommendation/Not applicable | e | See Notes See Notes |
| Hepatitis A (HepA) | | | | |
| Hepatitis B (HepB) | | 2, 3, or 4 doses dependin | ig on vaccine or condition | |
| Meningococcal A, C, W, Y (MenACWY) | | 1 or 2 deses depending on indication, s | ee notes for booster recommen | dations |
| Meningococcal B (MenB) | 2 or 3 19 through 23 years | 3 doses epending on vaccine and indic | ation, see notes for booster reco | ommendations |
| Haemophilus influenzae type b (Hib) | | 1 or 3 doses depen | ding on indication | |
| Recommended vaccination for adults lack documentation of vaccination, o | s who meet age requirement, or lack evidence of past infection | Recommended vaccination for adults with an additional risk factor or another indication | Recommended vaccination bas clinical decision-making | ed on shared No recommendation/ Not applicable |
| | Decomposed ad Adu | lt henre verige tiger. Cele e dude i theite d | Chatas 2022 (ada sau) | |

Recommended Adult Immunization Schedule, United States, 2023 (cdc.gov)
Table 2 Recommended Adult Immunization Schedule by Medical Condition or Other Indication, United States, 2023

| | | | | | | | Men who | | |
|---|------------------|---|---------------------------|---|--|-------------------------------|----------------------|--|--|
| Vaccine | Pregna | = Recommended vaccination for adult who meet age | Ch | disease | Diabetes | personnel ^b | have sex with men | | |
| | | requirement, lack documentation of vaccination, or lac | ck 🚽 | | | | | | |
| COVID-19 | | evidence of past infection | | | | | | | |
| IIV4 or RIV4 | | | | | | | | | |
| LAIV4 | | = Recommended vaccination for adults with an additional ris | sk <mark>tio</mark> | 'n | | 1 dose a | nnually | | |
| Tdan or Td | 1 dose Tda | factor or another indication | | | | | | | |
| Tuap of Tu | pregna | | , | years | | | | | |
| MMR | Contraindi | = Recommended vaccination based on share clinical decision | n- <mark>ng</mark> | on indicatio | on | | | | |
| VAR | Contraindi | making | 2 0 | 2 doses | | | | | |
| RZV | | = Precaution-vaccination might be indicated if benefit of | ata | age ≥50 yea | irs | | | | |
| HPV | Not Recomme | protection outweighs risk of adverse reaction | end | ending on age at initial vaccination or condition | | | | | |
| Pneumococcal (PCV15, PCV20, PPSV23) | | - Contraindicated or not recommended-vaccine should not k | PS PS | V23 OR 1 de | ose PCV20 (s | ee notes) | | | |
| НерА | | administered | 0C | es dependin | ig on vaccine | RE. | | | |
| НерВ | 3 dos (see no | *vaccinate after pregnancy | cor | ndition | | | | | |
| MenACWY | | - No recommendation (not applicable | | | | | | | |
| MenB | Precau | | bo | oster recon | nmendations | | | | |
| Hib | L | recipients only | | | | | | | |
| Recommended vaccination for adults who meet age requirement, lack documentation of vaccination, or lack evidence of past infection | | | Con reco sho *Va | ntraindicated or ommended-va ould not be adm occinate after pr | not ccine inistered. egnancy. | No recommen Not applicable | dation/ | | |

Notes **Recommended Adult Immunization Schedule, Unit**

 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)

 HPV vaccination recomm through age 26 years: 2- 0 on age at initial vaccinatio

Age 15 years or older at 3-dose series at 0, 1-2 mor intervals: dose 1 to dose 2 12 weeks / dose 1 to dose administered too soon)

Age 9–14 years at initial 1 dose or 2 doses less that 1 additional dose

Age 9–14 years at initial 2 doses at least 5 months series complete, no additi

 Interrupted schedules: If interrupted, the series doe

 No additional dose recom vaccine series has been co recommended dosing int

Shared clinical decision-

 Some adults age 27–45 years clinical decision-making, 2

Special situations

 Age ranges recommended catch-up vaccination or sh making also apply in spec

- Immunocompromising infection: 3-dose series, e vaccination at age 9 throu

- Pregnancy: Pregnancy te vaccination; HPV vaccinat until after pregnancy; no inadvertently vaccinated

See Addendum for new or updated ACIP vaccine recommendations 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-19immunization-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

 Hematopoietic stem cell transplant (HSCT): 3-dose series 4 weeks apart starting 6-12 months Hib vaccination history

Hepatitis A vaccination

(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

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)

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.

14 days before splenectomy

after successful transplant, regardless of

Routine vaccination

Not at risk but want protection from hepatitis A

- Chronic liver disease (e.g., persons with

HIV infection

Men who have sex with men

Injection or noninjection drug use

Persons experiencing homelessness Work with hepatitis A virus in research

Human papilloma **Routine vaccination**

Adult Immunization Schedule by Age

Print

Recommendations for Ages 19 Years or Older. United States. 2023 Addendum – Adult Recommended Immunization Schedule for ages 19 years or older, United States, 2023

| See Addendum for new or updat | Vaccines | | Recommendations | Effective Date of Recommendation* | |
|---|-------------------------|---|--|---|---|
| View addendum | Respiratory syncytial v | Addendu | Recommended Adult Immuniz | ation Schedule, United States, 2023 | |
| | | In addition to the record October 20, 2022. The Morbidity and Mortality | mmendations presented in the previous sections of following recommendations have been adopted by y Weekly Report (MMWR). | this Immunization Schedule, ACIP has approved the following recommendations by the CDC Director and are now official. Links are provided if these recommendations | majority vote since have been published in |
| Using the schedule | | Vaccines | Recommendations | | Effective Date of Recommendation* |
| To make vaccination recommendations | | COVID-19 (Moderna, Pfizer-BioNTech) | All persons ≥6 months of age should receive 2023–2024 For detailed information, see: www.cdc.gov/covidsched | 4 (monovalent, XBB containing) COVID-19 vaccines as authorized under EUA or approved by BLA. Jule | September 12, 2023 |
| Determine needed vaccines based of Assess for medical conditions and of | COVID 10 (Madaraa B | Respiratory syncytial virus (RSV) | Adults 60 years of age and older may receive a single do For detailed information, see: www.cdc.gov/mmwr/volu | ose of Respiratory Syncytial Virus (RSV) vaccine, using shared clinical decision-making. umes/72/wr/mm7229a4.htm?s_cid=mm7229a4_w | June 27, 2023 |
| Review special situations (Vaccination) Review contraindications and precations See Addendum for new or updated | COVID-19 (Moderna, P | Poliovirus (IPV) | Adults who are known or suspected to be unvaccinated vaccination series with inactivated polio vaccine (IPV). Adults who have received a primary series of trivalent o exposure may receive another dose of IPV. Available dated | d or incompletely vaccinated against polio should complete a primary ral polio vaccine (tOPV) or IPV in any combination and who are at increased risk of poliovirus ta do not indicate the need for more than a single lifetime booster dose with IPV for adults. | June 27, 2023 |
| | | Influenza (IIV4, cclV4, RIV4, LAIV4) | All persons ages ≥6 months with egg allergy should rect that is otherwise appropriate for the recipient's age and Affirm the updated MMWR Recommendations and Rep Recommendations of the Advisory Committee on Immu www.cdc.gov/mmwr/volumes/72/rr/rr7202a1.htm | reive influenza vaccine. Any influenza vaccine (egg based or non-egg based) I health status can be used. orts, "Prevention and Control of Seasonal Influenza with Vaccines: unization Practices—United States, 2023-24 Influenza Season" | June 27, 2023 |
| | | *The effective date is the d | late when the CDC director adopted the recommendation an | d when the ACIP recommendation became official | |
| | A | dult Immuniza | ation Schedule – Healthcare | e Providers CDC | |

General Best Practice Guidelines for Immunization

Updated August 1, 2023

Best Practices Guidance Kroger A, Bahta L, Long S, Sanchez P

Introduction Purpose and topics covered in this report...

History

History of development of: Timing and Spacing, Contraindications and Precautions, Preventing and Managing Adverse Reactions...

Timing and Spacing of Immunobiologics

Vaccine scheduling, supply and lapsed schedule, spacing of doses, simultaneous and nonsimultaneous administration, licensed combination vaccines, interchangeability of formulations, extra doses, conjugate vaccines...

Contraindications and Precautions

General principles, standards of valid contraindications and precautions, and conditions incorrectly perceived as contraindications...

Preventing and Managing Adverse Reactions

Benefit and risk communication, reporting adverse reactions, National Vaccine Injury Compensation Program...

Vaccine Administration

Infection control and sterile technique, route of administration, multiple and jet injections. alleviating discomfort and pain, clinical implications of nonstandard practices...

Altered Immunocompetence

General Best Practice Guidelines for Immunization

Print

Updated August 1, 2023

Printer friendly version 📕 [26 pages]

Updates

This section incorporates general content from the Infectious Diseases Society of America policy statement, 2013 IDSA Clinical Practice Guideline for Vaccination of the Immunocompromised Host (<u>1</u>), to which CDC provided input in November 2011. The evidence supporting this guidance is based on expert opinion and arrived at by consensus.

ACIP Altered Immunocompetence Guidelines for Immunizations | CDC

COVID-19 Vaccine Clinical Considerations

| Interim Clinical Considerations for Us Vaccines in the United States | e o | f COVID-19 Ages 12 years and older | | | | | |
|--|--------------------------|--|---|--|--------------------------------------|---|-------------------------------------|
| Print Summary of recent changes (last updated October 24, 2023): | | COVID-19 vaccination history prior to updated (2023–2024 Formula) mRNA vaccine* | Updated (2023– 2024 Formula) mRNA vaccine | Number of updated (2023–2024 Formula) mRNA doses indicated | Dosage (mL/ug) | Vaccine vial cap and label colors [§] | Interval between doses |
| Age transitions: Updated guidance for children who transition during the initial 4 years to age 5 years and children who are moderately or severely immunocol years to age 12 years to receive the age-appropriate dosage based on their age | Unvaccinated | Moderna 1 0.5 Da mL/50 ug ca | | Dark blue cap; blue label | _ | | |
| Interchangeability of COVID-19 vaccines: Clarification of circumstances in which doses from different manufacturers may be considered when doses from the s recommended. | 1 or more doses any mRNA | Pfizer-BioNTech Moderna | 1 | 0.3 mL/30 ug 0.5 | Gray cap; gray label Dark blue | At least 8 weeks | |
| | | | OR | | mL/50 ug | cap; blue label | after last dose |
| Reference Materials | Ge | | Pfizer-BioNTech | 1 | 0.3 mL/30 ug | Gray cap; gray label | At least 8 weeks after last dose |
| <u>COVID-19 Vaccination Recommendations Infographic</u> (Updated 10/13/2023) <u>COVID-19 Vaccination Recommendations Infographic</u> (<u>Immunocompromised</u>) (Updated 10/13/2023) | Re pa Wł | 1 or more doses Novavax or Janssen, including in combination with any mRNA vaccine dose(s) | Moderna OR | 1 | 0.5 mL/50 ug | Dark blue cap; blue label | At least 8 weeks after last dose |
| | | | Pfizer-BioNTech | 1 | 0.3 mL/30 ug | Gray cap; gray label | At least 8 weeks after last dose |

Interim Clinical Considerations for Use of COVID-19 Vaccines | CDC



Adult Case Study



Case Study #3: Buzz DOB: 2-4-56 Age: 67 years:

- Buzz is a 67-year-old patient with diabetes
- Has a history of chickenpox disease
- Has a winter home in Mexico
- Does not work in healthcare
- He has insurance that covers his immunizations

BUZZ'S MCIR RECORD:

| Series | Dose 1 | Dose 2 | Dose 3 | Dose 4 | Dose 5 | Dose 6+ |
|-------------------------|---|---|---|---|---|---------|
| DTP/DTaP/ DT/Td/Tdap | 06/10/1980 Td PF (adol/adult) 24yrs 4mos | 08/11/1980 Td PF (adol/adult) 24yrs 6mos | | | | |
| MMR | 08/20/1961 MMR 5yrs 6mos | | | | | |
| Hepatitis B | | | | | | |
| Hepatitis A | | | | | | |
| Seasonal Influenza | 08/10/2018 Influenza (Historical) 62yrs 6mos | 10/07/2019 Influenza (Historical) 63yrs 8mos | 10/15/2020 Influenza (Historical) 64yrs 8mos | 11/01/2021 Influenza (Historical) 65yrs 8mos | 10/21/2022 Influenza IIV4 High Dose (FluZone HD Quad) 66yrs 8mos | |
| Pneumococcal Adult | 08/10/2018 PPSV23 (Pneumovax) 62yrs 6mos | | | | | |
| SARS-CoV-2 | 09/19/2022 COVID-19 PFR Bivalent 30mcg/0.3mL 66yrs 7mos | | | | | |
| Zoster | 08/09/2019 Zoster ZVL (Zostavax) 63yrs 6mos | | | | | |
| RSV | | | | | | |
| Other Admin | istrations | | | | | |
| Series | | | | | | |
| Varicella | 08/09/2019 Zoster ZVL (Zostavax) 63yrs 6mos | | | | | |
| Dispensed V | accines / Bio | logics | | | | |
| Vaccine/Biolog | ic | | | | Date | Age |
| No Dispensed | Vaccines or B | iologics Found | 1 | | | |
| Non-Adminis | stered Doses | /Positive Im | munity | | | |
| Series/Antigen | | Date | Reas | on | Entered by | |
| Varicella | | 04/02/1962 | Immu | nity | MDHHS Nurse Ed | ucators |

Case Study #3: Buzz Question #1:

Which vaccines are recommended for Buzz to receive today?

- A. Td, Hep A, Flu, COVID-19, Shingrix (Zoster), Respiratory Syncytial Virus (RSV)
- B. Td, Hep A, Hep B, Flu, COVID-19
- C. Tdap, Hep B, Hep A, Flu, Pneumococcal, COVID-19, Shingrix (Zoster), RSV
- D. Tdap, Hep B, COVID-19, Flu, RSV
- E. None of the above

Question #1: Answer: C

- Tdap
- Hep B
- Hep A
- Flu
- Pneumococcal
- COVID-19
- Shingrix (Zoster)
- RSV



Discussion of answer

- Tdap needed
- 2nd MMR not recommended
- Hep B: Age 60 years or older with known risk factors for hepatitis B virus infection should complete a Hep B vaccine series
- Hep A: Winter home in Mexico, is recommended to have protection
- Flu*: is yearly and CDC prefers for those 65 years and older to get:
 - \circ HD-IIV4 **OR**
 - o allV4 OR
 - o RIV4
- *If none of these are available, give any flu vaccine that is age and dose appropriate
 - Pneumococcal:
 - One dose of PCV20 at least 1 year after the last PPSV23 dose, OR
 - $\circ~$ One dose of PCV15 at least 1 year after the last PPSV23
 - RSV: is a shared clinical decision-making discussion
 - COVID-19: Only needs one dose of COVID-19 2023-24 formula
 - Shingrix (Zoster): whole series needs to be administered even though he had a Zostavax

adult-combined-schedule.pdf (cdc.gov)



| | Hepatitis B vaccin | ation | |
|---|--|---|--|
| Hepatitis A vaccination Routine vaccination Not at risk but want protection from hepatitis A | Age 60 years or older with known hepatitis B virus infection should HepB vaccine series. Age 60 years or older without for hepatitic B virus infection methods. | own risk factors for d complete a known risk factors | |
| (identification of risk factor not required): 2-dose series HepA (Havrix 6–12 months apart or | HepB vaccine series. | Measles, mur | nps, and rubella vaccination |
| 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]) | | Routine vaccination No evidence of improve rubella: 1 dose Evidence of improve of imp | on munity to measles, mumps, or unity: Born before 1957 (health |
| Travel in countries with high or intermediate endemic hepatitis A (HepA-HepB [Twinrix] ma be administered on an accelerated schedule of 3 doses at 0, 7, and 21–30 days, followed by a booster dose at 12 months) | e Iy | care personnel, se of MMR vaccine, la or disease (diagno confirmation is no | e below), documentation of receipt aboratory evidence of immunity osis of disease without laboratory ot evidence of immunity) |

adult-combined-schedule.pdf (cdc.gov)

 COVID-19 vaccination recommendations have changed. Find the latest recommendations at www.cdc.gov/covidschedule

 Table 1
 Recommended Adult Immunization Schedule for ages 19 years or older, 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 annually | | | | |
| Influenza live, attenuated (LAIV4) | | 1 dose annu | ually | | | |
| Tetanus, diphtheria, pertussis (Tdap or Td) | 1 dose | e Tdap each pregnancy; 1 dose 1 dose Tdap, then Td | e Td/Tdap for wound management (see i or Tdap booster every 10 years | notes) | | |
| Measles, mumps, rubella (MMR) | | 1 or 2 doses de (if born | pending on indication in 1957 or later) | For healthcare personnel, see notes | | |
| Varicella (VAR) | 2 doses (if born in 1980) | or later) | 2 doses | | | |
| Zoster recombinant (RZV) | 2 doses for immunocompromising conditions (see notes) 2 dos | | | oses 🛠 | | |
| Human papillomavirus (HPV) | 2 or 3 doses depending on age at initial vaccination or condition | 27 through 45 years | | | | |
| Pneumococcal (PCV15, PCV20, PPSV23) | | See Notes | | | | |
| Hepatitis A (HepA) | | 2, 3, or 4 dose | s depending on vaccine | \bigstar | | |
| Hepatitis B (HepB) | | 2, 3, or 4 doses depe | ending on vaccine or condition | \bigstar | | |
| Meningococcal A, C, W, Y (MenACWY) | 1 or 2 doses depending on indication, see notes for booster recommendations | | | | | |
| Meningococcal B (MenB) | 2 or 3 dos 19 through 23 years | es depending on vaccine and | indication, see notes for booster recom | mendations | | |
| Recommended vaccination for adults | who meet age requirement, | Recommended vaccination for adults v | vith an Recommended vaccination base | ed on shared No recommendation | | |
| lack documentation of vaccination, or | lack evidence of past infection | additional risk factor or another indicat | ion clinical decision-making | Not applicable | | |
| | | | | | | |

adult-combined-schedule.pdf (cdc.gov)



Adults ≥65 years old

Complete pneumococcal vaccine schedules

| Prior vaccines | Option A | Pneumococcal vaccination | For guidance on determining which pneumococcal |
|--------------------------------------|-------------------------------|--|---|
| | •••••• | Routine vaccination | mobile app which can be downloaded here: www.co |
| | | Age 65 years or older who have: | gov/vaccines/vpd/pneumo/hcp/pneumoapp.html |
| None* | PCV20 | 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 | Special situations • Age 19–64 years with certain underlying medical conditions or other risk factors** who have |
| PPSV23 only | | 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. | 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 immunocompromision condition 4 conducts withan |
| at any age | ≥1 year > PCV20 | Previously received only PCV7: follow the recommendation above. | or cerebrospinal fluid leak |
| | V | Previously received only PCV13: 1 dose PCV20 at least 1 year after the PCV13 dose OR complete the | Previously received only PCV7: follow the recommendation above. |
| | | recommended PPSV23 series as described here www.cdc.gov/vaccines/vpd/pneumo/downloads/ pneumo-vaccine-timing.pdf. | Previously received only PCV13: 1 dose PCV20 at least 1 year after the PCV13 dose OR complete the recommended PPSV23 series as described here |
| PCV13 only | STugar DOV20 | Previously received only PPSV23: 1 dose PCV15 OR 1 dose PCV20 at least 1 year after the PPSV23 dose | www.cdc.gov/vaccines/vpd/pneumo/downloads/ pneumo-vaccine-timing.pdf. |
| at any age | 21 year F0420 | If PCV15 is used, it need not be followed by another dose of PPSV23. | • Previously received only PPSV23: 1 dose PCV15 (1 dose PCV20 at least 1 year after the PPSV23 dose. |
| | | Previously received both PCV13 and PPSV23 | If PCV15 is used, it need not be followed by anothe dose of PPSV23. |
| PCV13 at any age & PPSV23 at <65 yrs | ≥5 years PCV20 | 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 | 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/ |
| | | 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 nneumocorcal | pneumo-vaccine-timing.pdf. • For guidance on determining which pneumococcal vaccines a patient needs and when, please refer to t |
| Pneumococca | al Vaccine Timing for Adults | vaccine dose. | mobile app which can be downloaded here: www.cr gov/vaccines/vpd/pneumo/hcp/pneumoapp.html |
| greater than o | r equal to 65 years (cdc.gov) | | a tri tri tri |

Notes Recommended Adult Immunization Schedule, United States, 2023

nd when, please refer to the include chronic renal downloaded here: www.cdc. immunodeficiency, ia o/hcp/pneumoapp.html generalized malignan virus, Hodgkin disease

*Note: Immunocon

vpd/polio/hcp/recor

tain underlying medical actors** who have

a PCV13, PCV15, or us vaccination history 15 OR 1 dose PCV20. If d be followed by a dose of ear after the PCV15 dose. weeks between PCV15 dered for adults with an ondition,* cochlear implant,

PCV13: 1 dose PCV20 at 13 dose OR complete the ries as described here pd/pneumo/downloads/ odf.

PPSV23: 1 dose PCV15 OR ear after the PPSV23 dose. not be followed by another

myeloma, solid organ acquired asplenia, sic hemoglobinopathies. **Note: Underlying n risk factors include al lung disease, chronic cochlear implant, con CSF leak, diabetes me HIV, Hodgkin disease, immunosuppression, myeloma, nephrotic s or sickle cell disease Pc **Routine vaccinatio** Routine poliovirus va United States is not n **Special situations** Adults at increased to poliovirus with: No evidence of a co (i.e., at least 3 dose (1, 2, or 3 doses) to -Evidence of comple (i.e., at least 3 doses **IPV** booster For detailed informat

10:18 AM A LTE 💡 > BINA PneumoRecs VaxAdvisor Tool to help determine which pneumococcal vaccines children and adults need. **Getting Started** Enter a patient's age, pneumococcal vaccination history, and underlying medical conditions. Move through this tool to create customized recommendations for pneumococcal vaccination. Enter Tool î \triangle

PneumoRecs VaxAdvisor is available for download on iOS and Android mobile devices.

PneumoRecs VaxAdvisor: Vaccine Provider App | CDC

Adult Immunization Schedule – Healthcare Providers | CDC



Case Study #3: Buzz (question #1 cont.

After having a shared clinical decision-making discussion with his provider, Buzz decided he would get the following vaccines:

- Tdap
- Hep B (Twinrix)
- Hep A (Twinrix)
- <u>Flu</u>
- Pneumococcal
- RSV
- COVID-19
- Shingrix (Zoster)

ACIP Shared Clinical Decision-Making Recommendations | CDC

BUZZ'S MCIR RECORD:

| | Immunizations | | | | | Other |
|-------------------------|---|--|---|---|---|---|
| Series | Dose 1 | Dose 2 | Dose 3 | Dose 4 | Dose 5 | Dose 6+ |
| DTP/DTaP/ DT/Td/Tdap | 06/10/1980 Td PF (adol/adult) 24yrs 4mos | 08/11/1980 Td PF (adol/adult) 24yrs 6mos | 11/02/2023 Tdap (adol/adult) 67yrs 8mos | | | |
| MMR | 08/20/1961 MMR 5yrs 6mos | | | | | |
| Hepatitis B | 11/02/2023 Hep A-Hep B (Twinrix) 67yrs 8mos | | | | | |
| Hepatitis A | 11/02/2023 Hep A-Hep B (Twinrix) 67yrs 8mos | | | | | |
| Seasonal Influenza | 08/10/2018 Influenza (Historical) 62yrs 6mos | 10/07/2019 Influenza (Historical) 63yrs 8mos | 10/15/2020 Influenza (Historical) 64yrs 8mos | 11/01/2021 Influenza (Historical) 65yrs 8mos | 10/21/2022 Influenza IIV4 High Dose (FluZone HD Quad) 66yrs 8mos | 11/02/2023 Influenza IIV4 High Dose (FluZone HD Quad) 67yrs 8mos |
| Pneumococcal Adult | 08/10/2018 PPSV23 (Pneumovax) 62yrs 6mos | 11/02/2023 PCV20 (Prevnar 20) 67yrs 8mos | | | | |
| SARS-CoV-2 | 09/19/2022 COVID-19 PFR Bivalent 30mcg/0.3mL 66yrs 7mos | 11/02/2023 COVID-19 PFR Comirnaty 2023/24 30mcg 67yrs 8mos | | | | |
| Zoster | 08/09/2019 Zoster ZVL (Zostavax) 63yrs 6mos | 11/02/2023 Zoster RZV (Shingrix) 67yrs 8mos | | | | |
| RSV | 11/02/2023 RSV vaccine (Arexvy) 67yrs 8mos | | | | | |
| Other Admin | nistrations | | | | | |
| Series | | | | | | |
| Varicella | 08/09/2019 Zoster ZVL (Zostavax) 63yrs 6mos | | | | | |

Case Study #3: Buzz Question #2:

When does Buzz need to return for his **next** immunizations and which ones?

- A. 1 month for Hep B and Hep A (Twinrix)
- B. 2-3 months for MMR, RSV, COVID-19
- C. 3-6 months for MMR, Pneumococcal, RSV
- D. 12 months for Shingrix (Zoster)
- E. None of the above



Answer: A. 1 month for HepA/HepB (Twinrix)

- Td/Tdap: booster every 10 years
- Flu: only one flu shot per season
- Pneumococcal: He is complete
- RSV: At this time RSV is only a one dose recommendation
- COVID-19: He only needs 1 dose this season
- Shingrix (Zoster): 2nd and last dose is due
 2-6 months after the first dose

Vaccine Quick Looks (michigan.gov)

Thank you so much for being here today and for everything you do.

If you have any further questions, please contact us at: <u>checcimms@michigan.gov</u>

Stay Up-To-Date on Immunization Recommendations

- Stay up-to-date by joining the MDHHS Listserv and receive email updates
- To sign up, email Dara Barrera at

djbarrera@msms.org and ask to be added to the MDHHS Immunization Listserv Clusters of under vaccination, provider deserts, and preventing the next outbreak of vaccine preventable diseases in Michigan

Ryan Malosh, PhD MPH

Taylor Olsabeck, MS



Agenda



- Maps!
- Provider deserts
 - COVID
 - VFC
- Vaccine Preventable Disease Update
 - Cases
 - Chickenpox
 - H. influenzae outbreak in Detroit

Maps - Fine area estimation



MI Lighthouse





- A collaboration between the MDHHS, UM School of Public Health, UM School of Information, and the UM College of Engineering has created several vital tools used by public health professionals, health professionals, and the public to better understand the symptoms, risks, and available vaccination data throughout Michigan during the COVID-19 pandemic.
- MI Lighthouse is a vaccine visualization and analysis platform developed to provide local, highresolution information on COVID-19 vaccine coverage for Local Health Department staff.
- MI Lighthouse has been used by public health officials to enable data-informed decision-making about vaccine clinics, resource allocation, and response efforts since the introduction of the COVID-19 vaccines.
- The MI Lighthouse team is currently focused on expanding the scope of MI Lighthouse beyond COVID-19 to provide vaccine visualization and analytics for a wide range of vaccines, Starting with MMR.



Michigan Coverage Maps at Different Spatial Scales



- Interactive Mapping tools can be useful for exploring regional and census-tract level variation in MMR and Varicella Coverage at different spatial scales (e.g. county, census tract).
- For school buildings, a data table will appear with enrollment and MMR/VAR coverage from 2017-2021,
- The size of school points is proportional to their median enrollment over the study period, and the color is related to the MMR coverage in 2021.
- As a note tracts in white are censored (meaning denominator of 5 year olds of 10 or fewer children).

Data on MMR Coverage



Summary of Vaccination Coverage from IIS Data

| Year | IIS Denominator | 0-Dose MMR | 1-Dose MMR | 2-Dose MMR | 0-Dose MMR% | 1-Dose MMR% | 2-Dose MMR% |
|------|-----------------|--------------------|------------|----------------------|-------------|-------------|-------------|
| 2017 | 121679 | 507 <mark>1</mark> | 7861 | <mark>10874</mark> 7 | 4.2 | 6.5 | 89.4 |
| 2018 | 122743 | 5465 | 8577 | 108701 | 4.5 | 7.0 | 88.6 |
| 2019 | 124145 | 5051 | 9496 | 109598 | 4.1 | 7.6 | 88.3 |
| 2020 | 123516 | 6059 | 12010 | 105447 | 4.9 | 9.7 | 85.4 |
| 2021 | 122025 | 6944 | 12592 | 102489 | 5.7 | 10.3 | 84.0 |

Table 1. MMR Doses and Coverage - IIS Data, 2017-2021

County level maps



- County level maps are the first level of analysis
- Identifying areas with low coverage
- Allows for broad inferences (i.e. rural vs urban, border vs non-border)



Census tract level maps



MMR Coverage Maps



- Tract level maps can highlight specific areas in a county that have low coverage
- Tracts in high coverage counties have low coverage

Adding school data



- Can also add school building level data for MMR
- Future work will allow for ISD level aggregation

MMR Coverage Maps



Within counties you can see variation





Provider Deserts



COVID Vaccine Provider Deserts



Vaccine desert More than a 15 minute drive to the closest active COVID-19 vaccination site.

OO Potential vaccination sites within deserts To view, see "Find potential new sites" on the menu.



- Work with eTrueNorth to identify independent pharmacies and encourage them to enroll
- Identify opportunities for community outreach
- Work with vaccine champions

Count of VFC providers by county




Michigan counties with low coverage and lesser VFC providers (below median)



VPD Update

Health department: Measles case confirmed in Milwaukee resident

By: CBS 58 Newsroom

Morbidity and Mortality Weekly Report

Posted: Oct 10, 2023 2:12 PM CDT

Measles Outbreak — Central Ohio, 2022–2023

Notes from the Field

Elizabeth C. Tiller, MSPA^{1,2}; Nina B. Masters, PhD¹; Kelley L. Raines, MPH³; Adria D. Mathis, MSPH³; Stephen N. Crooke, PhD³; Rebecca C. Zwickl, MPH²; Gavin K. French²; Emily R. Alexy, MPH²; Elizabeth M. Koch, MD²; Naomi E. Tucker, MPH²; Elizabeth M. Wilson²; Tiffany S. Krauss, MSN²; Erica Leasure, MS⁴; Jeremy Budd⁴; Laurie M. Billing, MPH⁴; Courtney Dewart, PhD⁴⁻⁵; Kara Tarter, MPH⁴; Kristen Dickerson, PhD⁴; Radhika Iyer, MPH⁶; Alexandria N. Jones, MS⁶; Katia C. Halabi, MD⁷; Matthew C. Washam, MD⁷; David E. Sugerman, MD³; Mysheika W. Roberts, MD²

On November 5, 2022, Columbus Public Health, Ohio and the Ohio Department of Health were notified of two children aged 2 years who were admitted to a central Ohio hospital with rash, fever, cough, and congestion, suggestive of measles. Both children were undergoing medical evaluation and treatment for other etiologies before measles was considered in the differential diagnosis. Neither child had received measles, mumps, and rubella (MMR) vaccine, and neither had known contact with a person with measles. Each patient subsequently received a positive measles real-time reverse transcription–polymerase Health Public Health Laboratory performed RT-PCR testing of specimens from 193 persons during the outbreak; 74 (87%) measles cases were laboratory-confirmed,[†] and the remaining 11 (13%) were epidemiologically linked to confirmed cases. Among 65 genotyped specimens, all were genotype B3. The median patient age was 1 year (range = 6 months–15 years). Eighty (94%) patients had not received MMR vaccine. Sixty (71%) patients were aged ≥1 year and age-eligible for routine MMR vaccination,[§] but only three (5%) had documentation of receipt of 1 MMR vaccine dose at the time of infection[¶]; vaccination status of one (2%) patient was unknown.

Forty-four (52%) of the 85 measles patients experienced complications, including otitis media (33; 39%), diarrhea (22; 26%), and pneumonia (seven; 8%); 36 (42%) patients were hospitalized, predominately for dehydration.** The median length of hospitalization was 3 days (range = 1-7 days). Twelve hospitalized patients had coinfections with other respiratory pathogens (e.g., respiratory syncytial virus [RSV]).^{††} No deaths were reported.

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

<u>Print</u>



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

VPD Cases over time



| Disease | Pre-pandemic average (2016) | Total Cases, 2020 | Total Cases, 2021 | Total Cases, 2022 | Cases to Date, 2023 Sept. 1, 2023 |
|---|--------------------------------|----------------------|----------------------|----------------------|---|
| Congenital Rubella | 0 | 0 | 0 | 0 | 0 |
| Diphtheria | 0 | 0 | 0 | 0 | 0 |
| H. influenzae – Invasive in <15 year olds (serotype b) | 23.5 | 11 (1) | 14 (1) | 24 (0) | 13 (1) |
| Measles | 17 | 0 | 1 | 4 | 0 |
| Meningococcal disease | 5.75 | 3 | 6 | 5 | 6 |
| Mumps | 30 | 4 | 5 | 15 | 11 |
| Pertussis | 596.5 | 157 | 41 | 85 | 42 |
| Poliomyelitis | 0 | 0 | 0 | 0 | 0 |
| Rubella | | 1 | 5 | 4 | 3 |
| Tetanus | 1.5 | 0 | 1 | 1 | 1 |
| Varicella (associated with outbreaks) | 489.75 | 185 (15) | 178 (25) | 231 (25) | 220 (57) |

Varicella coverage by county



VAR Coverage Maps



Chickenpox cases by age 2023 to date





Chickenpox cases and outbreaks increasing





Detroit Hflu Outbreak May 2023

- Between May 1-May 8, Detroit Health Dept. was notified of 4 cases of invasive Haemophilus influenzae among students in the same school and grade.
 - Index patient deceased
 - 2 of 3 additional cases identified in same classroom as index case
 - All cases seen at hospital with sudden onset or rapid worsening of symptoms (fever, myalgia, lethargy, headache, vomiting, sore throat)
- In addition to Hi, 3 cases tested PCR positive for: adenovirus (2), rhinovirus/enterovirus (2), group A Streptococcus (2), RSV (1), SARS-CoV2 (1), and coronaviruses HKU1 and NL63 (1)



5/1

Epi-linked Non epi-linked

5/3

5/5

5/7

5/9 5/11

4/15 4/17 4/19 4/21 4/23 4/25 4/27 4/29

Detroit H Flu Continued



- Specimen from all 4 cases were sent to BOL for serotyping between 5/05-5/12 – all resulted as nontypeable
- Additional Whole Genome Sequencing (WGS) was performed by BOL and all specimen were found to share the same sequence type, indicating high probability of transmission.
- Nontypeable (unencapsulated) is the most common cause of invasive Hi disease, however secondary transmission is rare, and outbreaks are not often reported



Hi cases (<15 years) over time by serotype

'Other' includes nontypeable, beta-lactamase negative, type pending, or serogroups not marked as 'non-b.'

'Non-B' includes any cases marked as such + a serogroup, beta-lactamase negative, unable to grow, or *Haemophilus parainfluenza*.





- Fine area estimation of vaccine coverage can provide additional insights.
- Particularly during a VPD outbreak, these additional data can help drive decisions around resource allocation, priority communities, and vulnerable populations
- VPD cases and outbreaks are starting to return to pre-COVID levels, particularly for chickenpox.
- As we discussed this morning low immunization coverage leaves our communities and schools susceptible to outbreaks.