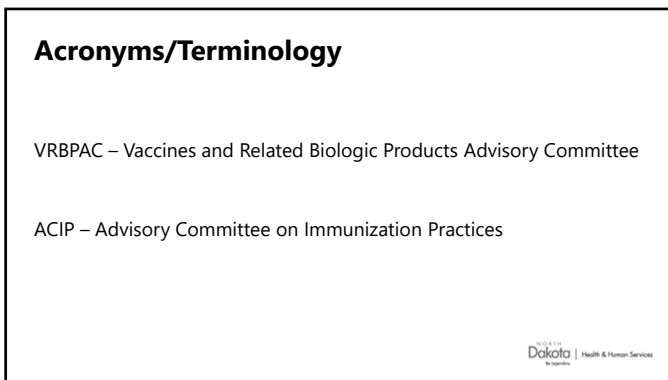




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ACIP February 2023 Topics



- Mpox Vaccine
- Influenza Vaccine
- Pneumococcal Vaccines
- Meningococcal Vaccines
- Polio Vaccine
- RSV Vaccines – Pediatric/Maternal
- RSV Vaccines – Adult
- Chikungunya Vaccine
- Dengue Vaccines
- Varicella
- COVID-19 Vaccines

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ACIP – Mpox Vaccine

- > 30,000 cases of mpox in the U.S. since May 2022
- Low numbers continuing
- Predominantly among men (gay, bisexual, and men who have sex with men)
- 2 vaccines available
 - Jynneos received an FDA EUA for pre- or post-exposure prophylaxis in children and intradermal administration in adults
 - ACAM2000 not being utilized in this outbreak
- Nearly 1.2 million doses of Jynneos administered during this outbreak
- Apparent equity issues

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ACIP – Mpox Vaccine

- No previous ACIP recommendation for Jynneos usage during outbreaks
- Efficacy data reviewed: 66%-83% after 2 doses regardless of route
- Safety data reviewed: no new concerns (myocarditis and pericarditis post-vaccination rates similar to general population)
- ACIP vote: recommend 2-dose Jynneos series (28 day separation) in people 18 and older at risk of mpox during an outbreak as defined by public health authorities
 - Subcutaneous administration recommended unless supply dictates intradermal administration
 - Utilization in individuals <18 years will be addressed at the June 2023 meeting
 - A more long-term strategy for Jynneos will be reviewed at the October 2023 meeting

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ACIP – Mpox Vaccine

Vaccine effectiveness of JYNNEOS against mpox ranges from 66%-83% for full vaccination and 36%-86% for partial vaccination

	Cases	Controls	Adjusted* VE (95% CI)
Full vaccination (2 doses)			
Epic Case-control study	25	335	66% (47%-79%)
Multi-jurisdictional case-control study	14	332	76% (48%-89%)
New York State case-control study	2	21	83% (22%-96%)
Partial vaccination (1 dose)			
Naval single-blind study	5	36	86% (29%-95%)
Epic Case-control study	146	1000	36% (22%-47%)
New York State case-control study	10	24	63% (22%-83%)

Vaccine Effectiveness (%)

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ACIP – Influenza Vaccine

- Early peak in U.S., late November/early December 2022
- Vast majority of cases have been Influenza A(H3N2) subtype
- Estimated 25 million illnesses, 280,000 hospitalizations, and 18,000 deaths
- Collected viruses susceptible to available antivirals



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ACIP – Influenza Vaccine

- Numerous studies of 2022-23 season vaccine efficacy, including NVSN, IVY, VISION Network, and Marshfield Clinic Research Institute
- Overall vaccine efficacy for 2022-23 season 49%
- Conveys significant protection in from hospitalization and in high-risk groups

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ACIP – Influenza Vaccine

Network – Population & Infection Severity	Efficacy
NVSN – Pediatric hospitalizations	68%
NVSN – Pediatric emergency department visits	42%
IVY – Adult hospitalizations	43%
VISION – Adult hospitalizations	39%
VISION – Adult emergency department/Urgent care visits	44%
Marshfield – Medically attended influenza A in children and working aged adults	54%
Marshfield – Symptomatic influenza A in children	71%

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ACIP – Pneumococcal Vaccines

Current recommendations:

- PCV13 – children
- PCV15 – children and adults
- PCV20 – adults and under FDA review for children with potential licensure as early as April 2023
- PPSV23 – some children based on their risks and adults with high-risk conditions

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ACIP – Pneumococcal Vaccines

CDC recommends pneumococcal vaccines for people at increased risk

Children

All children younger than 5 years old should receive PCV13 or PCV15.

Children 5 through 18 years old with certain medical conditions that increase their risk of pneumococcal disease should receive PCV13 or PCV15.

Children 2 through 18 years old with certain medical conditions should also receive PPSV23.

Adults

Adults who have never received a pneumococcal conjugate vaccine should receive PCV15 or PCV20 if they:

- Are 65 years and older
- Are 19 through 64 years old and have certain medical conditions or other risk factors

If PCV15 is used, it should be followed by a dose of PPSV23.

Adults who received an earlier pneumococcal conjugate vaccine (PCV13 or PCV7) should talk with a vaccine provider to learn about available options to complete their pneumococcal vaccine series.

Adults 65 years or older have the option to get PCV20 if they have already received:

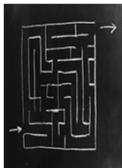
- PCV13 but not PCV15 or PCV20 at any age

and

- PPSV23 at or after the age of 65 years old

These adults can talk with their doctor and decide, together, whether to get PCV20.

Some people may need more than one dose of pneumococcal vaccines. Talk with your or your child's doctor about what is best for your specific situation.



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ACIP – Pneumococcal Vaccines

PCV20

- No anticipated impact to adult recommendations
- Reviewed possible schedules for children under 2 years – 4 doses vs. 3 doses of PCV13
- Discussed possibility of recommending PCV20 in place of PPSV23 in older children with underlying health conditions
- 20 is more than 13 or 15, so...?
- ACIP vote on recommendations anticipated after product licensure, likely in June 2023

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ACIP – Meningococcal Vaccines

- Decreased cases during pandemic, but numbers doubled in 2022
- 27 cases since 2019 of penicillin and ciprofloxacin resistant serogroup Y
- Serogroup B dominant in adolescents
- Currently MenACWY and MenB approved



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ACIP – Meningococcal Vaccines

- Pfizer MenABCWY pentavalent vaccine Phase 3 trial
- 2 dose series 6-12 months apart with a booster dose 4 years later
- ACIP Work Group considered the pentavalent vaccine noninferior to MenACWY + MenB, but some information lacking
- ACIP will continue to review at the June 2023 meeting and likely vote on any recommendations at the October 2023 meeting

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ACIP – Polio Vaccine

- Discussion surrounding possibility of an updated recommendation for using IPV in adults
 - Primary series or booster dose
 - Immunocompromised adults
 - Unvaccinated/incompletely vaccinated adults
- No anticipated impact to child recommendations
- Further discussion and possible vote at June 2023 meeting

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ACIP – Pediatric/Maternal RSV Vaccines



- United States season typically November-March
- Infects 97% of children by age 2
- Leading cause of hospitalizations in infants in the U.S.
- Pre-exposure prophylaxis available, but only 2% of infants meet palivizumab (Synagis®) indications
- No vaccine currently available for RSV

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ACIP – Pediatric/Maternal RSV Vaccines

- Nirsevimab – monoclonal antibody in joint development from Sanofi and AstraZeneca
- Potential recommendations:
 - All children born during RSV season
 - Children <8 months old during RSV season
 - High risk children during second RSV season
- FDA has third quarter 2023 as target action date for their decision
- Unknown if it will be included in the Vaccines For Children Program

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ACIP – Pediatric/Maternal RSV Vaccines

- Nirsevimab – a monoclonal antibody in joint development from Sanofi and AstraZeneca
- For the prevention of RSV lower respiratory tract disease in newborns and infants entering or during their first RSV season and for children up to 24 months who remain vulnerable to severe RSV disease through their second RSV season
- 79% efficacy against medically attended LRTI due to RSV, 80.6% efficacy against hospitalization, and 90% efficacy against ICU admission
- FDA has third quarter 2023 as target action date for their decision
- Cost, access, and administrative questions

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ACIP – Pediatric/Maternal RSV Vaccines

- Pfizer bivalent RSV Prefusion (RSVprf) vaccine product being studied for administration during pregnancy
- One dose at 24-36 weeks gestation to provide passive immunity to infants against LRTI and severe LRTI caused by RSV
- Phase 3 findings suggest a favorable safety profile and 82% efficacy against severe RSV up to 3 months of age and 69% at 6 months
- If licensed by the FDA, ACIP vote anticipated at October 2023 meeting

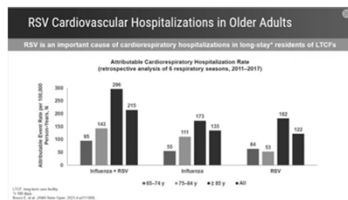


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ACIP – Adult RSV Vaccines

- Each year in the U.S. leads to 177,000 hospitalizations and 14,000 deaths among adults 65 years and older.



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ACIP – Adult RSV Vaccines

- Pfizer (bivalent RSVpreF) and GSK (adjuvanted RSVpreF3) each have a RSV vaccine in development for older adults
- In trials, both have shown efficacy against lower respiratory tract infections
 - GSK: 82.5% efficacy against RSV Lower Respiratory Tract Disease and 87.5% against medically attended RSV Lower Respiratory Tract Disease
 - Pfizer: 85.7% efficacy against RSV Lower Respiratory Tract Illness and 80.00% against medically attended RSV Lower Respiratory Tract Illness
- Safety: concerns about adverse events with both products, particularly Guillain-Barre syndrome
- Trials were conducted in adults 60 years and up, ACIP Work Group considering 65 years and up
- More data needed and products will be discussed at future meetings

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ACIP – Travel Vaccines – Chikungunya Virus

Chikungunya – virus spread via mosquito bite

- Symptoms can be prolonged
- Outbreaks occur in many places around the world
- Death rare
- Vaccine from Valneva may be licensed by FDA in August 2023
- ACIP vote likely at February 2024 meeting



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ACIP – Travel Vaccines – Dengue Virus


Dengue – virus spread via mosquito bite

- Symptoms can be prolonged symptoms
- Outbreaks occur in many places around the world
- About 1 in 20 people progress to severe dengue
- A vaccine is approved for children 9-16 years with laboratory-confirmed prior dengue infection who live in places where dengue is common
- New vaccine from Takeda being reviewed by FDA, possible long term protection for both naive and previously infected individuals
- ACIP will continue review at the June 2023 meeting and vote at the October 2023 meeting



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ACIP – 25 years of Varicella Vaccine

Varicella vaccine over the last 25 years:

- Has prevented up to 91 million cases of chickenpox
- Has prevented 238,000 hospitalizations for chickenpox
- Has prevented up to 2,446 deaths
- Societal cost savings of more than \$23 billion

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ACIP – COVID-19 Vaccines

Efficacy

- Continue to be highly effective with regards to serious outcomes
- Hospitalization rate in unvaccinated adults 16 times that of adults receiving a bivalent mRNA booster
- Hospitalization rate in vaccinated adults without a bivalent booster 2.6 times that of adults receiving a bivalent booster

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ACIP – COVID-19 Vaccines

Safety

- November 2022 possible safety signal for ischemic stroke in adults 65 years and up in Vaccine Safety Datalink (VSD)
 - No other vaccine surveillance system domestically or internationally reflected a similar safety signal and signal has since diminished
 - Further review showed signal was stronger when Pfizer bivalent booster and influenza vaccine given concurrently
 - No causal link proven, investigation continues
 - No change in vaccination recommendations
- Safety data for the bivalent booster do not show an increased risk of myocarditis in adolescent males

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ACIP – COVID-19 Vaccines

Simplification of recommendations under consideration

- Adopt a 1-dose schedule after early childhood
- End use of monovalent vaccines, utilizing bivalent for both primary series and booster
- NO changes at this time



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VRBPAC February-March 2023 Topics

- Safety and effectiveness of ABRYSVO, Pfizer's Respiratory Syncytial Virus Vaccine for adults 60 years and older
- Safety and effectiveness of AVEXVY, GSK's Respiratory Syncytial Virus Vaccine, Recombinant, Adjuvanted for adults 60 years and older



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VRBPAC February-March 2023

ABRYSVO
Continued
Monitoring

Pharmacovigilance Plan	
The Applicant will conduct passive and active surveillance activities for continued vaccine safety monitoring, including routine pharmacovigilance.	
Missing Information	Immunocompromised older adults
Surveillance Activities	• Planned postmarketing safety study in immunocompromised older adults
Important Potential Risks	• GBS and other immune-mediated demyelinating conditions • Cardiac disorders
Surveillance Activities	• Expedited reporting for all cases of GBS and other immune-mediated demyelinating conditions and Cardiac disorders • Aggregate analysis of GBS and other immune-mediated demyelinating conditions and Cardiac disorders in periodic safety reports • Planned postmarketing safety study to assess the risk of GBS and other immune-mediated demyelinating conditions among individuals vaccinated with ABRYSVO (RSVpreF)

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VRBPAC February-March 2023

ABRYSVO Efficacy

Summary - Efficacy

- VE to prevent first-episode LRTI-RSV with ≥ 2 and ≥ 3 symptoms were 66.7% (96.66% CI 28.8, 85.8) and 85.7% (96.66% CI 32.0, 98.7), respectively
 - Study success criterion met
- Descriptive point estimates for VE against LRTI-RSV appear preserved among participants ≥ 60 years and among participants with at least one at-risk condition, but were limited by small subpopulation sizes
- Preliminary VE against ARI-RSV was 62.1% (95% CI 37.1, 77.9)
- Data are not currently available on:
 - Duration of vaccine effectiveness
 - VE in immunocompromised and frail elderly individuals
 - VE in preventing severe LRTI cases
 - Data regarding concomitant administration with vaccines routinely recommended for use in this population

Patent (A)

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VRBPAC February-March 2023

ABRYSVO Safety

Summary - Safety

- Study included 34,284 participants (17,215 who received RSVpreF), of which 26,395 participants (77.0%) have had at least 6 months of follow-up
- Solicited local and systemic reactions were generally mild to moderate and of short duration
- Within 1 month after vaccination, a numerical imbalance was observed for events of atrial fibrillation (10 in RSVpreF group vs. 4 in placebo group). FDA review of these events is ongoing.
- Serious adverse events were balanced between the RSVpreF and placebo groups (2.3% in both groups)
- Three SAEs (hypersensitivity, GBS, and Miller Fisher Syndrome) were assessed by FDA as possibly related to RSVpreF, in agreement with the investigator's assessment
- Review of safety data from the 5 supportive clinical studies (~1,200 participants who received RSVpreF final formulation) did not reveal any other cases of GBS or other immune-mediated demyelinating condition post-vaccination, or any other safety signal

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VRBPAC February-March 2023

ABRYSVO

1. 7 yes
4 no
1 abstain
2. 7 yes
4 no
1 abstain

Voting Questions for VRBPAC

1. Are the available data adequate to support the safety of ABRYSVO (RSVpreF) when administered to individuals 60 years of age and older for the prevention of lower respiratory tract disease caused by RSV?
Please vote "Yes" or "No"
2. Are the available data adequate to support the effectiveness of ABRYSVO (RSVpreF) for the prevention of lower respiratory tract disease caused by RSV in individuals 60 years of age and older?
Please vote "Yes" or "No"

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VRBPAC February-March 2023

AVEXVY
Continued
Monitoring

Pharmacovigilance Plan

Missing information

Immunocompromised older adults

Surveillance Activities

- A post-authorization clinical trial evaluating AREXVY in immunocompromised individuals is being designed

Important potential risks

Potential Immune-Mediated Disorders (pIMDs)

Surveillance activities

Applicant will conduct passive surveillance activities for continued vaccine safety monitoring, including routine pharmacovigilance and:

- Expedited reporting for all cases of:
 - GBS, ADEM, and other immune-mediated demyelinating conditions and neurologic conditions
 - Supraventricular arrhythmias
- Aggregate analysis, in periodic safety reports, for:
 - GBS, ADEM, and other immune-mediated demyelinating conditions
 - Supraventricular arrhythmias

Note that the following are currently under discussion between FDA and the Applicant:

- Plans for a postmarketing safety study to assess the risk of GBS, ADEM, and other immune-mediated demyelinating conditions among individuals vaccinated with AREXVY.
- Determination of the inclusion of Cardiac Disorders as an important potential risk in the PVP.

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VRBPAC February-March 2023

AVEXVY
Efficacy

Summary: Efficacy

- VE against first occurrence of RT-PCR-confirmed RSV LRTD was 82.6% (96.95% CI 57.5, 94.1) in adults ≥60 YOA
- Subgroup analysis showed that VE was demonstrated for: RSV-A and RSV-B virus subtypes; age groups 60-69 YOA and 70-79 YOA; participants with at least 1 pre-existing comorbidity of interest, and RSV-ARI.
 - VE in participants ≥80 YOA inconclusive due to low number of cases
- VE demonstrated for Severe LRTD based on clinical symptomatology definition but the numbers of accrued cases meeting the definition based on supportive therapy were too small to estimate efficacy precisely.
- Data are not currently available on:
 - Duration of vaccine effectiveness
 - VE in immunocompromised individuals

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VRBPAC February-March 2023

AVEXVY
Safety

Summary: Safety

- A total of 15,745 RSVvPref3 recipients from four phase 3 studies were included in the Exposed Set. The median duration of follow-up was 7.2 months.
- RSVvPref3 is noted to have increased reactivity when compared to placebo, but the rates of Grade 3 reactions after vaccination in both groups were low (1.7%).
- Within 30 days post-vaccination a numerical imbalance was observed for events of atrial fibrillation in Study 006. FDA review of these events is ongoing.
- The frequency of SAEs reported up to 6 months post-vaccination was 4.0% and 4.5% in the RSVvPref3 and placebo groups.
- One (1) SAE (GBS) was considered by the study investigator and FDA to be related to vaccination.
- One (1) death due to ADEM considered by FDA as possibly related to FIU or RSVvPref3 vaccination.
- Up to the time of the DLPs at least one pIMD was reported by 0.4% and 0.3% of vaccine and placebo recipients, including 2 cases of ADEM in the Co-Ax group in Study 007.
- A safety update was submitted for an extended safety follow-up at Month 6-12, containing SAE and pIMD data, and FDA review of these data are ongoing at this time.

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Recommended Adult Immunization Schedule for ages 19 years or older

UNITED STATES
2023

1 Determine
the adult
immunization
schedule by
age (Table 1)

2 Assess
each adult
immunization
recommendation
for the
individual
(Table 2)

3 Review
contraindications
and
precautions
(Table 3)

4 Review
the immunizations
and
precautions
for each
individual
(Appendix)

Vaccines in the Adult Immunization Schedule*

How to use:

- Table 1** provides the recommended adult immunization schedule by age.
- Table 2** provides information on each vaccine.
- Table 3** provides information on each vaccine's contraindications and precautions.
- Appendix** provides information on each vaccine's contraindications and precautions.

Table 1. Recommended Adult Immunization Schedule by Age

Age	Vaccine
19-64 years	Td/DTaP vaccine
19-64 years	Tdap vaccine
19-64 years	MMV2 vaccine
19-64 years	MMV4 vaccine
19-64 years	MMV6 vaccine
19-64 years	MMV8 vaccine
19-64 years	MMV10 vaccine
19-64 years	MMV12 vaccine
19-64 years	MMV14 vaccine
19-64 years	MMV16 vaccine
19-64 years	MMV18 vaccine
19-64 years	MMV20 vaccine
19-64 years	MMV22 vaccine
19-64 years	MMV24 vaccine
19-64 years	MMV26 vaccine
19-64 years	MMV28 vaccine
19-64 years	MMV30 vaccine
19-64 years	MMV32 vaccine
19-64 years	MMV34 vaccine
19-64 years	MMV36 vaccine
19-64 years	MMV38 vaccine
19-64 years	MMV40 vaccine
19-64 years	MMV42 vaccine
19-64 years	MMV44 vaccine
19-64 years	MMV46 vaccine
19-64 years	MMV48 vaccine
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19-64 years	MMV208 vaccine
19-64 years	MMV210 vaccine
19-64 years	MMV212 vaccine
19-64 years	MMV214 vaccine
19-64 years	MMV216 vaccine
19-64 years	MMV218 vaccine
19-64 years	MMV220 vaccine
19-64 years	MMV222 vaccine
19-64 years	MMV224 vaccine

Table 1 Recommended Adult Immunization Schedule by Age Group, United States, 2023				
Vaccine	19–26 years	27–49 years	50–64 years	≥65 years
COVID-19		2- or 3-dose primary series and booster (See Notes)		
Influenza inactivated (IIV4) or Influenza recombinant (rIIV4)		1 dose annually		
Influenza live, attenuated (LIV4)		1 dose annually		
Tetanus, diphtheria, pertussis (Tdap or Td)		1 dose Tdap each pregnancy; 1 dose Td/Tdap for wound management (see notes)		
Measles, mumps, rubella (MMR)		1 or 2 doses depending on indication (if born in 1957 or later)		For healthcare personnel, see notes
Varicella (VAR)	2 doses (if born in 1980 or later)		2 doses	
Zoster recombinant (ZV)	2 doses for immunocompromising conditions (see notes)		2 doses	
Human papillomavirus (HPV)	2 or 3 doses depending on age at initial vaccination or condition	27 through 45 years		
Pneumococcal (PCV13, PCV20, PPV23)		1 dose PCV13 followed by PPV23 OR 1 dose PCV20 (see notes)		See Notes
Hepatitis A (HepA)		2, 3, or 4 doses depending on vaccine		See Notes
Hepatitis B (HepB)		2, 3, or 4 doses depending on vaccine or condition		
Meningococcal A, C, W, Y (MenACWY)		1 or 2 doses depending on indication, see notes for booster recommendations		
Meningococcal B (MenB)		2 or 3 doses depending on vaccine and indication, see notes for booster recommendations		
Neisseria meningitidis type b (MenB)		1 or 3 doses depending on indication		

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Table 2 Recommended Adult Immunization Schedule by Medical Condition or Other Indication, United States, 2023										
Vaccine	Pregnancy	Immune compromised (including HIV infection)	HIV infection CD4 percentage and count (CD4% and CD4 count)	Asplenia, splenectomy, or hyposplenism	End-organ transplant, or on immunosuppressive therapy	Heart or lung transplant, or on immunosuppressive therapy	Chronic liver disease	Diabetes	Health care personnel	Those who work with animals
COVID-19		See Notes								
IIV4 or rIIV4 (IIV4)		Contraindicated					1 dose annually			1 dose annually
Tdap or Td	1 dose Tdap each pregnancy	Contraindicated					1 dose Tdap, then Td or Tdap booster every 10 years			Precaution
MMR	Contraindicated ^a	Contraindicated						1 or 2 doses depending on indication		
VAR	Contraindicated ^a	Contraindicated						2 doses		
RVZ		2 doses at age ≥19 years					2 doses at age ≥50 years			
HPV	See Notes	3 doses through age 26 years					2 or 3 doses through age 26 years depending on age at initial vaccination or condition			
PCV13, PCV20, PPV23	See Notes						1 dose PCV13 followed by PPV23 OR 1 dose PCV20 (see notes)			
HepA							2, 3, or 4 doses depending on vaccine			
HepB	1 dose (see notes)						2, 3, or 4 doses depending on vaccine or condition			
MenACWY		1 or 2 doses depending on indication, see notes for booster recommendations								
MenB	Precaution	2 or 3 doses depending on vaccine and indication, see notes for booster recommendations								
Hib		2 doses (see Notes)					1 dose			
<div><div><div>■ Contraindicated^a or see notes for additional information on safety or contraindications on IIV4, rIIV4, Tdap, or Td</div><div>■ Based on clinical contraindications or contraindications on IIV4, rIIV4, Tdap, or Td</div><div>■ Based on clinical contraindications or contraindications on IIV4, rIIV4, Tdap, or Td</div><div>■ Based on clinical contraindications or contraindications on IIV4, rIIV4, Tdap, or Td</div><div>■ Based on clinical contraindications or contraindications on IIV4, rIIV4, Tdap, or Td</div><div>■ Based on clinical contraindications or contraindications on IIV4, rIIV4, Tdap, or Td</div><div>■ Based on clinical contraindications or contraindications on IIV4, rIIV4, Tdap, or Td</div><div>■ Based on clinical contraindications or contraindications on IIV4, rIIV4, Tdap, or Td</div><div>■ Based on clinical contraindications or contraindications on IIV4, rIIV4, Tdap, or Td</div><div>■ Based on clinical contraindications or contraindications on IIV4, rIIV4, Tdap, 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2023 Recommended Immunization Schedule Adult ages 19 years or older

Routine Immunization Schedule (table 1) updates:

- New COVID-19 vaccine row in yellow, indicating that it is routinely recommended for all adults
- MMR row – text in the column for ≥65 years referring providers to notes for considerations for healthcare providers
- Hepatitis A row – updated text “2, 3, or 4 doses depending on vaccine”

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2023 Recommended Immunization Schedule Adult ages 19 years or older

Immunization by Medical Indication Schedule (table 2) updates:

- New COVID-19 vaccine row in yellow, refers to notes regarding the immunocompromised and HIV positive
- Hepatitis A row – updated text “2, 3, or 4 doses depending on vaccine”

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2023 Recommended Immunization Schedule Adult ages 19 years or older

Vaccine Note updates:

- Edits to mirror the language between the Adult Schedule and the Child and Adolescent Schedule when possible
- Notes updated for COVID-19, HepB, Influenza, MMR, Meningococcal, Pneumococcal, Poliovirus, Tdap, and Zoster vaccines

Appendix updates:

- “Contraindications” column renamed to “Contraindicated or not recommended”
- Influenza (egg-based) row – history of egg allergy info moved from precautions column to notes
- Hep B row and human papillomavirus rows had language modifications regarding pregnancy

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

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Vaccines in the Child and Adolescent Immunization Schedule*

Vaccine	ACIP recommendation	U.S. & T. recommendation
COVID-19	2023-01-05	2023-01-05
Diphtheria, tetanus, and acellular pertussis vaccine	DTaP	DTaP
Hepatitis A vaccine	HepA	HepA
Hepatitis B vaccine	HepB	HepB
Influenza vaccine (inactivated)	Flu	Flu
Influenza vaccine (live attenuated)	Flu	Flu
Meningococcal conjugate vaccine	MenACWY	MenACWY
Meningococcal polysaccharide vaccine	MenPSV	MenPSV
Poliovirus vaccine (inactivated)	IPV	IPV
Poliovirus vaccine (live attenuated)	OPV	OPV
Tdap vaccine	Tdap	Tdap
Zoster vaccine	Zvax	Zvax

How to use the child and adolescent immunization schedule

1. Recommended routine immunization schedule for children and adolescents (Table 1)
2. Recommended routine immunization schedule for children and adolescents (Table 2)
3. Recommended routine immunization schedule for children and adolescents (Table 3)
4. Recommended routine immunization schedule for children and adolescents (Table 4)
5. Recommended routine immunization schedule for children and adolescents (Table 5)

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

Report

- Reportable cases of reportable vaccine-preventable diseases or outbreaks to your state or local health department.
- Clinically significant adverse events to the Vaccine Adverse Event Reporting System (VAERS) at vaers.hhs.gov or 800-858-7867.

Questions or comments

- Contact your state health department for more information (including contraindications and precautions).
- Contact your state health department for more information (including contraindications and precautions).
- Contact your state health department for more information (including contraindications and precautions).
- Contact your state health department for more information (including contraindications and precautions).

Helpful information

- <https://www.cdc.gov/vaccines/imz/downloads/2023-01-05/>
- <https://www.cdc.gov/vaccines/imz/downloads/2023-01-05/>
- <https://www.cdc.gov/vaccines/imz/downloads/2023-01-05/>
- <https://www.cdc.gov/vaccines/imz/downloads/2023-01-05/>

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2023 Recommended Immunization Schedule Child and Adolescent ages 18 years or younger

Routine Immunization Schedule (table 1) updates:

- New COVID-19 vaccine row in yellow, indicating that it is routinely recommended for 6 months to 18 years
- Pneumococcal conjugate row – PCV 15 added
- IPV row – prompts healthcare providers to review the Notes section for people age 18 years

Catch-up Immunization Schedule (table 2) updates:

- Pneumococcal conjugate row – language for the minimum interval between doses 3 and 4 revised

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2023 Recommended Immunization Schedule Child and Adolescent ages 18 years or younger

Immunization by Medical Indication Schedule (table 3) updates:

- New COVID-19 vaccine row in yellow, refers to notes regarding the immunocompromised and HIV positive

Vaccine Note updates:

- Edits to mirror the language between the Adult Schedule and the Child and Adolescent Schedule when possible
- Notes updated for COVID-19, Dengue, HepB, Influenza, MMR, MenACWY, MenB, Pneumococcal, and Poliovirus vaccines

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2023 Recommended Immunization Schedule Child and Adolescent ages 18 years or younger

Appendix updates:

- "Contraindications" column renamed to "Contraindicated or not recommended"
- Influenza (egg-based) row – history of egg allergy with symptoms other than hives info moved from precautions column to notes
- Dengue row – lack of lab confirmation of previous dengue virus infection is a contraindication added
- Hep B row and human papillomavirus rows had language modifications regarding pregnancy
- MMR row – MMRV added, precaution for using MMRV of personal or family history of seizure added
- Varicella row – precautions for MMR/MMRV should be reviewed if using MMRV

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Post-Test

- Post-test
 - Nurses interested in continuing education credit, visit https://ndhealth.co1.qualtrics.com/jfe/form/SV_6LH8Fmg6rIRNOig
 - Successfully complete the five-question post-test to receive your certificate
 - Credit for this session will not expire until April 22, 2023.
- This presentation will be posted to our website: www.hhs.nd.gov/immunizations

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Immunization Update:

ACIP and VRBPAC Meeting Reviews

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**Immunization Update:
ACIP and VRBPAC Meeting Reviews**

References:

www.cdc.gov

IZ Express. "Immunize.org summarizes ACIP's February 22-24 meeting recommending mpox vaccine for adults during outbreaks and discussing RSV, pneumococcal disease, meningococcal disease, future COVID-19 vaccine plans, and more." Issue 1,680: March 1, 2023.

[Vaccines and Related Biological Products Advisory Committee February 28 - March 1, 2023 Meeting Announcement - 02/28/2023 | FDA](#)

[Today's ACIP meeting Cliff notes - by Katelyn Jetelina \(substack.com\)](#)

Howell, Molly. "Advisory Committee on Immunization Practices (ACIP) Notes, February 22-24, 2023."

Schaffner, William and Carrico, Ruth. "Respiratory Syncytial Virus: It's Not Just for Kids - A Review of the Data Clinicians Need to Know." [Respiratory Syncytial Virus: It's Not Just for Kids - A Review of the Data Clinicians Need to Know \(medscape.org\)](#) Feb 6, 2023.

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