



# **2018 Adult Immunization Schedule Updates**

**National Adult and Influenza Immunization Summit**  
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# 2018 Adult Immunization Schedule Updates

- Recommended use of recombinant zoster vaccine
- Recommended use of MMR in mumps outbreak setting
- Updated ACIP recommendations in prevention of hepatitis B
- FDA licensure of conjugate 1018-adjuvanted hepatitis B vaccine

# Recommended Immunization Schedule for Adults Aged 19 Years or Older, United States, 2018

In February 2018, the *Recommended Immunization Schedule for Adults Aged 19 Years or Older, United States, 2018* became effective, as recommended by the Advisory Committee on Immunization Practices (ACIP) and approved by the Centers for Disease Control and Prevention (CDC). The adult immunization schedule was also approved by the American College of Physicians, the American Academy of Family Physicians, the American College of Obstetricians and Gynecologists, and the American College of Nurse-Midwives.

CDC announced the availability of the 2018 adult immunization schedule in the *Morbidity and Mortality Weekly Report (MMWR)*.<sup>1</sup> The schedule is published in its entirety in the *Annals of Internal Medicine*.<sup>2</sup>

The adult immunization schedule consists of figures that summarize routinely recommended vaccines for adults by age groups and medical conditions and other indications, footnotes for the figures, and a table of vaccine contraindications and precautions. Note the following when reviewing the adult immunization schedule:

- The figures in the adult immunization schedule should be reviewed with the accompanying footnotes.
- The figures and footnotes display indications for which vaccines, if not previously administered, should be administered unless noted otherwise.
- The table of contraindications and precautions identifies populations and situations for which vaccines should not be used or should be used with caution.
- When indicated, administer recommended vaccines to adults whose vaccination history is incomplete or unknown.
- Increased interval between doses of a multidose vaccine series does not diminish vaccine effectiveness; it is not necessary to restart the vaccine series or add doses to the series because of an extended interval between doses.
- Combination vaccines may be used when any component of the combination is indicated and when the other components of the combination are not contraindicated.
- The use of trade names in the adult immunization schedule is for identification purposes only and does not imply endorsement by the ACIP or CDC.

Special populations that need additional considerations include:

- Pregnant women. Pregnant women should receive the tetanus, diphtheria, and acellular pertussis vaccine (Tdap) during pregnancy and the influenza vaccine during or before pregnancy. Live vaccines (e.g., measles, mumps, and rubella vaccine [MMR]) are contraindicated.
- Asplenia. Adults with asplenia have specific vaccination recommendations because of their increased risk for infection by encapsulated bacteria. Anatomical or functional asplenia includes congenital or acquired asplenia, splenic dysfunction, sickle cell disease and other hemoglobinopathies, and splenectomy.
- Immunocompromising conditions. Adults with immunosuppression should generally avoid live vaccines. Inactivated vaccines (e.g., pneumococcal vaccines) are generally acceptable. High-level immunosuppression includes HIV infection with a CD4 cell count <200 cells/ $\mu$ L, receipt of daily corticosteroid therapy with  $\geq 20$  mg of prednisone or equivalent for  $\geq 14$  days, primary immunodeficiency disorder (e.g., severe combined immunodeficiency or complement component deficiency), and receipt of cancer chemotherapy. Other immunocompromising conditions and immunosuppressive medications to consider when vaccinating adults can be found in *IDSA Clinical Practice Guideline for Vaccination of the Immunocompromised Host*.<sup>3</sup> Additional information on vaccinating immunocompromised adults is in *General Best Practice Guidelines for Immunization*.<sup>4</sup>

Additional resources for health care providers include:

- Details on vaccines recommended for adults and complete ACIP statements at [www.cdc.gov/vaccines/hcp/acip-recs/index.html](http://www.cdc.gov/vaccines/hcp/acip-recs/index.html)
- Vaccine Information Statements that explain benefits and risks of vaccines at [www.cdc.gov/vaccines/hcp/vis/index.html](http://www.cdc.gov/vaccines/hcp/vis/index.html)
- Information and resources on vaccinating pregnant women at [www.cdc.gov/vaccines/adults/rec-vac/pregnant.html](http://www.cdc.gov/vaccines/adults/rec-vac/pregnant.html)
- Information on travel vaccine requirements and recommendations at [www.cdc.gov/travel/destinations/list](http://www.cdc.gov/travel/destinations/list)
- CDC Vaccine Schedules App for immunization service providers to download at [www.cdc.gov/vaccines/schedules/hcp/schedule-app.html](http://www.cdc.gov/vaccines/schedules/hcp/schedule-app.html)
- Adult Vaccination Quiz for self-assessment of vaccination needs based on age, health conditions, and other indications at [www2.cdc.gov/nip/adultimmsched/default.asp](http://www2.cdc.gov/nip/adultimmsched/default.asp)
- *Recommended Immunization Schedule for Children and Adolescents Aged 18 Years or Younger* at [www.cdc.gov/vaccines/schedules/hcp/child-adolescent.html](http://www.cdc.gov/vaccines/schedules/hcp/child-adolescent.html)

Report suspected cases of reportable vaccine-preventable diseases to the local or state health department, and report all clinically significant postvaccination events to the Vaccine Adverse Event Reporting System at [www.vaers.hhs.gov](http://www.vaers.hhs.gov) or by telephone, 800-822-7967. All vaccines included in the adult immunization schedule except 23-valent pneumococcal polysaccharide and zoster vaccines are covered by the Vaccine Injury Compensation Program. Information on how to file a vaccine injury claim is available at [www.hrsa.gov/vaccinecompensation](http://www.hrsa.gov/vaccinecompensation) or by telephone, 800-338-2382. Submit questions and comments to CDC through [www.cdc.gov/cdc-info](http://www.cdc.gov/cdc-info) or by telephone, 800-CDC-INFO (800-232-4636), in English and Spanish, 8:00am–8:00pm ET, Monday–Friday, excluding holidays.

The following abbreviations are used for vaccines in the adult immunization schedule (in the order of their appearance):

IIV	inactivated influenza vaccine
RIV	recombinant influenza vaccine
Tdap	tetanus toxoid, reduced diphtheria toxoid, and acellular pertussis vaccine
Td	tetanus and diphtheria toxoids
MMR	measles, mumps, and rubella vaccine
VAR	varicella vaccine
RZV	recombinant zoster vaccine
ZVL	zoster vaccine live
HPV vaccine	human papillomavirus vaccine
PCV13	13-valent pneumococcal conjugate vaccine
PPSV23	23-valent pneumococcal polysaccharide vaccine
HepA	hepatitis A vaccine
HepA-HepB	hepatitis A vaccine and hepatitis B vaccine
HepB	hepatitis B vaccine
MenACWY	serogroups A, C, W, and Y meningococcal vaccine
MenB	serogroup B meningococcal vaccine
Hib	<i>Haemophilus influenzae</i> type b vaccine

1. MMWR Morb Mortal Wkly Rep. 2018;66(5):xx–xx. Available at [www.cdc.gov/mmwr/volumes/67/xxxxxxxxxx](http://www.cdc.gov/mmwr/volumes/67/xxxxxxxxxx).

2. Ann Intern Med. 2018;168:xxx–xxx. Available at [annals.org/aim/article/doi/10.7326/M17-3439](http://annals.org/aim/article/doi/10.7326/M17-3439).

3. Clin Infect Dis. 2014;58:e44–100. Available at [www.idsociety.org/Templates/Content.aspx?id=32212256011](http://www.idsociety.org/Templates/Content.aspx?id=32212256011).


4. Kroger et al. Available at [www.cdc.gov/vaccines/hcp/acip-recs/general-recs/index.html](http://www.cdc.gov/vaccines/hcp/acip-recs/general-recs/index.html).





**Figure 1. Recommended immunization schedule for adults aged 19 years or older by age group, United States, 2018**

This figure should be reviewed with the accompanying footnotes. This figure and the footnotes describe indications for which vaccines, if not previously administered, should be administered unless noted otherwise.

Vaccine	19–21 years	22–26 years	27–49 years	50–64 years	≥65 years
Influenza <sup>1</sup>	1 dose annually				
Tdap <sup>2</sup> or Td <sup>2</sup>	1 dose Tdap, then Td booster every 10 yrs				
MMR <sup>3</sup>	1 or 2 doses depending on indication (if born in 1957 or later)				
VAR <sup>4</sup>	2 doses				
RZV <sup>5</sup> (preferred) or ZVL <sup>5</sup>				2 doses RZV (preferred) or 1 dose ZVL	
HPV–Female <sup>6</sup>	2 or 3 doses depending on age at series initiation				
HPV–Male <sup>6</sup>	2 or 3 doses depending on age at series initiation				
PCV13 <sup>7</sup>					1 dose
PPSV23 <sup>7</sup>	1 or 2 doses depending on indication				1 dose
HepA <sup>8</sup>	2 or 3 doses depending on vaccine				
HepB <sup>9</sup>	3 doses				
MenACWY <sup>10</sup>	1 or 2 doses depending on indication, then booster every 5 yrs if risk remains				
MenB <sup>10</sup>	2 or 3 doses depending on vaccine				
Hib <sup>11</sup>	1 or 3 doses depending on indication				

 Recommended for adults who meet the age requirement, lack documentation of vaccination, or lack evidence of past infection

 Recommended for adults with other indications

 No recommendation

**Figure 2. Recommended immunization schedule for adults aged 19 years or older by medical condition and other indications, United States, 2018**

This figure should be reviewed with the accompanying footnotes. This figure and the footnotes describe indications for which vaccines, if not previously administered, should be administered unless noted otherwise.

Vaccine	Pregnancy <sup>1-6</sup>	Immuno-compromised (excluding HIV infection) <sup>3,7,11</sup>	HIV infection CD4+ count (cells/ $\mu$ L) <sup>3,7,9-10</sup>		Asplenia, complement deficiencies <sup>7,10,11</sup>	End-stage renal disease, on hemodialysis <sup>7,9</sup>	Heart or lung disease, alcoholism <sup>7</sup>	Chronic liver disease <sup>7,9</sup>	Diabetes <sup>7,9</sup>	Health care personnel <sup>3,4,9</sup>	Men who have sex with men <sup>6,8,9</sup>	
			<200	$\geq$ 200								
Influenza <sup>1</sup>												1 dose annually
Tdap <sup>2</sup> or Td <sup>2</sup>	1 dose Tdap each pregnancy											1 dose Tdap, then Td booster every 10 yrs
MMR <sup>3</sup>		contraindicated										1 or 2 doses depending on indication
VAR <sup>4</sup>		contraindicated										2 doses
RZV <sup>5</sup> (preferred) or ZVL <sup>5</sup>												2 doses RZV at age $\geq$ 50 yrs (preferred) or 1 dose ZVL at age $\geq$ 60 yrs
HPV-Female <sup>6</sup>												3 doses through age 26 yrs
HPV-Male <sup>6</sup>												3 doses through age 26 yrs
PCV13 <sup>7</sup>												2 or 3 doses through age 26 yrs
PPSV23 <sup>7</sup>												2 or 3 doses through age 21 yrs
HepA <sup>8</sup>												1 dose
HepB <sup>9</sup>												1, 2, or 3 doses depending on indication
MenACWY <sup>10</sup>												2 or 3 doses depending on vaccine
MenB <sup>10</sup>												3 doses
Hib <sup>11</sup>												1 or 2 doses depending on indication, then booster every 5 yrs if risk remains
												2 or 3 doses depending on vaccine
												3 doses HSCT recipients only
												1 dose

Recommended for adults who meet the age requirement, lack documentation of vaccination, or lack evidence of past infection
  Recommended for adults with other indications
  Contraindicated
  No recommendation

## Footnotes. Recommended immunization schedule for adults aged 19 years or older, United States, 2018

### 1. Influenza vaccination

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/flu.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/flu.html)

#### General Information

- Administer 1 dose of age-appropriate inactivated influenza vaccine (IIV) or recombinant influenza vaccine (RIV) annually
- Live attenuated influenza vaccine (LAIV) is not recommended for the 2017–2018 influenza season
- A list of currently available influenza vaccines is available at [www.cdc.gov/flu/protect/vaccine/vaccines.htm](http://www.cdc.gov/flu/protect/vaccine/vaccines.htm)

#### Special populations

- Administer age-appropriate IIV or RIV to:
  - **Pregnant women**
  - Adults with **hives-only egg allergy**
  - Adults with **egg allergy other than hives** (e.g., angioedema or respiratory distress): Administer IIV or RIV in a medical setting under supervision of a health care provider who can recognize and manage severe allergic conditions

### 2. Tetanus, diphtheria, and pertussis vaccination

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/tdap-td.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/tdap-td.html)

#### General Information

- Administer to adults who previously did not receive a dose of tetanus toxoid, reduced diphtheria toxoid, and acellular pertussis vaccine (Tdap) as an adult or child (routinely recommended at age 11–12 years) 1 dose of Tdap, followed by a dose of tetanus and diphtheria toxoids (Td) booster every 10 years
- Information on the use of Tdap or Td as tetanus prophylaxis in wound management is available at [www.cdc.gov/mmwr/preview/mmwrhtml/rr5517a1.htm](http://www.cdc.gov/mmwr/preview/mmwrhtml/rr5517a1.htm)

#### Special populations

- **Pregnant women:** Administer 1 dose of Tdap during each pregnancy, preferably in the early part of gestational weeks 27–36

### 3. Measles, mumps, and rubella vaccination

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/mmr.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/mmr.html)

#### General Information

- Administer 1 dose of measles, mumps, and rubella vaccine (MMR) to adults with no evidence of immunity to measles, mumps, or rubella
- Evidence of immunity is:
  - Born before 1957 (except for health care personnel, see below)
  - Documentation of receipt of MMR
  - Laboratory evidence of immunity or disease
- Documentation of a health care provider-diagnosed disease without laboratory confirmation is not considered evidence of immunity

#### Special populations

- **Pregnant women and nonpregnant women of childbearing age** with no evidence of immunity to rubella: Administer 1 dose of MMR (if pregnant, administer MMR after pregnancy and before discharge from health care facility)

- **HIV infection and CD4 cell count  $\geq 200$  cells/ $\mu$ L for at least 6 months** and no evidence of immunity to measles, mumps, or rubella: Administer 2 doses of MMR at least 28 days apart
- **Students in postsecondary educational institutions, international travelers, and household contacts of immunocompromised persons:** Administer 2 doses of MMR at least 28 days apart (or 1 dose of MMR if previously administered 1 dose of MMR)

- **Health care personnel born in 1957 or later** with no evidence of immunity: Administer 2 doses of MMR at least 28 days apart for measles or mumps, or 1 dose of MMR for rubella (if born before 1957, consider MMR vaccination)
- Adults who **previously received  $\leq 2$  doses of mumps-containing vaccine and are identified by public health authority to be at increased risk for mumps in an outbreak:** Administer 1 dose of MMR
- MMR is contraindicated for pregnant women and adults with severe immunodeficiency

### 4. Varicella vaccination

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/varicella.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/varicella.html)

#### General Information

- Administer to adults without evidence of immunity to varicella 2 doses of varicella vaccine (VAR) 4–8 weeks apart if previously received no varicella-containing vaccine (if previously received 1 dose of varicella-containing vaccine, administer 1 dose of VAR at least 4 weeks after the first dose)
- Evidence of immunity to varicella is:
  - U.S.-born before 1980 (except for pregnant women and health care personnel, see below)
  - Documentation of receipt of 2 doses of varicella or varicella-containing vaccine at least 4 weeks apart
  - Diagnosis or verification of history of varicella or herpes zoster by a health care provider
  - Laboratory evidence of immunity or disease

#### Special populations

- Administer 2 doses of VAR 4–8 weeks apart if previously received no varicella-containing vaccine (if previously received 1 dose of varicella-containing vaccine, administer 1 dose of VAR at least 4 weeks after the first dose) to:
  - **Pregnant women without evidence of immunity:** Administer the first of the 2 doses or the second dose after pregnancy and before discharge from health care facility
  - **Health care personnel without evidence of immunity**
- Adults with **HIV infection and CD4 cell count  $\geq 200$  cells/ $\mu$ L:** May administer, based on individual clinical decision, 2 doses of VAR 3 months apart
- VAR is contraindicated for pregnant women and adults with severe immunodeficiency

### 5. Zoster vaccination

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/shingles.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/shingles.html)

#### General Information

- Administer 2 doses of recombinant zoster vaccine (RZV) 2–6 months apart to adults aged 50 years or older regardless of past episode of herpes zoster or receipt of zoster vaccine live (ZVL)

- Administer 2 doses of RZV 2–6 months apart to adults who previously received ZVL at least 2 months after ZVL
- For adults aged 60 years or older, administer either RZV or ZVL (RZV is preferred)

#### Special populations

- ZVL is contraindicated for pregnant women and adults with severe immunodeficiency

### 6. Human papillomavirus vaccination

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/hpv.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/hpv.html)

#### General Information

- Administer human papillomavirus (HPV) vaccine to **females through age 26 years and males through age 21 years** (males aged 22 through 26 years may be vaccinated based on individual clinical decision)
- The number of doses of HPV vaccine to be administered depends on age at initial HPV vaccination
  - **No previous dose of HPV vaccine:** Administer 3-dose series at 0, 1–2, and 6 months (minimum intervals: 4 weeks between doses 1 and 2, 12 weeks between doses 2 and 3, and 5 months between doses 1 and 3; repeat doses if given too soon)
  - **Aged 9–14 years at HPV vaccine series initiation and received 1 dose or 2 doses less than 5 months apart:** Administer 1 dose
  - **Aged 9–14 years at HPV vaccine series initiation and received 2 doses at least 5 months apart:** No additional dose is needed

#### Special populations

- Adults with **immunocompromising conditions (including HIV infection)** through age 26 years: Administer 3-dose series at 0, 1–2, and 6 months
- **Men who have sex with men** through age 26 years: Administer 2- or 3-dose series depending on age at initial vaccination (see above); if no history of HPV vaccine, administer 3-dose series at 0, 1–2, and 6 months
- **Pregnant women** through age 26 years: HPV vaccination is not recommended during pregnancy, but there is no evidence that the vaccine is harmful and no intervention needed for women who inadvertently receive HPV vaccine while pregnant; delay remaining doses until after pregnancy; pregnancy testing is not needed before vaccination

### 7. Pneumococcal vaccination

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/pneumo.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/pneumo.html)

#### General Information

- Administer to immunocompetent adults aged 65 years or older 1 dose of 13-valent pneumococcal conjugate vaccine (PCV13), if not previously administered, followed by 1 dose of 23-valent pneumococcal polysaccharide vaccine (PPSV23) at least 1 year after PCV13; if PPSV23 was previously administered but not PCV13, administer PCV13 at least 1 year after PPSV23
- When both PCV13 and PPSV23 are indicated, administer PCV13 first (PCV13 and PPSV23 should not be administered during the same visit); additional information on vaccine timing is available at [www.cdc.gov/vaccines/vpd/pneumo/downloads/pneumo-vaccine-timing.pdf](http://www.cdc.gov/vaccines/vpd/pneumo/downloads/pneumo-vaccine-timing.pdf)

### Special populations

- Administer to adults aged 19 through 64 years with the following chronic conditions 1 dose of PPSV23 (at age 65 years or older, administer 1 dose of PCV13, if not previously received, and another dose of PPSV23 at least 1 year after PCV13 and at least 5 years after PPSV23):
  - Chronic heart disease (excluding hypertension)
  - Chronic lung disease
  - Chronic liver disease
  - Alcoholism
  - Diabetes mellitus
  - Cigarette smoking
- Administer to adults aged 19 years or older with the following indications 1 dose of PCV13 followed by 1 dose of PPSV23 at least 8 weeks after PCV13, and a second dose of PPSV23 at least 5 years after the first dose of PPSV23 (if the most recent dose of PPSV23 was administered before age 65 years, at age 65 years or older, administer another dose of PPSV23 at least 5 years after the last dose of PPSV23):
  - Immunodeficiency disorders (including B- and T-lymphocyte deficiency, complement deficiencies, and phagocytic disorders)
  - HIV infection
  - Anatomical or functional asplenia (including sickle cell disease and other hemoglobinopathies)
  - Chronic renal failure and nephrotic syndrome
- Administer to adults aged 19 years or older with the following indications 1 dose of PCV13 followed by 1 dose of PPSV23 at least 8 weeks after PCV13 (if the dose of PPSV23 was administered before age 65 years, at age 65 years or older, administer another dose of PPSV23 at least 5 years after the last dose of PPSV23):
  - Cerebrospinal fluid leak
  - Cochlear implant

## 8. Hepatitis A vaccination

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/hepa.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/hepa.html)

### General Information

- Administer to adults who have a specific risk (see below), or lack a risk factor but want protection, 2-dose series of single antigen hepatitis A vaccine (HepA; Havrix at 0 and 6–12 months or Vaqta at 0 and 6–18 months; minimum interval: 6 months) or a 3-dose series of combined hepatitis A and hepatitis B vaccine (HepA-HepB) at 0, 1, and 6 months; minimum intervals: 4 weeks between first and second doses, 5 months between second and third doses

### Special populations

- Administer HepA or HepA-HepB to adults with the following indications:
  - Travel to or work in countries with high or intermediate hepatitis A endemicity
  - Men who have sex with men
  - Injection or noninjection drug use
  - Work with hepatitis A virus in a research laboratory or with nonhuman primates infected with hepatitis A virus
  - Clotting factor disorders
  - Chronic liver disease

- Close, personal contact with an international adoptee (e.g., household or regular babysitting) during the first 60 days after arrival in the United States from a country with high or intermediate endemicity (administer the first dose as soon as the adoption is planned)
- Healthy adults through age 40 years who have recently been exposed to hepatitis A virus; adults older than age 40 years may receive HepA or HepA-HepB if hepatitis A immunoglobulin cannot be obtained

## 9. Hepatitis B vaccination

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/hepb.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/hepb.html)

### General Information

- Administer to adults who have a specific risk (see below), or lack a risk factor but want protection, 3-dose series of single antigen hepatitis B vaccine (HepB) or combined hepatitis A and hepatitis B vaccine (HepA-HepB) at 0, 1, and 6 months (minimum intervals: 4 weeks between doses 1 and 2 for HepB and HepA-HepB; between doses 2 and 3, 8 weeks for HepB and 5 months for HepA-HepB)

### Special populations

- Administer HepB or HepA-HepB to adults with the following indications:
  - Chronic liver disease (e.g., hepatitis C infection, cirrhosis, fatty liver disease, alcoholic liver disease, autoimmune hepatitis, alanine aminotransferase [ALT] or aspartate aminotransferase [AST] level greater than twice the upper limit of normal)
  - HIV infection
  - Percutaneous or mucosal risk of exposure to blood (e.g., household contacts of hepatitis B surface antigen [HBsAg]-positive persons; adults younger than age 60 years with diabetes mellitus or aged 60 years or older with diabetes mellitus based on individual clinical decision; adults in predialysis care or receiving hemodialysis or peritoneal dialysis; recent or current injection drug users; health care and public safety workers at risk for exposure to blood or blood-contaminated body fluids)
  - Sexual exposure risk (e.g., sex partners of HBsAg-positive persons; sexually active persons not in a mutually monogamous relationship; persons seeking evaluation or treatment for a sexually transmitted infection; and men who have sex with men [MSM])
  - Receive care in settings where a high proportion of adults have risks for hepatitis B infection (e.g., facilities providing sexually transmitted disease treatment, drug-abuse treatment and prevention services, hemodialysis and end-stage renal disease programs, institutions for developmentally disabled persons, health care settings targeting services to injection drug users or MSM, HIV testing and treatment facilities, and correctional facilities)
  - Travel to countries with high or intermediate hepatitis B endemicity

## 10. Meningococcal vaccination

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/mening.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/mening.html)

Special populations: Serogroups A, C, W, and Y meningococcal vaccine (MenACWY)

- Administer 2 doses of MenACWY at least 8 weeks apart and revaccinate with 1 dose of MenACWY every 5 years, if the risk remains, to adults with the following indications:
  - Anatomical or functional asplenia (including sickle cell disease and other hemoglobinopathies)
  - HIV infection
  - Persistent complement component deficiency
  - Eculizumab use
- Administer 1 dose of MenACWY and revaccinate with 1 dose of MenACWY every 5 years, if the risk remains, to adults with the following indications:
  - Travel to or live in countries where meningococcal disease is hyperendemic or epidemic, including countries in the African meningitis belt or during the Hajj
  - At risk from a meningococcal disease outbreak attributed to serogroup A, C, W, or Y
  - Microbiologists routinely exposed to *Neisseria meningitidis*
  - Military recruits
  - First-year college students who live in residential housing (if they did not receive MenACWY at age 16 years or older)

General Information: Serogroup B meningococcal vaccine (MenB)

- May administer, based on individual clinical decision, to young adults and adolescents aged 16–23 years (preferred age is 16–18 years) who are not at increased risk 2-dose series of MenB-4C (Bexsero) at least 1 month apart or 2-dose series of MenB-FHbp (Trumenba) at least 6 months apart
- MenB-4C and MenB-FHbp are not interchangeable

### Special populations: MenB

- Administer 2-dose series of MenB-4C at least 1 month apart or 3-dose series of MenB-FHbp at 0, 1–2, and 6 months to adults with the following indications:
  - Anatomical or functional asplenia (including sickle cell disease)
  - Persistent complement component deficiency
  - Eculizumab use
  - At risk from a meningococcal disease outbreak attributed to serogroup B
  - Microbiologists routinely exposed to *Neisseria meningitidis*

## 11. *Haemophilus influenzae* type b vaccination

[www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/hib.html](http://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/hib.html)

### Special populations

- Administer *Haemophilus influenzae* type b vaccine (Hib) to adults with the following indications:
  - Anatomical or functional asplenia (including sickle cell disease) or undergoing elective splenectomy: Administer 1 dose if not previously vaccinated (preferably at least 14 days before elective splenectomy)
  - Hematopoietic stem cell transplant (HSCT): Administer 3-dose series with doses 4 weeks apart starting 6 to 12 months after successful transplant regardless of Hib vaccination history

**Table. Contraindications and precautions for vaccines recommended for adults aged 19 years or older\***

The Advisory Committee on Immunization Practices (ACIP) recommendations and package inserts for vaccines provide information on contraindications and precautions related to vaccines. Contraindications are conditions that increase chances of a serious adverse reaction in vaccine recipients and the vaccine should not be administered when a contraindication is present. Precautions should be reviewed for potential risks and benefits for vaccine recipients.

**Contraindications and precautions for vaccines routinely recommended for adults**

Vaccine(s)	Contraindications	Precautions
All vaccines routinely recommended for adults	• Severe reaction, e.g., anaphylaxis, after a previous dose or to a vaccine component	• Moderate or severe acute illness with or without fever

**Additional contraindications and precautions for vaccines routinely recommended for adults**

Vaccine(s)	Additional Contraindications	Additional Precautions
IIV <sup>1</sup>		<ul style="list-style-type: none"> <li>History of Guillain-Barré syndrome within 6 weeks after previous influenza vaccination</li> <li>Egg allergy other than hives, e.g., angioedema, respiratory distress, lightheadedness, or recurrent emesis; or required epinephrine or another emergency medical intervention (IIV may be administered in an inpatient or outpatient medical setting and under the supervision of a health care provider who is able to recognize and manage severe allergic conditions)</li> </ul>
RIV <sup>1</sup>		<ul style="list-style-type: none"> <li>History of Guillain-Barré syndrome within 6 weeks after previous influenza vaccination</li> </ul>
Tdap, Td	<ul style="list-style-type: none"> <li>For pertussis-containing vaccines: encephalopathy, e.g., coma, decreased level of consciousness, or prolonged seizures, not attributable to another identifiable cause within 7 days of administration of a previous dose of a vaccine containing tetanus or diphtheria toxoid or acellular pertussis</li> </ul>	<ul style="list-style-type: none"> <li>Guillain-Barré syndrome within 6 weeks after a previous dose of tetanus toxoid-containing vaccine</li> <li>History of Arthus-type hypersensitivity reactions after a previous dose of tetanus or diphtheria toxoid-containing vaccine. Defer vaccination until at least 10 years have elapsed since the last tetanus toxoid-containing vaccine</li> <li>For pertussis-containing vaccine, progressive or unstable neurologic disorder, uncontrolled seizures, or progressive encephalopathy (until a treatment regimen has been established and the condition has stabilized)</li> </ul>
MMR <sup>2</sup>	<ul style="list-style-type: none"> <li>Severe immunodeficiency, e.g., hematologic and solid tumors, chemotherapy, congenital immunodeficiency or long-term immunosuppressive therapy<sup>3</sup>, human immunodeficiency virus (HIV) infection with severe immunocompromise</li> <li>Pregnancy</li> </ul>	<ul style="list-style-type: none"> <li>Recent (within 11 months) receipt of antibody-containing blood product (specific interval depends on product)<sup>4</sup></li> <li>History of thrombocytopenia or thrombocytopenic purpura</li> <li>Need for tuberculin skin testing<sup>5</sup></li> </ul>
VAR <sup>2</sup>	<ul style="list-style-type: none"> <li>Severe immunodeficiency, e.g., hematologic and solid tumors, chemotherapy, congenital immunodeficiency or long-term immunosuppressive therapy<sup>3</sup>, HIV infection with severe immunocompromise</li> <li>Pregnancy</li> </ul>	<ul style="list-style-type: none"> <li>Recent (within 11 months) receipt of antibody-containing blood product (specific interval depends on product)<sup>4</sup></li> <li>Receipt of specific antiviral drugs (acyclovir, famciclovir, or valacyclovir) 24 hours before vaccination (avoid use of these antiviral drugs for 14 days after vaccination)</li> </ul>
ZVL <sup>2</sup>	<ul style="list-style-type: none"> <li>Severe immunodeficiency, e.g., hematologic and solid tumors, chemotherapy, congenital immunodeficiency or long-term immunosuppressive therapy<sup>3</sup>, HIV infection with severe immunocompromise</li> <li>Pregnancy</li> </ul>	<ul style="list-style-type: none"> <li>Receipt of specific antiviral drugs (acyclovir, famciclovir, or valacyclovir) 24 hours before vaccination (avoid use of these antiviral drugs for 14 days after vaccination)</li> </ul>
HPV vaccine		<ul style="list-style-type: none"> <li>Pregnancy</li> </ul>
PCV13	<ul style="list-style-type: none"> <li>Severe allergic reaction to any vaccine containing diphtheria toxoid</li> </ul>	

- For additional information on use of influenza vaccines among persons with egg allergy, see: CDC. Prevention and control of seasonal influenza with vaccines: recommendations of the Advisory Committee on Immunization Practices—United States, 2016–17 influenza season. MMWR. 2016;65(RR-5):1–54. Available at [www.cdc.gov/mmwr/volumes/65/rr/r6505a1.htm](http://www.cdc.gov/mmwr/volumes/65/rr/r6505a1.htm).
- MMR may be administered together with VAR or ZVL on the same day. If not administered on the same day, separate live vaccines by at least 28 days.
- Immunosuppressive steroid dose is considered to be daily receipt of 20 mg or more prednisone or equivalent for 2 or more weeks. Vaccination should be deferred for at least 1 month after discontinuation of immunosuppressive steroid therapy. Providers should consult ACIP recommendations for complete information on the use of specific live vaccines among persons on immune-suppressing medications or with immune suppression because of other reasons.
- Vaccine should be deferred for the appropriate interval if replacement immune globulin products are being administered. See: Best practices guidance of the Advisory Committee on Immunization Practices (ACIP). Available at [www.cdc.gov/vaccines/hcp/acip-recs/general-recs/index.html](http://www.cdc.gov/vaccines/hcp/acip-recs/general-recs/index.html).
- Measles vaccination may temporarily suppress tuberculin reactivity. Measles-containing vaccine may be administered on the same day as tuberculin skin testing, or should be postponed for at least 4 weeks after vaccination.

\* Adapted from: CDC. Table 6. Contraindications and precautions to commonly used vaccines. General recommendations on immunization: recommendations of the Advisory Committee on Immunization Practices. MMWR. 2011;60(No. RR-2):40–1 and from: Hamborsky J, Kroger A, Wolfe S, eds. Appendix A. Epidemiology and prevention of vaccine preventable diseases. 13th ed. Washington, DC: Public Health Foundation, 2015. Available at [www.cdc.gov/vaccines/pubs/pinkbook/index.html](http://www.cdc.gov/vaccines/pubs/pinkbook/index.html).

**Abbreviations of vaccines**

IIV	inactivated influenza vaccine	VAR	varicella vaccine	HepA	hepatitis A vaccine
RIV	recombinant influenza vaccine	RZV	recombinant zoster vaccine	HepA-HepB	hepatitis A and hepatitis B vaccines
Tdap	tetanus toxoid, reduced diphtheria toxoid, and acellular pertussis vaccine	ZVL	zoster vaccine live	HepB	hepatitis B vaccine
Td	tetanus and diphtheria toxoids	HPV vaccine	human papillomavirus vaccine	MenACWY	serogroups A, C, W, and Y meningococcal vaccine
MMR	measles, mumps, and rubella vaccine	PCV13	13-valent pneumococcal conjugate vaccine	MenB	serogroup B meningococcal vaccine
		PPSV23	23-valent pneumococcal polysaccharide vaccine	Hib	<i>Haemophilus influenzae</i> type b vaccine

# 2018 Adult Immunization Updates – Zoster Vaccination

- Recombinant zoster vaccine (RZV) licensed by FDA on October 20
- Administer 2 doses RZV 2–6 mos apart to adults  $\geq 50$ y regardless of past herpes zoster or receipt of zoster vaccine live (ZVL)
- Administer RZV 2–6 mos apart to adults who previously received ZVL at least 2 mos after ZVL
- For adults  $\geq 60$ y, administer either RZV or ZVL (RZV is preferred)
- Administer RZV to age-eligible adults with chronic health conditions including diabetes, chronic heart/lung/liver/kidney ds, asplenia, complement deficiencies
- Pending considerations on use of RZV in immunocompromising conditions including HIV infection
- No data on pregnant women (consider delay)

# 2018 Adult Immunization Updates – Mumps Vaccination

- Routine mumps vaccination recommendations
  - Children – 2 doses MMR during K-12 (12–15 mos and 4–6 yrs for measles)
  - Adults – 2 doses MMR for high risk (students at post-high school educational institutions, health care personnel, international travelers)
- Multiple outbreaks of mumps and high numbers of reported cases since 2015, many among young adults who received 2 doses MMR
- Updated ACIP recommendations on use of MMR during mumps outbreak
  - Administer 1 dose MMR to persons who previously received  $\leq 2$  doses mumps-containing vaccine and identified by public health authority to be at increased risk during mumps outbreak

# ACIP Updates Hepatitis B Prevention

- New or updated ACIP recommendations for children
  - Universal HepB within 24 hrs of birth for medically stable infants weighing  $\geq 2,000$  g
  - Test HBsAg(+) pregnant women for hepatitis B virus deoxyribonucleic acid (HBV DNA)
  - Test postvaccination serology for infants whose mother's HBsAg status unknown indefinitely (e.g., when a parent or person with lawful custody surrenders an infant confidentially shortly after birth)
  - Single-dose revaccination for infants born to HBsAg(+) women when not respond to initial vaccine series
  - Removal of permissive language for delaying birth dose after hospital discharge
- Relevant for adults
  - Vaccinate persons with chronic liver disease (hepatitis C virus [HCV] infection, 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)

# New Hepatitis B Vaccine for Adults

- FDA licensed (Nov 9, 2017) single antigen HepB (HEPLISAV-B, Dynavax Technologies Corp.) for all HBV subtypes for  $\geq 18y$
- 5<sup>th</sup> inactivated HepB in U.S. (Engerix-B, Recombivax HB, Pediarix, Twinrix)
- Contains yeast-derived recombinant HBsAg with 1018 adjuvant (immunostimulatory sequences) that binds Toll-like receptor 9 to stimulate directed immune response
- 2 doses 1 month apart

# Heplisav-B – Seroprotection and Safety

## ■ Immunogenicity

- 90.0%–100% vs. 70.5%–90.2% in comparison group (3 doses Engerix-B)
- Diabetes Type II: 90.0% vs. 65.1% (3 doses Engerix-B)
- Chronic kidney disease: 89.9% (3 doses) vs. 81.1% (4 double doses Engerix-B)

## ■ Safety and reactogenicity

- Mild and serious adverse events similar
  - Mild: 45.6% vs. 45.7% (Engerix-B)
  - Serious: 5.4% vs. 6.3% (Engerix-B)
- Cardiovascular events
  - 0.27% vs. 0.14% (Engerix-B)
- Potentially immune-mediated adverse events (e.g., granulomatosis with polyangiitis, Grave's disease)
  - 0.1%–0.2% vs. 0%–0.7% (Engerix-B)

Jackson S, Lentino J, Kopp J, et al. Immunogenicity of a two-dose investigational hepatitis B vaccine, HBsAg-1018, using a toll-like receptor 9 agonist adjuvant compared with a licensed hepatitis B vaccine in adults. *Vaccine* 2017; 36:668-74

Janssen R, Bennett S, Namini H, et al. Immunogenicity and Safety of Two Doses of Investigational Heplisav Compared to Three Doses of Licensed Hepatitis B Vaccine (Engerix-B) in Two Phase 3 Trials. *Journal of Hepatology* 2013; 58(Suppl 1):S574

HEPLISAV-B™ [Hepatitis B Vaccine (Recombinant), Adjuvanted] package insert. <https://www.fda.gov/downloads/BiologicsBloodVaccines/Vaccines/ApprovedProducts/UCM584762.pdf>

# Pending ACIP Deliberation on Heplisav-B

- Use of Heplisav-B in prevention of hepatitis B for  $\geq 18$ y up for ACIP vote in February 2018
- Other considerations



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

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