VACCINATE ADULTS!

from the Immunization Action Coalition — www.immunize.org

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Ask the Experts

The Immunization Action Coalition extends thanks to our experts, medical officer Andrew T. Kroger, MD, MPH, and nurse educator Donna L. Weaver, RN, MN, both with the National Center for Immunization and Respiratory Diseases at the Centers for Disease Control and Prevention (CDC).

Tdap and Td vaccines

For a person entering a long-term-care facility at age 70 or older, if we cannot document that the resident has had a primary series of three doses of tetanus-containing vaccine, is the right course of action upon admission to give a Tdap first, then a Td in 1 to 2 months, followed by a Td in 6 to 12 months, and then a Td booster every 10 years?

Your understanding of the general Td/Tdap recommendation is correct, and this is the schedule that should be followed for persons 7 years old and older who have never received tetanus-containing vaccine or who cannot provide documentation of prior vaccination. Be sure to document doses administered so a primary series does not need to be repeated in the future.

Use Standing Orders to Increase Coverage Rates and Protect Patients

Using standing orders protocols for vaccination in your medical practice allows appropriately trained healthcare professionals – who are permitted to do so under state law – to assess a patient's need for vaccination, determine if there are contraindications or precautions, and then to administer vaccine without obtaining a physician's written or verbal order for an individual patient.

Numerous studies have shown that standing orders, carried out by nurses or other qualified healthcare professionals, are one of the most consistently effective means for increasing vaccination rates and reducing missed opportunities for vaccination, thereby improving quality of care.

CDC's Advisory Committee on Immunization Practices (ACIP) has recommended the use of standing orders to increase adult vaccination rates since 2000. See www.cdc.gov/mmwr/preview/mmwrhtml/rr4901a2.htm.

The Community Preventive Services Task Force (Task Force) recommends standing orders for vaccinations based on strong evidence of effectiveness in improving vaccination rates among adults and children, when used alone or with additional interventions, and across a range of settings and populations. See www.thecommunity guide.org/vaccines/standingorders.html.

Exactly who is authorized to administer vaccines under standing orders varies by state law. To find out which medical personnel are permitted to administer vaccines under standing orders in your state, contact your state immunization program manager. Contact information is available at www.immunize.org/coordinators.

If you are interested in starting a standing orders program in your practice setting, the Immunization Action Coalition (IAC) has materials

If a patient is receiving a tetanus-containing vaccine after an injury and there is no history of any prior tetanus vaccine (e.g., an Amish person who has previously declined vaccination), how much tetanus protection will one dose provide? Also, what is the time frame that the tetanus toxoid needs to be given following an injury?

One dose of tetanus toxoid-containing vaccine (Tdap or Td) provides little or no protection. That is why tetanus immune globulin (TIG) is also recommended in this situation. See "The Pink Book" section titled Wound Management at www.cdc.gov/vaccines/pubs/pinkbook/tetanus.html. As

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available that help make standing orders easy to implement. "Using Standing Orders for Administering Vaccines: What You Should Know" is a one-page article describing the basics of standing orders. See www.immunize.org/catg.d/p3066.pdf.

Standing Orders Templates for Routinely Recommended Vaccines Are Available on IAC's Website

IAC has created standing orders templates for all routinely recommended vaccines for administration to children, teens, and adults. They are all available online and are modifiable in any way you choose to suit your practice's needs. These standing orders templates are based on ACIP vaccine recommendations and are reviewed for technical accuracy by CDC staff. IAC updates its standing orders protocols whenever ACIP makes changes in vaccine recommendations.

You can find IAC's standing orders templates on IAC's Standing Orders web page at www.immunize. org/standing-orders. A few examples follow:

- "Standing Orders for Administering Influenza Vaccine to Adults"
 (www.immunize.org/catg.d/p3074.pdf)
- "Standing Orders for Administering Influenza Vaccine to Children and Adolescents"

(www.immunize.org/catg.d/p3074a.pdf)

- "Standing Orders for Administering Tdap/Td Vaccine to Adults" (www.immunize.org/catg.d/p3078.pdf)
- "Medical Management of Vaccine Reactions in Children and Teens" (www.immunize.org/catg.d/p3082a.pdf)
- "Medical Management of Vaccine Reactions in Adult Patients" (www.immunize.org/catg.d/p3082.pdf)

Access all IAC's standing orders templates at www.immunize.org/standing-orders.

To be notified when new or revised templates become available, subscribe to IAC's free weekly news service, *IAC Express*, at www.immunize. org/subscribe, which is sent to more than 50,000 healthcare professionals every Wednesday. ◆

Immunization questions?

- ► Email nipinfo@cdc.gov
- Call your state health department (phone numbers at www.immunize. org/coordinators)

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online at www.immunize.org/va Immunization Action Coalition

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far as timing, the toxoid and TIG should be given as soon as possible.

Hepatitis B vaccine

I work in occupational health and have some patients who are off schedule for their hepatitis B vaccine series. They came back for dose #2 in 4 to 6 months rather than getting it 1 month later. In this situation, what is the correct timing for dose #3? And how long must the interval be between doses before I am required to restart the series?

The minimal intervals for hepatitis B vaccine are at least 4 weeks between doses #1 and #2, at least 8 weeks between doses #2 and #3, and at least 16 weeks between doses #1 and #3. Since in your cases 16 weeks or more have elapsed since dose #1, you should schedule dose #3 to be given 8 weeks after dose #2. It is not necessary to restart the series because of an extended interval between doses, no matter how long.

MMR vaccine

Would you consider a healthcare worker with two documented doses of MMR vaccine to be immune, even if the serology for one or more of the antigens comes back negative?

Yes. Healthcare personnel (HCP) with two documented doses of MMR vaccine are considered to be immune, regardless of the results of a subsequent serologic test for measles, mumps, or rubella. Documented age-appropriate vaccination supersedes the results of subsequent serologic testing. HCP who do not have documentation of MMR vaccination and whose serologic test is interpreted as "indeterminate" or "equivocal" should be considered not immune and should receive two doses of MMR. ACIP does not recommend serologic testing after vaccination. For more information, see ACIP's recommendations on the use of MMR at www.cdc.gov/mmwr/pdf/rr/rr6204.pdf, page 22.

Meningococcal vaccines

I have an otherwise healthy 26-year-old patient with HIV infection who received one dose of meningococcal conjugate vaccine (MCV4, MenACWY:

Menactra, Sanofi Pasteur; Menveo, GSK) three years ago. Should he receive one or two doses now? Will he need booster doses later?

It is not necessary to start the MenACWY series again. Give the person one dose of MenACWY vaccine now. This dose represents a delayed second dose in the primary series, which is recommended in patients with HIV regardless of the presence of another condition that increases the risk of meningococcal disease. In the absence of another condition that increases the risk of meningococcal disease (such as asplenia), booster doses are not recommended.

Are microbiologists recommended to receive meningococcal B vaccine? And if so, how frequently?

ACIP recommends that microbiologists who work with meningococcus bacteria in a laboratory receive both MenB vaccine and MenACWY vaccine. MenB can be given at the same time as any other vaccine. You can administer either two doses of Bexsero 4 weeks apart, or three doses of Trumenba on a 0-, 1–2-, and 6-month schedule. In April 2016, the FDA also approved a 2-dose schedule for Trumenba with doses given at 0 and 6 months. There is currently no recommendation for a booster dose of MenB vaccine for any age or risk group.

Pneumococcal vaccines

Does a patient younger than age 65 years who smokes marijuana on a daily basis, but doesn't smoke cigarettes, need to receive pneumococcal polysaccharide (PPSV) vaccine?

No. ACIP does not identify people who smoke marijuana but not cigarettes as being at increased risk for pneumococcal disease or as being in a risk group for PPSV (Pneumovax 23, Merck) vaccination.

Is a patient younger than age 65 years who recently had a prostatectomy with lymph node dissection for prostate cancer a candidate for PPSV? The patient is believed to be cancer-free and is on no chemotherapy.

In the absence of "generalized malignancy" (which is generally considered to mean disseminated cancer) or immunosuppression, a recent history of prostate cancer surgery alone is not an indication for PPSV.

Ask the Experts...continued on page 3 ▶

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IAC's "Ask the Experts" team from the Centers for Disease Control and Prevention





Andrew T. Kroger, MD, MPH

Donna L. Weaver, RN, MN

I have patients who are in their 70s and 80s and remember getting a pneumococcal vaccine a few years ago. Should we assume that this was PPSV? Should I assume that it was given before the 65th birthday?

You can accept a patient's verbal report of PPSV* and it is reasonable to assume that PPSV was the pneumococcal vaccine that was administered. If the patient's history suggests that this dose was given on or after age 65 years, it can be counted as the one dose recommended for this age group. If it has been a year or longer since this dose, pneumococcal conjugate vaccine (PCV, Prevnar 13, Pfizer) should be administered now. If there is any question about the age at which the dose was given, it is reasonable to give PCV now then give a dose of PPSV in 1 year.

*Note: a personal report (undocumented) of receipt of a vaccination is acceptable only for PPSV and influenza vaccines. All other vaccines must be documented with a written, dated record.

Zoster vaccine

If a patient received dose #1 of varicella vaccine at age 60 years, should we administer zoster vaccine as dose #2?

The action taken depends on why varicella vaccine was given in the first place. If it was given because the

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person tested negative for varicella antibody, then the next dose should be varicella vaccine. If the varicella vaccine was given in error (i.e., without serologic testing), then zoster vaccine (Zostavax, Merck) should be given.

A dose of zoster vaccine was inadvertently given to a patient receiving chemotherapy for colon cancer. We realize this was an error, so please advise us on what to do now.

Zoster vaccine is given to people who presumably had chickenpox earlier in life and so have immunity to varicella virus. The cancer chemotherapy will not change the person's immunity to varicella virus. However, the patient should be monitored for the next two weeks for symptoms that might indicate an adverse reaction, such as fever and rash. If symptoms suggestive of varicella develop, the patient can be started on antiviral therapy, such as acyclovir.

Healthcare personnel

Which vaccines are recommended for healthcare personnel (HCP)?

ACIP recommends that people working in healthcare settings be vaccinated against influenza, hepatitis B, measles, mumps, rubella, varicella, and pertussis. For measles, mumps, rubella, and varicella, serologic evidence of immunity is an acceptable substitute for documentation of vaccination. In addition, microbiologists working in a laboratory should receive meningococcal conjugate and meningococcal serogroup B vaccines. In rare cases, some laboratory personnel should also receive polio and typhoid vaccines. For more information, see www.cdc.gov/mmwr/pdf/rr/ rr6007.pdf.

Should HCP be vaccinated routinely against hepatitis A?

No. A number of studies have shown that HCP are not at increased risk of hepatitis A virus (HAV) infection because of their occupation. However, if he/she is going to work (or vacation) in a country with a high or intermediate endemic rate of HAV infection, he/she is at risk of HAV infection and should be vaccinated. The only HCP for whom hepatitis A vaccine is routinely recommended are those who work with primates or live HAV.

I am a nurse who received the hepatitis B vaccine series more than 10 years ago and had a positive follow-up antibody titer for anti-HBs (at least 10 mIU/mL) at that time. At present, my titer is negative (less than 10 mIU/ mL). What should I do now?

Nothing needs to be done. Data show that vaccineinduced anti-HBs levels might decline over time; however, immune memory (anamnestic anti-HBs response) remains intact following immunization. People with anti-HBs concentrations that decline to less than 10 mIU/mL are still protected against HBV infection. For HCP with normal immune status who

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have demonstrated adequate anti-HBs (at least 10 mIU/mL) following full vaccination, booster doses of vaccine or periodic anti-HBs testing are not recommended.

Storage and handling

How long is a vaccine viable if it has been stored in the refrigerator in a syringe?

Disposable syringes are meant for administration of immunobiologics, not for storage. CDC recommends that vaccines that have been drawn into syringes in the work setting be discarded at the end of the clinic day. Manufacturer-filled syringes that have not been activated (i.e., have not had the needle guard removed or a needle attached) may be kept and used until their expiration date.

Miscellaneous questions

I understand that certain vaccines contain aluminum and/or formaldehyde. Could you

provide me with the research showing the "safe" amounts of aluminum and formaldehyde that can be injected per pound of body weight?

Aluminum is common in the environment and is ingested every day in larger amounts than the amount in vaccines. Formaldehyde is a compound that the human body normally produces in higher amounts than are injected with vaccines.

The FDA performed two studies directly related to your question which demonstrate the amounts injected with vaccines are well below what are considered toxic levels. The following citation is about aluminum in vaccines: www.fda.gov/Biologics-BloodVaccines/ScienceResearch/ucm284520.htm

Regarding formaldehyde, the FDA did the following study, and estimated that less formaldehyde is received from vaccines by infants than infants produce themselves: www.accessdata.fda.gov/scripts/publications/search_result_record.cfm?id=45918.

Information on vaccine additives is available on the CDC website at www.cdc.gov/vaccines/vac-gen/additives.htm and on the FDA website at www.fda.gov/biologicsbloodvaccines/safetyavailability/vaccinesafety/ucm187810.htm.

To find more than

Ask the Experts Q&As answered by CDC experts,

visit

www.immunize.org/ askexperts

About IAC's Question of the Week

Each week, *IAC Express* highlights a new, topical, or important-to-reiterate Q&A. This feature is a cooperative venture between IAC and CDC. William L. Atkinson, MD, MPH, IAC's associate director for immunization education, chooses a new Q&A to feature every week from a set of Q&As prepared by experts at CDC's National Center for Immunization and Respiratory Diseases.

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Laminated U.S. Immunization Schedules

Purchase IAC's laminated versions of the 2016 U.S. immunization schedules for adults and children (0–18 years old). Both are laminated and washable for heavy-duty use, complete with essential footnotes, and printed in color for easy reading.

More information and discount pricing options are available online at www.immunize.org/shop/laminated-schedules.asp or see the order form on page 16.



To submit an "Ask the Experts" question...

You can email your questions about immunization to us at admin@immunize.org. IAC will respond to your inquiry. Because we receive hundreds of emails each month, we cannot guarantee that we will use your question in "Ask the Experts." IAC works with CDC to compile new Q&As for our publications based on commonly asked questions. Most of the questions are thus a composite of several inquiries.

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If you find an error, please notify us immediately by sending an email message to admin@immunize.org. We publish notification of significant errors in our email announcement service, *IAC Express*. Be sure you're signed up for this service. To subscribe, visit www.immunize.org/subscribe.

Vaccine Highlights

Recommendations, schedules, and more

Editor's note: The information in Vaccine Highlights is current as of May 23, 2016.

Next ACIP meetings

The Advisory Committee on Immunization Practices (ACIP) is comprised of 15 national experts who advise CDC on the appropriate use of vaccines.

ACIP meets three times a year in Atlanta; meetings are open to the public and viewable online via live webcast. The next meetings will be held on June 22–23 and Oct. 19–20. For more information, visit www.cdc.gov/vaccines/acip.

ACIP periodically issues recommendations on the use of vaccines; they are published and readily available in the *Morbidity and Mortality Weekly Report (MMWR)*. Clinicians who vaccinate should have a current set for reference. Here are sources:

- Download from IAC's website: www.immunize. org/acip
- Download from CDC's website: www.cdc.gov/ vaccines/hcp/acip-recs

VIS news

On March 31, CDC released final versions of the HPV9 (Gardasil 9) and the Meningococcal ACWY Vaccine Information Statements (VISs). Neither differs significantly from the previous VIS, but both are now final, rather than interim, editions. CDC encourages providers to begin using these VISs immediately, but stocks of the previous editions may still be used until supplies are exhausted. The two new VISs, along with many translations, are available here:

- HPV 9 www.immunize.org/vis/vis_hpv_gardasil_9.asp
- Meningococcal ACWY www.immunize. org/vis/vis_meningococcal_mcv_mpsv.asp

To access all VISs in more than 35 languages, visit IAC's VIS web section at www.immunize.org/vis.

FDA vaccine news

On April 14, FDA approved a change for Trumenba MenB vaccine (Pfizer) to include a two-dose schedule (doses administered at 0 and 6 months) as well as a modification of the three-dose schedule from administration at 0, 2, and 6 months to administration at 0, 1–2, and 6 months. Trumenba is approved for use in individuals age 10 through 25 years for active immunization to prevent invasive disease caused by *Neisseria meningitidis* serogroup B. The updated package insert is available at http://labeling.pfizer.com/ShowLabeling.aspx?id=1796.

In April, FDA published an update on distribution of yellow fever vaccine during the current shortage. According to the FDA's "CBER-Regulated Products: Current Shortages" website, YF-VAX (Sanofi Pasteur) is being provided only to customers who have patients planning to travel within 30 days to an area where yellow fever vaccine is required or recommended. This measure is necessary to responsibly manage the limited U.S. supply of YF-VAX. To obtain vaccine, providers must call Sanofi Pasteur (800-822-2463) and speak to a customer service representative.

CDC news

CDC's 47th National Immunization Conference will be held Sept. 13–15, in Atlanta. For more information, visit www.cdc.gov/vaccines/events/nic/index.html.

Measles outbreak news

As of the time of this writing (May 10), the measles outbreak in Shelby County (Memphis), Tennessee, stands at seven cases. On April 21, the Shelby County Health Department (SCHD) received confirmation of measles infection in two county residents who both developed rash on April 9 and were hospitalized briefly before recovering. The two local patients lived in different parts of the county and had no connection to one another; neither had traveled or reported contact with any ill travelers in the weeks prior to illness onset. Genotyping of the virus suggested the same source. Since then, five additional persons have been diagnosed with measles; four of the five were known contacts of the two initial cases.

The date of rash onset and isolation of the most recent case was May 4; known contacts to this case will be monitored through May 25. Of the cases, SCHD reports that six were unimmunized; one had received one dose of vaccine.

SCHD, with assistance from the Tennessee Department of Health, has followed up on hundreds of contacts exposed in healthcare and other settings. SCHD has set up a hotline for public calls, posted measles educational information to its website, and begun auditing student records in area school systems in preparation for the possibility of a school exposure. Education efforts also have been aimed at ensuring healthcare providers recognize and respond properly to a suspected measles case and that they have appropriate documentation of immunity to measles before an occupational exposure occurs.

For more information, visit www.schdresponse.com/content/measlesoutbreak. •

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Adenovirus	MMR
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For a ready-to-print version of this table for posting in your practice, go to www.immunize. org/catg.d/p2029.pdf.



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Recommended Adult Immunization Schedule – United States, 2016

Note: These recommendations <u>must</u> be read with the footnotes that follow containing number of doses, intervals between doses, and other important information.

Figure 1. Recommended immunization schedule for adults ages 19 years and older, by vaccine and age group¹

Vaccine	19–21 years	22-26 years	27–49 years	50–59 years	60-64 years	≥65 years
Influenza ^{2,*}		1 dose annually				
Tetanus, diphtheria, pertussis (Td/Tdap) ^{3,*}		Substitute	Tdap for Td once,	then Td booster	every 10 yrs	
Varicella ^{4,*}			2 d	oses		
Human papillomavirus (HPV) Female ^{5,*}	3 do	ses				
Human papillomavirus (HPV) Male ^{5,*}	3 do	ses				
Zoster ⁶					1 d	ose
Measles, mumps, rubella (MMR) ^{7,*}	1 or 2 doses depending on indication					
Pneumococcal 13-valent conjugate (PCV13) ^{8,*}					1.	lose
Pneumococcal 23-valent polysaccharide (PPSV23) ⁸			1 or 2 doses depe	ending on indica	tion	1 dose
Hepatitis A ^{9,*}			2 or 3 doses depe	ending on vaccin	e	
Hepatitis B ^{10,*}	3 doses					
Meningococcal 4-valent conjugate (MenACWY) or polysaccharide (MPSV4) ^{11,*}	1 or more doses depending on indication					
Meningococcal B (MenB) ¹¹	2 or 3 doses depending on vaccine					
Haemophilus influenzae type b (Hib) ^{12,*}		1 or 3 doses depending on indication				

Figure 2. Vaccines that might be indicated for adults ages 19 years and older based on medical and other indications1

Indication▶		Immuno- compromising conditions (excluding HIV infec-	HIV infection (cells/μL) ^{4,6,7}		Men who	Kidney failure, end-stage renal disease, on	Heart disease, chronic lung disease, chronic	Asplenia and persistent complement component defi-	Chronic liver		Healthcare
Vaccine ▼	Pregnancy	tion]) ^{4,6,7,8,13}	<200	≥200	men (MSM)	hemodialysis	alcoholism	ciencies ^{8,11,12}	disease	Diabetes	personnel
Influenza ^{2,*}						I dose annua	ally				
Td/Tdap³,*	1 dose Tdap each pregnancy				Substitu	te Tdap for Td on	ce, then Td booste	er every 10 yrs			
Varicella ^{4,*}	(Contraindicated					2 de	oses			
HPV Female ^{5,*}		3 doses through age 26 yrs						3 doses through	age 26 yrs		
HPV Male ^{5,*}		3 doses through age 26 yrs				3 doses through age 21 yrs					
Zoster ⁶	(Contraindicated 1 dose									
MMR ^{7,*}	(Contraindicated				1	or 2 doses deper	nding on indication	n		
PCV13 ^{8,*}						1 dose					
PPSV23 ⁸					1, 2, or 3	doses depending	on indication				
Hepatitis A ^{9,*}					2 or 3	doses depending	on vaccine				
Hepatitis B ^{10,*}						`	3 doses				
MenACWY or MPSV4 ^{11,*}					1 or more	doses depending	g on indication				
MenB ¹¹		2 or 3 doses depending on vaccine									
Hib ^{12,*}		3 doses post-HSCT recipients only					1 dose				
documentation	d for all persons of vaccination, or	nsation Program. who meet the age rlack evidence of pass of past episode	ast infection; z				s with a risk facto le, or other indication		o recommendatio	on	Contraindica

These schedules indicate the recommended age groups and medical indications for which administration of currently licensed vaccines is commonly recommended for adults ages 19 years and older, as of February 2016. For all vaccines being recommended on the Adult Immunization Schedule: a vaccine series does not need to be restarted, regardless of the time that has elapsed between doses. Licensed combination vaccines may be used whenever any components of the combination are indicated and when the vaccine's other components are not contraindicated. For detailed recommendations on all vaccines, including those used primarily for travelers or that are issued during the year, consult the manufacturers' package inserts and the complete statements from the Advisory Committee on Immunization Practices (www.cdc.gov/vaccines/hcp/acip-recs/index.html). Use of trade names and commercial sources is for identification only and does not imply endorsement by the U.S. Department of Health and Human Services.

The recommendations in this schedule were approved by the Centers for Disease Control and Prevention's (CDC) Advisory Committee on Immunization Practices (ACIP), the American Academy of Family Physicians (AAFP), the American College of Physicians (ACP), American College of Obstetricians and Gynecologists (ACOG), and American College of Nurse-Midwives (ACNM).

1. Additional Information

- Additional guidance for the use of the vaccines described in this supplement is available at www.cdc.gov/vaccines/hcp/acip-recs/index.html.
- Information on vaccination recommendations when vaccination status is unknown and other general immunization information can be found in the General Recommendations on Immunization at www.cdc.gov/mmwr/preview/ mmwrhtml/rr6002a1.htm.
- Information on travel vaccine requirements and recommendations (e.g., for hepatitis A and B, meningococcal, and other vaccines) is available at wwwnc. cdc.gov/travel/destinations/list.
- Additional information and resources regarding vaccination of pregnant women can be found at www.cdc.gov/vaccines/adults/rec-vac/pregnant.html.

2. Influenza vaccination.

- Annual vaccination against influenza is recommended for all persons age 6 months or older. A list of currently available influenza vaccines can be found at www.cdc.gov/flu/protect/vaccine/vaccines.htm.
- Persons age 6 months and older, including pregnant women can receive the inactivated influenza vaccine (IIV). An age-appropriate IIV formulation should be used.
- Intradermal IIV is an option for persons ages 18 through 64 years.
- High-dose IIV is an option for persons ages 65 years or older.
- Live-attenuated influenza vaccine (LAIV [FluMist]) is an option for healthy, non-pregnant persons ages 2 through 49 years.
- Recombinant influenza vaccine (RIV [Flublok]) is approved for persons ages 18 years or older.
- RIV, which does not contain any egg protein, may be administered to persons ages 18 years or older with egg allergy of any severity; IIV may be used with additional safety measures for persons with hives-only allergy to eggs.
- Health care personnel who care for severely immunocompromised persons who require care in a protected environment should receive IIV or RIV; health care personnel who receive LAIV should avoid providing care for severely immunosuppressed persons for 7 days after vaccination.

3. Tetanus, diphtheria, and acellular pertussis (Td/Tdap) vaccination.

- Administer 1 dose of Tdap vaccine to pregnant women during each pregnancy (preferably during 27 to 36 weeks' gestation), regardless of interval since prior Td or Tdap vaccination.
- Persons age 11 years or older who have not received Tdap vaccine or for whom vaccine status is unknown should receive a dose of Tdap followed by tetanus and diphtheria toxoids (Td) booster doses every 10 years thereafter. Tdap can be administered regardless of interval since the most recent tetanus or diphtheria toxoid-containing vaccine.
- Adults with an unknown or incomplete history of completing a 3-dose primary vaccination series with Td-containing vaccines should begin or complete a primary vaccination series including a Tdap dose.
- For unvaccinated adults, administer the first 2 doses at least 4 weeks apart and the third dose 6 to 12 months after the second.
- For incompletely vaccinated (i.e., less than 3 doses) adults, administer remaining doses.
- Refer to the ACIP statement for recommendations for administering Td/Tdap as prophylaxis in wound management (see footnote 1).

4. Varicella vaccination.

- All adults without evidence of immunity to varicella (as defined below) should receive 2 doses of single-antigen varicella vaccine or a second dose if they have received only 1 dose.
- Vaccination should be emphasized for those who have close contact with
 persons at high risk for severe disease (e.g., health care personnel and
 family contacts of persons with immunocompromising conditions) or are at
 high risk for exposure or transmission (e.g., teachers; child care employees;
 residents and staff members of institutional settings, including correctional
 institutions; college students; military personnel; adolescents and adults
 living in households with children; nonpregnant women of childbearing age;
 and international travelers).

- Pregnant women should be assessed for evidence of varicella immunity.
 Women who do not have evidence of immunity should receive the first dose of varicella vaccine upon completion or termination of pregnancy and before discharge from the health care facility. The second dose should be administered 4 to 8 weeks after the first dose.
- Evidence of immunity to varicella in adults includes any of the following: 1) documentation of 2 doses of varicella vaccine at least 4 weeks apart; 2) U.S.-born before 1980, except health care personnel and pregnant women; 3) history of varicella based on diagnosis or verification of varicella disease by a health care provider; 4) history of herpes zoster based on diagnosis or verification of herpes zoster disease by a health care provider; or 5) laboratory evidence of immunity or laboratory confirmation of disease.

5. Human papillomavirus (HPV) vaccination.

- Three HPV vaccines are licensed for use in females (bivalent HPV vaccine [HPV2], quadrivalent HPV vaccine [HPV4], and 9-valent HPV vaccine [HPV9]), and two HPV vaccines are licensed for use in males (HPV4 and HPV9).
- For females, HPV2, HPV4 or HPV9 is recommended in a 3-dose series for routine vaccination at age 11 or 12 years, and for those age 13 through 26 years, if not previously vaccinated.
- For males, HPV4 or HPV9 is recommended in a 3-dose series for routine vaccination at age 11 or 12 years and for those age 13 through 21 years, if not previously vaccinated. Males age 22 through 26 years may be vaccinated.
- HPV vaccination is recommended for men who have sex with men through age 26 years who did not get any or all doses when they were younger.
- Vaccination is recommended for immunocompromised persons (including those with HIV infection) through age 26 years for those who did not get any or all doses when they were younger.
- A complete HPV vaccination series consists of 3 doses. The second dose should be administered 4 to 8 weeks (minimum interval of 4 weeks) after the first dose; the third dose should be administered 24 weeks after the first dose and 16 weeks after the second dose (minimum interval of at least 12 weeks).
- HPV vaccines are not recommended for use in pregnant women. However, pregnancy testing is not needed before vaccination. If a woman is found to be pregnant after initiating the vaccination series, no intervention is needed; the remainder of the 3-dose series should be delayed until completion or termination of pregnancy.

6. Zoster vaccination.

- A single dose of zoster vaccine is recommended for adults age 60 years or older regardless of whether they report a prior episode of herpes zoster.
 Although the vaccine is licensed by the U.S. Food and Drug Administration for use among and can be administered to persons age 50 years or older, ACIP recommends that vaccination begin at age 60 years.
- Persons age 60 years and older with chronic medical conditions may be vaccinated unless their condition constitutes a contraindication, such as pregnancy or severe immunodeficiency.

7. Measles, mumps, rubella (MMR) vaccination.

- Adults born before 1957 are generally considered immune to measles and mumps. All adults born in 1957 or later should have documentation of 1 or more doses of MMR vaccine unless they have a medical contraindication to the vaccine or laboratory evidence of immunity to each of the three diseases. Documentation of provider-diagnosed disease is not considered acceptable evidence of immunity for measles, mumps, or rubella.
- Measles component: A routine second dose of MMR vaccine, administered
 a minimum of 28 days after the first dose, is recommended for adults who:
 1) are students in postsecondary educational institutions, 2) work in a health
 care facility, or 3) plan to travel internationally. Persons who received inactivated (killed) measles vaccine or measles vaccine of unknown type from
 1963–1967 should be revaccinated with 2 doses of MMR vaccine.
- Mumps component: A routine second dose of MMR vaccine, administered
 a minimum of 28 days after the first dose, is recommended for adults who:
 1) are students in a postsecondary educational institution, 2) work in a health
 care facility, or 3) plan to travel internationally. Persons vaccinated before

- 1979 with either killed mumps vaccine or mumps vaccine of unknown type who are at high risk for mumps infection (e.g., persons who are working in a health care facility) should be considered for revaccination with 2 doses of MMR vaccine.
- Rubella component: For women of childbearing age, regardless of birth year, rubella immunity should be determined. If there is no evidence of immunity, women who are not pregnant should be vaccinated. Pregnant women who do not have evidence of immunity should receive MMR vaccine upon completion or termination of pregnancy and before discharge from the health care facility.
- Health care personnel born before 1957: For unvaccinated health care
 personnel born before 1957 who lack laboratory evidence of measles, mumps,
 and/or rubella immunity or laboratory confirmation of disease, health care
 facilities should consider vaccinating personnel with 2 doses of MMR vaccine
 at the appropriate interval for measles and mumps or 1 dose of MMR vaccine
 for rubella.

8. Pneumococcal vaccination.

- General information
 - Adults are recommended to receive 1 dose of 13-valent pneumococcal conjugate vaccine (PCV13) and 1, 2, or 3 doses (depending on indication of 23-valent pneumococcal polysaccharide vaccine (PPSV23).
 - PCV13 should be administered at least 1 year after PPSV23.
 - PPSV23 should be administered at least 1 year after PCV13, except among adults with immunocompromising conditions, anatomical or functional asplenia, or cerebrospinal fluid leak or cochlear implant, for whom the interval is at least 8 weeks; the interval between PPSV23 is at least 5 years.
 - No additional dose of PPSV23 is indicated for adults vaccinated with PPSV23 at age 65 years or older.
 - When indicated, PCV13 and PPSV23 should be administered to adults whose pneumococcal vaccination history is incomplete or unknown.
- Adults age 65 years and older (immunocompetent) who
 - Have not received PCV13 or PPSV23: Administer PCV13 followed by PPSV23 at least 1 year after PCV13.
 - Have not received PCV13 but have received a dose of PPSV23 at age 65 years or older: Administer PCV13 at least 1 year after PPSV23.
 - Have not received PCV13 but have received 1 or more doses of PPSV23 before age 65 years: Administer PCV13 at least 1 year after the most recent dose of PPSV23. Administer a dose of PPSV23 at least 1 year after PCV13 and at least 5 years after most recent dose of PPSV23.
 - Have received PCV13 but not PPSV23 before age 65 years: Administer PPSV23 at least 1 year after PCV13.
 - Have received PCV13 and 1 or more doses of PPSV23 before age 65 years: Administer PPSV23 at least 1 year after PCV13 and at least 5 years after the most recent dose of PPSV23.
- Adults age 19 years or older with immunocompromising conditions or anatomical or functional asplenia (defined below) who
 - Have not received PCV13 or PPSV23: Administer PCV13 followed by PPSV23 at least 8 weeks after PCV13. Administer a second dose of PPSV23 at least 5 years after the first dose of PPSV23.
 - Have not received PCV13 but have received 1 dose of PPSV23: Administer PCV13 at least 1 year after the PPSV23. Administer a second dose of PPSV23 at least 8 weeks after PCV13 and at least 5 years after the first dose of PPSV23.
 - Have not received PCV13 but have received 2 doses of PPSV23: Administer PCV13 at least 1 year after the most recent dose of PPSV23.
 - Have received PCV13 but not PPSV23: Administer PPSV23 at least 8

- weeks after PCV13. Administer a second dose of PPSV23 at least 5 years after the first dose of PPSV23.
- Have received PCV13 and 1 dose of PPSV23: Administer a second dose of PPSV23 at least 8 weeks after PCV13 and 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 a dose of PPSV23 at least 8 weeks after PCV13 and at least 5 years after the last dose of PPSV23.

Immunocompromising conditions that are indications for pneumococcal vaccination are: Congenital or acquired immunodeficiency (including B- or T-lymphocyte deficiency, complement deficiencies, and phagocytic disorders excluding chronic granulomatous disease), HIV infection, chronic renal failure, nephrotic syndrome, leukemia, lymphoma, Hodgkin disease, generalized malignancy, multiple myeloma, solid organ transplant, and iatrogenic immunosuppression (including long-term systemic corticosteroids and radiation therapy).

Anatomical or functional asplenia that are indications for pneumococcal vaccination are: Sickle cell disease and other hemoglobinopathies, congenital or acquired asplenia, splenic dysfunction, and splenectomy. Administer pneumococcal vaccines at least 2 weeks before immunosuppressive therapy or an elective splenectomy, and as soon as possible to adults who are newly diagnosed with asymptomatic or symptomatic HIV infection

- Adults age 19 years and older with cerebrospinal fluid leaks or cochlear implants: administer PCV13 followed by PPSV23 at least 8 weeks after PCV13; no additional dose of PPSV23 is indicated if age is younger than 65 years. If 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.
- Adults age 19 through 64 years who have chronic heart disease (including congestive heart failure and cardiomyopathies, excluding hypertension), chronic lung disease (including chronic obstructive lung disease, emphysema, and asthma), chronic liver disease (including cirrhosis), alcoholism, or diabetes mellitus, or who smoke cigarettes: administer PPSV23. At age 65 years or older, administer PCV13 at least 1 years after PPSV23, followed by another dose of PPSV23 at least 1 year after PCV13 and at least 5 years after the last dose of PPSV23.
- Routine pneumococcal vaccination is not recommended for American Indian/ Alaska Native or other adults unless they have the indications as above; however, public health authorities may consider recommending the use of pneumococcal vaccines for American Indians/Alaska Natives or other adults who live in areas with increased risk for invasive pneumococcal disease.

9. Hepatitis A vaccination.

- Vaccinate any person seeking protection from hepatitis A virus (HAV) infection and persons with any of the following indications:
 - men who have sex with men
 - persons who use injection or noninjection illicit drugs;
 - persons working with HAV-infected primates or with HAV in a research laboratory setting;
 - persons with chronic liver disease and persons who receive clotting factor concentrates;
 - persons traveling to or working in countries that have high or intermediate endemicity of hepatitis A (see footnote 1); and
 - unvaccinated persons who anticipate close personal contact (e.g., house-hold or regular babysitting) with an international adoptee during the first 60 days after arrival in the United States from a country with high or intermediate endemicity of hepatitis A (see footnote 1). The first dose of the 2-dose hepatitis A vaccine series should be administered as soon as adoption is planned, ideally 2 or more weeks before the arrival of the adoptee.
- Single-antigen vaccine formulations should be administered in a 2-dose schedule at either 0 and 6 to 12 months (Havrix), or 0 and 6–18 months (Vaqta). If the combined hepatitis A and hepatitis B vaccine (Twinrix) is used,

 administer 3 doses at 0, 1, and 6 months; alternatively, a 4-dose schedule may be used, administered on days 0, 7, and 21–30, followed by a booster dose at 12 months.

10. Hepatitis B vaccination.

- Vaccinate any person seeking protection from hepatitis B virus (HBV) infection and persons with any of the following indications:
 - sexually active persons who are not in a long-term, mutually monogamous relationship (e.g., persons with more than 1 sex partner during the previous 6 months); persons seeking evaluation or treatment for a sexually transmitted disease (STD); current or recent injection drug users; and men who have sex with men;
 - health care personnel and public safety workers who are potentially exposed to blood or other infectious body fluids;
 - persons who are younger than age 60 years with diabetes as soon as feasible after diagnosis; persons with diabetes who are age 60 years or older at the discretion of the treating clinician based on the likelihood of acquiring HBV infection, including the risk posed by an increased need for assisted blood glucose monitoring in long-term care facilities, the likelihood of experiencing chronic sequelae if infected with HBV, and the likelihood of immune response to vaccination;
 - persons with end-stage renal disease (including patients receiving hemodialysis), persons with HIV infection, and persons with chronic liver disease;
 - household contacts and sex partners of hepatitis B surface antigenpositive persons, clients and staff members of institutions for persons with developmental disabilities, and international travelers to regions with high or intermediate levels of endemic HBV infection (see footnote 1); and
 - all adults in the following settings: STD treatment facilities, HIV testing
 and treatment facilities, facilities providing drug abuse treatment and
 prevention services, health care settings targeting services to injection
 drug users or men who have sex with men, correctional facilities, endstage renal disease programs and facilities for chronic hemodialysis
 patients, and institutions and nonresidential day care facilities for persons
 with developmental disabilities.
- Administer missing doses to complete a 3-dose series of hepatitis B vaccine to those persons not vaccinated or not completely vaccinated. The second dose should be administered at least 1 month after the first dose; the third dose should be administered at least 2 months after the second dose (and at least 4 months after the first dose). If the combined hepatitis A and hepatitis B vaccine (Twinrix) is used, give 3 doses at 0, 1, and 6 months; alternatively, a 4-dose Twinrix schedule may be used, administered on days 0, 7, and 21–30, followed by a booster dose at 12 months.
- Adult patients receiving hemodialysis or with other immunocompromising conditions should receive 1 dose of 40 mcg/mL (Recombivax HB) administered on a 3-dose schedule at 0, 1, and 6 months or 2 doses of 20 mcg/mL (Engerix-B) administered simultaneously on a 4-dose schedule at 0, 1, 2, and 6 months.

11. Meningococcal vaccination.

- General information
 - Serogroup A, C,W, and Y meningococcal vaccine is available as a conjugate (MenACWY [Menactra, Menveo]) or a polysaccharide (MPSV4 [Menomune]) vaccine.
 - Serogroup B meningococcal (MenB) vaccine is available as a 2-dose series of MenB-4C vaccine (Bexsero) administered at least 1 month apart or a 3-dose series of MenB-FHbp (Trumenba) vaccine administered at 0, 2, and 6 months; the two MenB vaccines are not interchangeable (i.e., the same MenB vaccine product must be used for all doses).
 - MenACWY vaccine is preferred for adults with serogroup A, C, W, and Y meningococcal vaccine indications who are age 55 years or younger and for adults age 56 years or older 1) who were vaccinated previously with MenACWY vaccine and are recommended for revaccination or 2) for whom multiple doses of vaccine are anticipated; MPSV4 vaccine is

- preferred for adults age 56 years or older who have not received Men-ACWY vaccine previously and who require a single dose only (e.g., persons at risk because of an outbreak).
- Revaccination with MenACWY vaccine every 5 years is recommended for adults previously vaccinated with MenACWY or MPSV4 vaccine who remain at increased risk for infection (e.g., adults with anatomical or functional asplenia or persistent complement component deficiencies, or microbiologists who are routinely exposed to isolates of Neisseria meningitidis.)
- MenB vaccine is approved for use in persons ages 10 through 25 years; however, because there is no theoretical difference in safety for persons older than age 25 years compared to those ages 10 through 25 years, MenB vaccine is recommended for routine use in persons ages 10 years and older who are at increased risk for serogroup B meningococcal disease.
- There is no recommendation for MenB revaccination at this time.
- MenB vaccine may be administered concomitantly with MenACWY vaccine, but at a different anatomic site, if feasible.
- HIV infection is not an indication for routine vaccination with MenACWY or MenB vaccine; if an HIV-infected person of any age is to be vaccinated, administer 2 doses of MenACWY vaccine at least 2 months apart.
- Adults with anatomical or functional asplenia or persistent complement component deficiencies: Administer 2 doses of MenACWY vaccine at least 2 months apart and revaccinate every 5 years. Also, administer a series of MenB vaccine.
- Microbiologists who are routinely exposed to isolates of Neisseria meningitidis: Administer a single dose of MenACWY vaccine; revaccinate with MenACWY vaccine every 5 years if the person remains at risk for infection. In addition, administer a series of Men B vaccine.
- Persons at risk because of a meningococcal disease outbreak: If the outbreak
 is attributable to serogroup A, C, W, or Y, administer a single dose of MenACWY vaccine; if the outbreak is attributable to serogroup B, administer a
 series of MenB vaccine.
- Persons who travel to or live in countries in which meningococcal disease is hyperendemic or epidemic: Administer a single dose of MenACWY vaccine and revaccinate with MenACWY vaccine every 5 years if an increased risk of infection remains (see footnote 1); MenB vaccine is not recommended because meningococcal disease in these countries is generally not caused by serogroup B.
- Military recruits: Administer a single dose of MenACWY vaccine.
- First-year college students ages 21 years or younger who live in residence halls: Administer a single dose of MenACWY vaccine if they have not received a dose on or after their 16th birthday.
- Young adults ages 16 through 23 years (preferred age range is 16 through 18 years): May be vaccinated with a series of MenB vaccine to provide shortterm protection against most strains of serogroup B meningococcal disease.

12. Haemophilus influenzae type b (Hib) vaccination.

- One dose of Hib vaccine should be administered to persons who have anatomical or functional asplenia or sickle cell disease or are undergoing elective splenectomy if they have not previously received Hib vaccine. Hib vaccination 14 or more days before splenectomy is suggested.
- Recipients of hematopoietic stem cell transplant (HSCT) should be vaccinated with a 3-dose regimen 6–12 months after a successful transplant, regardless of vaccination history; at least 4 weeks should separate doses.
- Hib vaccine is not recommended for adults with HIV infection since their risk for Hib infection is low.

13. Immunocompromising conditions.

Inactivated vaccines (e.g., pneumococcal, meningococcal, and inactivated influenza vaccines) generally are acceptable and live vaccines generally should be avoided in persons with immune deficiencies or immunocompromising conditions. Information on specific conditions is available at www.cdc. gov/vaccines/hcp/acip-recs/index.html.

Guide to Contraindications and Precautions to Commonly Used Vaccines in Adults^{1,*,†}

Vaccine	Contraindications ¹	Precautions ¹
Influenza, inactivated (IIV)² Influenza, recombinant (RIV)²	For IIV, severe allergic reaction (e.g., anaphylaxis) after a previous dose of any influenza vaccine; or to a vaccine component, including egg protein For RIV, severe allergic reaction (e.g., anaphylaxis) after a previous dose of RIV or to a vaccine component. RIV does not contain any egg protein²	Moderate or severe acute illness with or without fever History of Guillain-Barré Syndrome (GBS) within 6 weeks of previous influenza vaccination Adults with egg allergy of any severity may receive RIV; adults with hivesonly allergy to eggs may receive IIV with additional safety measures²
Influenza, live attenuated (LAIV) ^{2,3}	Severe allergic reaction (e.g., anaphylaxis) to any component of the vaccine, or to a previous dose of any influenza vaccine In addition, ACIP recommends that LAIV not be used in the following populations: pregnant women; immunosuppressed adults; adults with egg allergy of any severity; adults who have taken influenza antiviral medications (amantadine, rimantadine, zanamivir, or oseltamivir) within the previous 48 hours; avoid use of these antiviral drugs for 14 days after vaccination	Moderate or severe acute illness with or without fever History of GBS within 6 weeks of previous influenza vaccination Asthma in persons age 5 years and older Other chronic medical conditions (e.g., other chronic lung diseases, chronic cardiovascular disease [excluding isolated hypertension], diabetes, chronic renal or hepatic disease, hematologic disease, neurologic disease, and metabolic disorders)
Tetanus, diphtheria, pertussis (Tdap) Tetanus, diphtheria (Td)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component 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 Tdap, diphtheria and tetanus toxoids and pertussis (DTP), or diphtheria and tetanus toxoids and acellular pertussis (DTaP) vaccine	Moderate or severe acute illness with or without fever GBS within 6 weeks after a previous dose of tetanus toxoid-containing vaccine 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 For pertussis-containing vaccines: progressive or unstable neurologic disorder, uncontrolled seizures, or progressive encephalopathy until a treatment regimen has been established and the condition has stabilized
Varicella (Var) ³	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component Known severe immunodeficiency (e.g., from hematologic and solid tumors, receipt of chemotherapy, congenital immunodeficiency, or long-term immunosuppressive therapy ⁴ or patients with human immunodeficiency virus [HIV] infection who are severely immunocompromised) Pregnancy	Noderate or severe acute illness with or without fever Recent (within 11 months) receipt of antibody-containing blood product (specific interval depends on product) ⁵ Receipt of specific antivirals (i.e., acyclovir, famciclovir, or valacyclovir) 24 hours before vaccination; avoid use of these antiviral drugs for 14 days after vaccination
Human papillomavirus (HPV)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component	Moderate or severe acute illness with or without fever Pregnancy
Zoster (HZV) ³	Severe allergic reaction (e.g., anaphylaxis) to a vaccine component Known severe immunodeficiency (e.g., from hematologic and solid tumors, receipt of chemotherapy, or long-term immunosuppressive therapy, or patients with HIV infection who are severely immunocompromised) Pregnancy	Moderate or severe acute illness with or without fever Receipt of specific antivirals (i.e., acyclovir, famciclovir, or valacyclovir) 4 hours before vaccination; avoid use of these antiviral drugs for 14 days after vaccination
Measles, mumps, rubella (MMR) ³	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component Known severe immunodeficiency (e.g., from hematologic and solid tumors, receipt of chemotherapy, congenital immunodeficiency, or long-term immunosuppressive therapy, or patients with HIV infection who are severely immunocompromised) Pregnancy	Moderate or severe acute illness with or without fever Recent (within 11 months) receipt of antibody-containing blood product (specific interval depends on product) ⁵ History of thrombocytopenia or thrombocytopenic purpura Need for tuberculin skin testing ⁶
Pneumococcal: conjugate (PCV13), polysaccharide (PPSV23)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component (including, for PCV13, to any vaccine containing diphtheria toxoid-containing vaccine	Moderate or severe acute illness with or without fever
Hepatitis A (HepA) Hepatitis B (HepB)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component	Moderate or severe acute illness with or without fever
Meningococcal: conjugate (MenACWY), serogroup B (MenB), polysaccharide (MPSV4)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component	Moderate or severe acute illness with or without fever
Haemophilus influen- zae type b (Hib)	Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component	Moderate or severe acute illness with or without fever

FOOTNOTES

- 1. Vaccine package inserts and the full ACIP recommendations for these vaccines should be consulted for additional information on vaccine-related contraindications and precautions and for more information on vaccine $excipients. Events or conditions \ listed \ as \ precautions \ should \ b \dot{e} \ reviewed \ carefully. \ Benefits \ of \ and \ risks \ for \ administration \ administration \ and \ risks \ for \ administration \ administratio$ istering a specific vaccine to a person under these circumstances should be considered. If the risk from the vaccine is believed to outweigh the benefit, the vaccine should not be administered. If the benefit of vaccination is believed to outweigh the risk, the vaccine should be administered. A contraindication is a condition in a recipient that increases the chance of a serious adverse reaction. Therefore, a vaccine should not be administered when a contraindication is present.
- 2. For more information on use of influenza vaccines among persons with egg allergies and a complete list of conditions that CDC considers to be reasons to avoid receiving LAIV, see CDC. "Prevention and Control of Seasonal Influenza with Vaccines: Recommendations of the Advisory Committee on Immunization Practices (ACIP)— United States, 2015-16 Influenza Season. MMWR 2015;64(30):818-25.
- 3. LAIV, MMR, varicella, or zoster vaccines can be administered on the same day. If not administered on the same day, live vaccines should be separated by at least 28 days.
- 4. Immunosuppressive steroid dose is considered to be 2 or more weeks of daily receipt of 20 mg prednisone or the equivalent. Vaccination should be deferred for at least 1 month after discontinuation of such therapy. Providers should consult ACIP recommendations for complete information on the use of specific live vaccines among

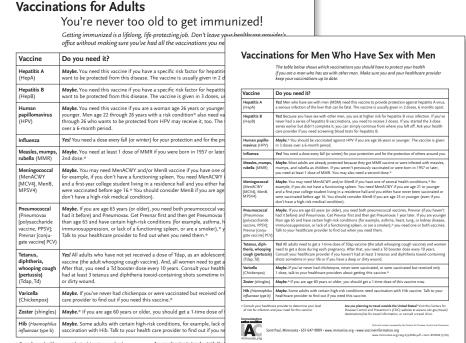
- persons on immune-suppressing medications or with immune suppression because of other reasons.
- Vaccine should be deferred for the appropriate interval if replacement immune globulin products are being administered (see Table 5 in CDC. "General Recommendations on Immunization: Recommendations of the Advisory Committee on Immunization Practices [ACIP]." MMWR 2011;60 (No. RR-2), available at www.cdc.gov/ vaccines/hcp/acip-recs/index.html.
- 6. Measles vaccination might suppress tuberculin reactivity temporarily. Measles-containing vaccine may be administered on the same day as tuberculin skin testing. If testing cannot be performed until after the day of MMR vaccination, the test should be postponed for at least 4 weeks after the vaccination. If an urgent need exists to skin test, do so with the understanding that reactivity might be reduced by the vaccine.
- * Adapted from "Table 6. Contraindications and Precautions to Commonly Used Vaccines" found in: CDC. "General Recommendations on Immunization: Recommendations of the Advisory Committee on Immunization Practices (ACIP)." MMWR 2011;60(No. RR-2), p. 40-41, and from Hamborsky J, Kroger A, Wolfe C, 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.
- Regarding latex allergy, consult the package insert for any vaccine administered.

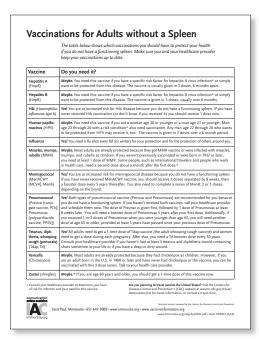
Technical content reviewed by the Centers for Disease Control and Prevention

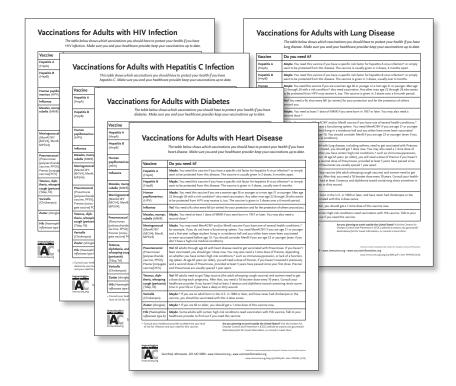
Patient Schedules for All Adults and for High-Risk Adults

These documents reflect current ACIP recommendations.

Download, make copies, and hand them to your patients.







Vaccinations for Adults – You're Never Too Old to Get Immunized www.immunize.org/catg.d/p4030.pdf

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Vaccinations for Adults without a Spleen www.immunize.org/catg.d/p4047.pdf

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Vaccinations for Adults with Heart Disease www.immunize.org/catg.d/p4044.pdf

Vaccinations for Adults with Lung Disease www.immunize.org/catg.d/p4045.pdf

IAC's Popular Summary of ACIP Recommendations for Adult Immunization

Make copies on cardstock and keep as handy references in exam rooms!

For more than 20 years, IAC has been publishing its summary of ACIP adult vaccine recommendations, updating it yearly or more often, and sending it to CDC for technical review. The adult summary has been reprinted in state immunization newsletters, textbooks, and is one of the most frequently downloaded documents from IAC's website. If you haven't seen it or used it, please try it out!

Summary of Recommendations for Adult Immunization (Age 19 years and older)

(Page 1 of 5)

Vaccine name and route	People for whom vaccination is recommended	Schedule for vaccination administration (any vaccine can be given with another unless otherwise noted)	Contraindications and precautions (mild illness is not a contraindication)
Influenza Inactivated Influenza vaccine (IIV*) Give IM or ID (intradermally) * includes recombinant influenza vaccine (RIV3) Live attenuated influenza vaccine (LAIV) Give NAS (intranasally)	For people through age 18 years, consult "Summary of Recommendations for Child/Teen Immunization" at www.immunize. org/catg.d/p2010.pdf. • Vaccination is recommended for all adults. • LAIV (Flumist) is approved only for healthy nonpregnant people age 2–49 yrs. • Adults age 18 through 64 yrs may be given any intramuscular IIV product (Fluzone, Fluvirin, Afluria, Flucelvax), or the intradermal IIV product (Fluzone Intradermal), or RIV3 (FluBlok). • Adults age 18 through 64 yrs may be given intramuscular IIV (Afluria) with a needle and syringe or using a jet injector (Stratis). • Adults age 65 yrs and older may be given standard-dose IIV, or high-dose IIV (Fluzone High-Dose), or RIV3. NOTE: Health care personnel who care for severely immunocompromised persons (i.e., those who require care in a protective environment) should receive IIV rather than LAIV. For information on other contraindications and precautions to LAIV, see far right column.	Give 1 dose every year in the fall or winter. Begin vaccination services as soon as vaccine is available and continue until the supply is depleted. Continue to give vaccine to unvaccinated adults throughout the influenza season (including when influenza activity is present in the community) and at other times when the risk of influenza exists. If 2 or more of the following live virus vaccines are to be given – LAIV, MMR, Var, HZV, and/or yellow fever – they should be given on the same day. If they are not given on the same day, space them by at least 28d.	Contraindications Previous severe allergic reaction (e.g., anaphylaxis) to this vaccine, to any of its components, including egg protein. Adults with egg allergy of any severity may receive RIV or, adults who experience only hives with exposure to eggs may receive other IIV with additional safety precautions (i.e., observe patient for 30 minutes after receipt of vaccine for signs of a reaction). For LAIV only: pregnancy; immunosuppression; receipt of specific antivirals (i.e., amantadine, rimantadine, zanamivir, or oseltamivir) within the previous 48hrs. Avoid use of these anti-viral drugs for 14d after vaccination. Precautions Moderate or severe acute illness. History of Guillain-Barré syndrome (CBS) within 6 wks following previous influenza vaccination. For LAIV only: Chronic pulmonary (including asthma), cardiovascular (except hypertension), renal, hepatic, neurologic, hematologic, or metabolic (including diabetes) disorders; immunosuppression (including that caused by medications or HIV).
Td, Tdap (Tetanus, diphtheria, pertussis) Give IM	For people through age 18 years, consult "Summary of Recommendations for Child/Teen Immunization" at www.immunize.org/catg.d/ p2010.pdf. • All people who lack written documentation of a primary series consisting of at least 3 doses of tetanus- and diphtheriatoxoid-containing vaccine. • A booster dose of Td or Tdap may be needed for wound management, so consult ACIP recommendations.¹ For Tdap only • Adults who have not already received Tdap or whose Tdap history is not known. • Health care personnel of all ages. • Give Tdap to pregnant women during each pregnancy (preferred during 27–36 weeks' gestation), regardless of the interval since prior Td or Tdap.	For people who are unvaccinated or behind, complete the primary Td series (3 doses with an interval of 1–2m between dose #1 and #2, and an interval of 6–12m between dose #2 and #3); substitute a one-time dose of Tdap for one of the doses in the series, preferably the first. Give Td booster every 10 yrs after the primary series has been completed. Tdap should be given regardless of interval since previous Td.	Contraindications Previous severe allergic reaction (e.g., anaphylaxis) to this vaccine or to any of its components. For Tdap only, history of encephalopathy not attributable to an identifiable cause, within 7d following DTP/DTaP, or Tdap. Precautions Moderate or severe acute illness. History of Guillain-Barré syndrome within 6wks following previous dose of tetanus-toxoid-containing vaccine. History of arthus reaction following a prior dose of tetanusor diphtheria-toxoid-containing vaccine (including MCV4); defervaccination until at least 10 yrs have elapsed since the last tetanus toxoid-containing vaccine. For pertussis-containing vaccines only, progressive or unstable neurologic disorder, uncontrolled seizures, or progressive encephalopathy until a treatment regimen has been established and the condition has stabilized.

1 CDC. Preventing Tetanus, Diphtheria, and Pertussis Among Adults: Use of Tetanus Toxoid, Reduced Diphtheria Toxoid and Acellular Pertussis Vaccine. Recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR

This document was adapted from the recommendations of the Advisory Committee on Immunization Practices (ACIP). To obtain copies of these recommendations, visit CDC's website at www.cdc.gov/vaccines/hcp/ACIP-recs/index.html or visit the Immunization Action Coalition (IAC) website at www.immunize.org/acip.

This table is revised periodically. Visit IAC's website at www.immunize.org/adultrules to make sure you have the most current version.

For the purposes of calculating intervals between doses, 4 weeks = 28 days. Intervals of 4 months or greater are determined by calendar months.

A vaccine series does not need to be restarted, regardless of the time that has elapsed between doses.

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www.immunize.org/catg.d/p2011.pdf · Item #P2011 (9/15)

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> HEALTHCARE PROFESSIONALS: Be sure to recommend zoster vaccine for your patients age 60 and older.

> > 'n severe immuno-

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14 days after

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- Contraindications and Precautions
- Do not give zoster vaccine to a person who has had an anaphylactic onent of the vaccine.

Herpes Zoster (Shingles) Vaccine Pocket Guide

Who Should Be Vaccinated

A one-time vaccination against herpes zoster (also known as shingles) is routinely recommended for all people age 60 yrs and older who do

ints with Patients

not have a valid contraindication to the vaccine.

Pneumococcal Vaccine Pocket Guide

Routine Vaccination with PCV13 and PPSV

Children: Administer pneumococcal conjugate vaccine (PCV13) to all infants and children at ages 2, 4, and 6 mos with a booster at age 12-15 mos. For incomplete or unvaccinated children, catch-up vaccination should occur through age 59 mos.

Adults age 65 years (or older):

- Administer a 1-time dose of PCV13 (if not previously received).
- Administer a dose of pneumococcal polysaccharide vaccine (PPSV) at least 1 yr after PCV13.

Risk-Based Vaccination with PCV13 and PPSV

A dose of PPSV is recommended for all people age 2 through 64 yrs with any of the following conditions:

- a. Cigarette smokers age 19 yrs and older
- b. Chronic cardiovascular disease (e.g., congestive heart failure, cardiomyopathy)
- c. Chronic pulmonary disease (including asthma in people age 19 yrs and older)
- d. Diabetes mellitus, alcoholism, or chronic liver disease
- e. Candidate for or recipient of cochlear implant
- f. Cerebrospinal fluid leak
- g. Functional or anatomic asplenia (e.g., sickle cell disease, splenectomy) h. Immunocompromising conditions (e.g., congenital or acquired
- immunodeficiency, HIV infection, leukemia, lymphoma, Hodgkin disease, generalized malignancy, iatrogenic immunosuppression, multiple myeloma, or on immunosuppressive therapy, including long-term systemic corticosteroids, radiation therapy)
- i. Solid organ transplantation; for bone marrow transplantation patients, see www.cdc.gov/vaccines/pubs/hemato-cell-transplts.htm j. Chronic renal failure or nephrotic syndrome
- A second dose of PPSV is recommended for children and adults through age 64 yrs who are at highest risk of serious pneumococcal disease or likely to have a rapid decline in pneumococcal antibody levels (categories

g-j above) at least 5 yrs after dose #1. Note: Administer an additional dose of PPSV to all adults at age 65 yrs (or older). Give it at least 5 yrs after any previous PPSV.

A 1-time dose of PCV13 is recommended for previously unvaccinated people age 6 through 64 yrs who meet any of the criteria in categories

Created by the Immunization Action Coalition (www.immunize.org) Supported by an educational grant from Merck.

inistration

SV intramuscularly (22-25g; needle age/body mass [1–1½"]); PPSV may ously (23–25g, 5/s" needle).

Use of PCV13 and PPSV

CV13 (if not previously received) after PCV13. For those with conditions r 8 wks later

eviously received PCV13 and who ait at least 8 wks after the dose of

eed PPSV dose #2 should receive it

age 64 yrs who need PCV13 and PPSV, 5V at least 8 wks later. If previously ren through age 18 yrs PCV13 at least of PPSV; give adults age 19 through

ov/mmwr/preview/mmwrhtml/

ecautions

tients with a history of an anaphylactic CV13, PPSV, or one of their components. the same visit; see "Intervals for SV" section above.

PSV (in all) and PCV13 (in adults) are site for 1–2 days. For PCV13 in young s are decreased appetite and irritability.

monly presents as a serious infection (bacteremia), or brain (meningitis). r invasive pneumococcal disease (bac-early 35,000 cases and 4,200 deaths. n occurs in older people, as well as in lition (e.g., immunosuppression,

n't remember receiving pneumococcal

risk for developing shingles if they have had an earlier chickenpox. Studies have shown that over 99% of adults 60 yrs and older have been previously infected with hether or not they recall the episode.

people will develop shingles at some time in their lives vaccinated against it. The older you are, the greater leveloping shingles.

Illy starts as a blistering rash which can become severe. be extremely painful and can cause serious problems, ness if it affects the eye. Other serious problems that may ingles include skin infections, muscle weakness, loss of hearing.

experience long-term nerve pain that can last for months. d can interfere with eating and sleeping, and be

Points for Healthcare Providers

used by the reactivation of the dormant chickenpox virus ells of people who have been infected with chickenpox. oster vaccine efficacy decreases with age, getting vacciintly reduces the risk of developing post-herpetic neuralgia ps. The oldest adults are at the highest risk of postalgia, and they are least able to tolerate the condition or ns used to control pain

vaccine is routinely recommended for people age 60 nould be covered by most health insurance plans. For in Medicare, it is currently available under Part D, the rug benefit.

r patients to purchase the vaccine from their pharmacy it to your office for administration. This "brown ht significantly jeopardize the stability of the vaccine.

nization Action Coalition (www.immunize.org) cational grant from Merck.

NOTE: These technical guides are for staff use only and are not suitable for use by patients.

 Both pneumococcal vaccines are covered under Medicare Part B for people 65 yrs and older when given at least 1 yr apart.

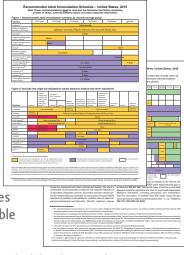
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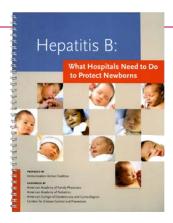
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AAFP, AAP, ACOG, and CDC endorse administering hepatitis B vaccine at birth prior to hospital discharge, and all four provided a review of this guide.



The Vaccine Handbook: A Practical Guide for Clinicians

This fifth edition is an extensive update for 2015.

The Vaccine Handbook: A Practical Guide for Clinicians ("The Purple Book") is a uniquely comprehensive source of practical, up-to-date information for vaccine providers and educators. Its author, Gary S. Marshall, MD, has drawn together the latest vaccine science and guidance into a concise, user-friendly, practical resource for the private office, public health clinic, academic medical center, and hospital.

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The fifth edition contains a foreword by Deborah L. Wexler, MD, executive director, Immunization Action Coalition, which has partnered with the publisher, Professional Communications, Inc. (PCI), to promote *The Vaccine Handbook*.



FROM THE FOREWORD:

The Purple Book belongs in the hands of every medical student, physician-in-training, doctor, nursing student, and nurse who provides vaccines to patients, regardless of patient age or medical specialty. It is my honor to introduce the Fifth Edition to you. This essential reference beautifully supports all of us in our efforts to move forward in protecting our patients from the consequences of preventable diseases.

DEBORAH L. WEXLER, MD Executive Director Immunization Action Coalition

About the Author



Gary Marshall, MD, is professor of pediatrics at the University of Louisville School of Medicine in Kentucky, where he serves as chief of the division of

pediatric infectious diseases and director of the Pediatric Clinical Trials Unit. In addition to being a busy clinician, he is nationally known for his work in the areas of vaccine research, advocacy, and education.

The Reviews

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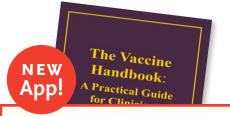
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