

EPIC[®] Immunization Update For Healthcare Providers in Training 2023 Update School Health Children, Adolescents, & Adults

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EPIC[®] is presented by:

Georgia Chapter - American Academy of Pediatrics
Ga. Dept. of Public Health/Immunization Program *In Cooperation with:*Georgia Academy of Family Physicians
Georgia Chapter - American College of Physicians
Georgia OB/Gyn Society

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Faculty Disclosure Information

- In accordance with ACCME* and ANCC-COA* Standards, all faculty members are required to disclose to the program audience any real or apparent conflict of interest to the content of their presentation.
- This presentation will include the most current ACIP recommendations for frequently used vaccines but is not a comprehensive review of all available vaccines.
- Some ACIP recommendations for the use of vaccines have not currently been approved by the FDA.
- Detailed information regarding all ACIP Recommendations is available at www.cdc.gov/vaccines/acip/recs/index.html

*American Nurses Credentialing Center Commission on Accreditation

Objectives

At the end of this presentation, you will be able to:

- Recall the role vaccines have played in preventing diseases
- Discuss the importance of vaccines for children, adolescents and adults
- Summarize the most recent CDC recommendations for storage and handling of vaccines
- List at least 2 reliable sources for immunization information

RESULTS OF EXPOSURE TO A VACCINE PREVENTABLE DISEASE



GOALS OF VACCINATING



Community Immunity Formerly known as "Herd Immunity"*



*Presentation from Immunize Georgia, September 9, 2016 by Walt A. Orenstein, MD, Professor of Medicine Global, Health, Epidemiology and Pediatrics Emory Department of Medicine, Associate Director, Emory Vaccine Center Director, Vaccine Policy and Development, Emory University, Atlanta, GA

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Vaccination Terminology (1)

Active Immunity

- Protection produced by the person's own immune system
- Usually permanent from disease
- May require multiple doses of a vaccine

Passive Immunity

- Protection transferred from another person or animal (e.g from mom to baby or Immunoglobulin treatments)
- Temporary protection that wanes with time

Epidemiology and Prevention of Vaccine-Preventable Diseases. 14th Edition, 2021, https://www.cdc.gov/vaccines/pubs/pinkbook/index.html

Vaccination Terminology (2)*

<u>Antigen</u>

• A live or inactivated substance (e.g., protein, polysaccharide) capable of producing an immune response

<u>Antibody</u>

 Protein molecules (immunoglobulin) produced by B lymphocytes to help eliminate an antigen

Vaccines*

<u>Vaccine</u> - A product that interacts with the immune system to produce <u>active</u> immunity against a disease without the risk of the disease and its potential complications.

Live, Attenuated

- Measles, Mumps & Rubella (MMR)
- Varicella
- LAIV
- Rotavirus
- Yellow fever

Inactivated

- Toxoids (DTaP, Td, Tdap)
- Whole (Hepatitis A, IPV)
- Split (Influenza IIV)
- Recombinant vaccines (Hepatitis B, 9vHPV, FluBlok, Shingrix, Men B)
- Polysaccharide vaccines -- (PPSV23)
- Conjugated vaccines (Hib, PCV13, MCV4)
- COVID-19

Vaccines Work!

CDC statistics demonstrate dramatic declines in vaccine-preventable diseases when compared with the pre-vaccine era

DISEASE	PRE-VACCINE ERA ESTIMATED ANNUAL MORBIDITY ¹	MOST RECENT REPORTS OR ESTIMATES OF U.S. CASES	PERCENT DECREASE
Diphtheria	21,053	2 ²	>99%
H. influenzae (invasive, <5 years of age)	20,000	14 ^{2,3}	>99%
Hepatitis A	117,333	(est) 24,900 ⁴	79%
Hepatitis B (acute)	66,232	(est) 21,600 ⁴	67%
Measles	530,217	1,287 ²	>99%
Meningococcal disease (all serotypes)	2,8865	329 ²	89%
Mumps	162,344	3,509²	98%
Pertussis	200,752	15,662 ²	92%
Pneumococcal disease (invasive, <5 years of age)	16,069	1,7007	93%
Polio (paralytic)	16,316	0 ²	100%
Rotavirus (hospitalizations, <3 years of age)	62,500 ⁸	30,625 ⁹	51%
Rubella	47,745	4 ²	>99%
Congenital Rubella Syndrome	152	0²	100%
Smallpox	29,005	0 ²	100%
Tetanus	580	19²	96%
Varicella	4,085,120	102,128 ¹⁰	>98%

https://www.immunize.org/catg. d/p4037.pdf

Advisory Committee on Immunization Practices (ACIP)

- 15 voting members with expertise in one or more of the following:
 - Vaccinology
 - Immunology
 - Infectious diseases
 - Pediatrics
 - Internal Medicine
 - Preventive medicine
 - Public health
 - Consumer perspectives and/or social and community aspects
 of immunization programs
- ACIP develops recommendations and schedules for the use of licensed vaccines



Indications Recommendations Requirements



Indication

Information about the appropriate use of the vaccine

Recommendation

- ACIP statement that broadens and further delineates the Indication found in the package insert
- Basis for standards for best practice
- All ACIP Recommendations can be found at: https://www.cdc.gov/vaccines/hcp/acip-recs/index.html

Requirement

 Mandate by a state that a particular vaccine must be administered and documented before entrance to childcare and/or school 5/6/2023 13

2023 Childhood and Adolescent **Immunization Schedules**

- Recommended Schedule for Children Ages 0-18 Years •
- Catch-up Schedule ٠
- Vaccines that might be indicated for children and adolescents • aged 18 years or younger based on medical indications
- Footnotes \bullet

READ THE FOOTNOTES TO ACCESS SPECIFIC VACCINE ADMINISTRATION DETAILS!







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2023 Recommended Immunization Schedule for Adults Aged ≥19 Years*

- Recommended adult schedule by age group
- Recommended immunization schedule for adults aged 19 years or older by medical condition and other indications
- Contraindications and Precautions
- Footnotes

READ THE FOOTNOTES TO ACCESS SPECIFIC VACCINE ADMINISTRATION DETAILS!

Table 1 Recommended Adult Immunization Schedule by Age Group, United States, 2021						
Vaccine	19–26 years	27-49 years		50-64 years	≥65 years	
Influenza inactivated (IIV) or Influenza recombinant (RIV4)	1 dose annually					
Influenza live, attenuated (LAIV4)	tor 1 dose annually					
Tetanus, diphtheria, pertussis	1 dose Tdap each pregnancy; 1 dose Td/Tdap for wound management (see notes)					
Measles, mumps, rubella (MMR)	i dove izagi, tien i dori laga booster every i lyears 1 or 2 doses depending on indication (if born in 1957 or later)					
Varicella (VAR)	2 doses (if born in 1980 or later)		2 doses			
Zoster recombinant (RZV)				2 do	oses	
Human papillomavirus (HPV)	2 or 3 doses depending on age initial vaccination or condition	at 27 through 45 years				
Pneumococcal conjugate (PCV13)	1 dose 1 dose					
Pneumococcal polysaccharide (PPSV23)	1 or 2 doses depending on indication			1 dose		
Hepatitis A (HepA)	2 or 3 doses depending on vaccine					
Hepatitis B (HepB)	2 or 3 doses depending on vaccine					
Meningococcal A, C, W, Y (MenACWY)	1 or 2 doses depending on indication, see notes for booster recommendations					
Meningococcal B (MenB)	2 or 3 doses depending on vaccine and indication, see notes for booster recommendations 19 through 23 years					
Haemophilus influenzae type b (Hib)	1 or 3 doses depending on indication					
Recommended successor for data who were appropriate the second se						



Vaccine Schedules Varying From ACIP/AAP/AAFP Recommendations

Alternate Schedules

- Dr. Bob's Selective Vaccine Schedule
- Dr. Bob's Alternative Vaccine Schedule
- Parent-derived schedules
- Parent/caretaker refusal of all vaccines

Concerns re: alternate schedules

- Alternate or delayed schedules have not been tested
- No studies to prove they are safer

If any of these Alternate Schedules are requested, the health care provider and staff must spend additional time educating the parent/caretaker about the appropriate use of vaccines.

MAKING HEADLINES

When a vaccine works, it prevents a disease. Prevention does not make headlines.

The possibility that a vaccine has an adverse effect, true or false, DOES MAKE A GOOD STORY.





Vaccine Risk Perception

Many parents of young children are not familiar with vaccine-preventable diseases and perceive the risks of vaccines outweigh the benefits

Myths and Concerns

- Myth: Immune system overload
- Children get too many shots at one visit
- Vaccines have side effects (adverse reactions)
- Myth: Immunity from the disease is better than immunity from a vaccine (ie. chicken pox)
- Myth: Vaccines cause autism

Response to Vaccine Safety Concerns

- Vaccines are among the most thoroughly tested and safest things we put into our bodies
- Refusing a vaccine means taking the risks of the disease and spreading the disease to others
- "Natural immunity" (from disease) may come with complications, permanent damage, or death
- In Georgia, an unimmunized student may be prohibited from attending school during an epidemic*
- Consistent reproducible research has shown that autism is NOT caused by:
 - Thimerosal
 - Multiple vaccines at one time
 - MMR vaccine

Talking with Parents about Vaccines*

- Start conversations early (prenatal visits)
- Use language and examples parents can understand
- Give written information (VIS) prior to the immunization visit
- Provide your recommendations
- Draw upon your experiences as a health care provider
- Solicit and welcome questions
- Recognize that some parents may be more interested in discussing vaccines than others

Resources to encourage Childhood Vaccinations

Resources

Use these resources to promote routine childhood vaccinations in your practice.



The Catch-up Immunization Schedule

Children and teens can catch up on vaccinations even if they start late or are more than one month behind schedule. Check out this catch-up schedule and share with parents to help get their child caught up on routine vaccination.





Resources to Encourage Childhood Vaccinations



VFC Flyer for Parents



Vaccines for Children (VFC) Program



Reminders & Recall Systems



Anti-Vaccine Movement

- Promotes the idea that there is less evidence of disease today and immunizations are no longer needed
- Sends confusing & conflicting information
- Uses stories, personal statements, and books to play on the emotional side of concerned parents
- Encourage parents/patients to:
 - Get the facts
 - Consider the source
 - Discuss their concerns with you



Global Vaccine Awareness League







Resources for Factual & Responsible Vaccine Information



VACCINE PREVENTABLE DISEASES







Diphtheria

Tetanus



Vaccines Containing Diphtheria & Tetanus Toxoid plus Pertussis Antigens

ACIP recommends:

DTaP – 2 months through 6 years (Multiple doses)

Tdap

- Children and adolescents starting at 11 or 12 years of age
- Any adult who has not received a dose
- Either Tdap or Td can be used for routine booster every 10 yrs.
- Either vaccine can be used for tetanus prophylaxis for wound management

Haemophilus influenzae type b (Hib)*

ACIP recommends Hib vaccine:3 or 4 doses for children2 through 15 months of age



One dose of Hib for unimmunized persons 5 through 18 years who have asplenia, sickle cell disease, or HIV infection.

One dose of Hib may be given to adults with immunocompromising conditions.

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ACIP recommends: Inactivated Polio Vaccine (IPV) 4 doses 2 months through 6 years Persons traveling to countries experiencing polio outbreaks may require a booster dose

Measles, Mumps, Rubella Measles (M) Mumps (M)





Source: American Academy of Pediatrics Red Book On Line Visual Library

Source: Creative Commons

Rubella (R)







Congenital Rubella (R)

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MEASLES

- Incubation period---11 to 12 days from exposure to onset of symptoms
- Symptoms: fever, cough, coryza, conjunctivitis, maculopapular rash and Koplik spots
- Complications: otitis media, pneumonia, croup, diarrhea, encephalitis and death
- Subacute sclerosing panencephalitis (SSPE) is a progressive neurological disorder that is rare but always fatal. It usually occurs 7-10 years after measles infection.



Source: Immunization Action Coalition

MMR Vaccine

ACIP recommendations:

Children: 2 doses of MMR:

- Dose 1 @ 12 through 15 months of age
- Dose 2 @ 4 through 6 years of age

Second dose can be given 28 days after first dose, if necessary.

<u>Adults:</u>

- At least 1 dose MMR for unvaccinated adults
- 2 doses MMR for students entering colleges, universities, technical and vocational schools, and other post-high-school educational institutions
- 2 doses MMR for measles and mumps and 1 dose MMR for rubella for healthcare personnel
- Travelers to foreign countries should be appropriately immunized with MMR before leaving U.S.
- Infants 6-12 mos. of age traveling abroad should receive 1 dose of MMR. This dose must be repeated at age 12 -15 months of age and a second dose at least 4 weeks later.
- A 3rd MMR may be recommended in the instance of a public health-declared mumps outbreak.

MMR Vaccine

- Antibodies develop in approximately 95% of children vaccinated at age 12 months and over 99% of children who receive 2 doses
- Immunity long-term and probably lifelong in most persons
 What about adults?
- Evidence of Immunity: Generally, persons can be considered immune to measles if they were:
 - born before 1957,
 - have serologic evidence of measles immunity (equivocal test results should be considered negative),
 - laboratory confirmation of disease,
 - have documentation of adequate vaccination for measles.
- Healthcare providers and health departments should not accept verbal reports of vaccination without written documentation as presumptive evidence of immunity.

Measles Containing Vaccines

- <u>MMR-II</u>
- PRIORIX (GSK). ACIP Recommended June 2022
 - First licensed in Germany in 1997 and approved in over 100 countries
 - Contains equivalent vaccine virus strains as MMR II (Merck)
 - No significant differences found in safety or side effects when comparing Priorix to MMR-II.
 - PRIORIX and M-M-R II are fully interchangeable.
 - ACIP General Best Practices states a preference that doses of vaccine in a series come from the same manufacturer; however, vaccination should not be deferred when the manufacturer of the previously administered vaccine is unknown or when the vaccine from the same manufacturer is unavailable
 - Studies have shown that PRIORIX is safe and immunogenic when administered as a second dose after M-M-R II





Varicella* (Chickenpox)





ACIP recommends: Two doses of varicella vaccine for everyone who has not had chickenpox

Acceptable Evidence of Varicella Immunity

- Written documentation of age-appropriate vaccination
- Laboratory evidence of immunity or laboratory confirmation of varicella disease
- U.S.-born before 1980
 - Does not apply to healthcare personnel or pregnant people
- Healthcare provider diagnosis or verification of varicella disease
- History of herpes zoster based on healthcare provider diagnosis

Vaccines containing Varicella antigen

• Varicella

 ACIP recommends first dose at 12 through 15 months of age and second dose at 4 through 6 years

• MMRV

- Combination vaccine with measles, mumps, rubella, and varicella antigens
- Contains 7 times the varicella component of single antigen varicella vaccine
- Approved for use in children 12 mos. through 12 years of age
Herpes Zoster

- Herpes zoster (HZ), or shingles, occurs through reactivation of latent varicella-zoster virus
- Typically characterized by prodromal pain and an acute vesicular eruption (rash) accompanied by moderate to severe pain
- One in three persons will develop zoster during their lifetime
- Post-herpetic neuralgia (PHN) is a common consequence of zoster. PHN is defined as nerve pain persisting longer than 3 mos. after disappearance of the rash.
- Risk for zoster and PHN increases with age







Shingrix[®](RZV) from GSK*

- As of November 18, 2020, Zostavax (ZVL) is no longer available for use in the United States
- Shingrix (RZV) is the only currently licensed Zoster vaccine in the United States

<u>Efficacy</u>

- > 91% in preventing zoster in all vaccinated persons in licensed age groups
- > 88% in preventing PHN
- At least 85% vaccine effectiveness >4 years post-vaccination in persons 70 years and older

$Shingrix_{\mathbb{R}}(RZV)$ from GSK*

ACIP Recommendations

- RZV (recombinant zoster vaccine) is recommended by the ACIP for the prevention of shingles and related complications.
- RZV is recommended for **immunocompetent** adults 50 years and older who previously received ZVL and **immunocompromised adults 19 years and older**.
- Two doses of RZV are recommended, regardless of prior history of herpes zoster disease or previous receipt of zoster vaccine live vaccine (ZVL).
- RZV may be given ≥2 months after prior receipt of ZVL. People who have received ZVL should be revaccinated with a 2-dose series of RZV vaccine.
- RZV may be administered to patients who previously received varicella vaccine.
- RZV may be administered while patients are taking antiviral medications.
- RZV can be administered at the same visit as other vaccines

https://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/shingles.html And https://www.cdc.gov/mmwr/volumes/71/wr/mm7103a2.htm and https://www.cdc.gov/vaccines/schedules/hcp/imz/adult.html#note-varicella







Photo courtesy AAP

Pneumococcal Disease

- Infection with pneumococcal bacteria may cause pneumonia, bacteremia, meningitis, and otitis media resulting in thousands of hospitalizations and deaths each year in the United States
- Multi-drug resistant pneumococci are common

Pneumococcal Conjugate Vaccine (PCV13, PCV15) ACIP Recommendations

<u>Children</u>

- All children PCV13 or PCV15: 4-dose series at 2, 4, 6 months and 12-15 months
- On June 22, 2022, the ACIP recommended use of PCV15 as an option for pneumococcal conjugate vaccination of persons aged <19 years, according to currently recommended PCV13 dosing and schedules.
- April 2023: PCV20 licensed by FDA for use in children. Not yet recommended by ACIP as of 5/11/2023
- For older children and adolescents (2 years through 18 years) with underlying medical conditions, see detailed recommendations at https://www.cdc.gov/vaccines/schedules/hcp/imz/child-adolescent.html#note-pneumo

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https://www.cdc.gov/vaccines/pubs/pinkbook/pneumo.html and https://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/pneumo.html and https://www.cdc.gov/mmwr/volumes/71/wr/mm7137a3.htm

Pneumococcal Polysaccharide Vaccine (PPSV23)

ACIP Recommendations:

- For children and adolescents 2 years through 18 years at high risk for pneumococcal disease and
- Adults 19 years through 64 years old with certain chronic medical conditions or other risk factors
- For all adults 65 years and older

See Summary of recommendations of PPSV23 and timing at: https://www.cdc.gov/vaccines/vpd/pneumo/hcp/who-when-tovaccinate.html

FDA Recommended Influenza Antigens for 2022-2023 Season in the U.S.

 Egg-based influenza vaccines 	Cell culture–based inactivated (ccIIV4) and recombinant (RIV4) influenza vaccines
 an influenza A/Victoria/2570/2019	 an influenza A/Wisconsin/588/2019
(H1N1)pdm09-like virus an influenza A/Darwin/9/2021	(H1N1)pdm09-like virus an influenza A/Darwin/6/2021
(H3N2)-like virus an influenza	(H3N2)-like virus an influenza
B/Austria/1359417/2021 (Victoria	B/Austria/1359417/2021 (Victoria
lineage)- like virus, and an influenza B/Phuket/3073/2013	lineage)- like virus, an influenza B/Phuket/3073/2013
(Yamagata lineage)-like virus	(Yamagata lineage)-like virus

ACIP recommends annual influenza vaccine for all persons 6 months of age and older who do not have contraindications.

Live, Attenuated Influenza Vaccine (LAIV4)*

FluMist® MedImmune (Nasal Spray)

• Licensed for healthy persons 2 through 49 years of age

LAIV4 MAY be used in the 2022-2023 season.

Contraindications to LAIV include:

- Children 2-4 yrs. of age with a diagnosis of asthma
- Persons receiving aspirin-containing medications potential risk for Reye syndrome
- Persons who are immunocompromised, by medication or disease, have a CSF leak or cochlear implant, or asplenia
- Close contacts and caregivers of severely immunosuppressed persons
- Persons who have received influenza antiviral medications within the previous days (dependent on antiviral)
- Persons with a cranial CSF leak; people with cochlear implants
- Persons with a severe allergic reaction to any component of the vaccine or to a previous dose of any influenza vaccine (exception for allergy to egg)
- Pregnancy

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History of egg allergy and egg-based Influenza vaccines

- Persons with a history of egg allergy who have experienced <u>only</u> urticaria (hives) after exposure to egg should receive influenza vaccine.
- Any licensed, recommended influenza vaccine (i.e., any IIV4, RIV4, or LAIV4) that is otherwise appropriate for the recipient's age and health status can be used.
- Persons who report having had reactions to egg involving symptoms other than urticaria (e.g., angioedema or swelling, respiratory distress, lightheadedness, or recurrent vomiting) or who required epinephrine or another emergency medical intervention can also receive any licensed, recommended influenza vaccine (i.e., any IIV4, RIV4, or LAIV4) that is otherwise appropriate for their age and health status.
- If a vaccine other than ccIIV4 or RIV4 is used, the selected vaccine should be administered in an inpatient or outpatient medical setting, including but not necessarily limited to hospitals, clinics, health departments, and physician offices. Vaccine administration should be supervised by a health care provider who is able to recognize and manage severe allergic reactions.

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Co-administration

- Inactivated influenza vaccines(IIV4s) and RIV4 may be administered simultaneously or sequentially with other inactivated vaccines or live vaccines. Injectable vaccines that are given concomitantly should be administered at separate anatomic sites.
- LAIV4 can be administered simultaneously with other live or inactivated vaccines. However, if two live vaccines are not given simultaneously, then after administration of one live vaccine (such as LAIV4), at least 4 weeks should pass before another live vaccine is administered
- Guidance concerning administration of COVID-19 vaccines with other vaccines indicates that these vaccines may be given with other vaccines, including influenza vaccines. Providers should be aware of the potential for increased reactogenicity with coadministration and should consult the CDC guidance as more information becomes available. (This is more likely with the adjuvanted or high dose IIV4s which are recommended in persons 65 years and older.

Hepatitis A



Photo Courtesy Immunization Action Coalition

- Fecal-Oral transmission
- Food borne outbreaks
- Adults average 27 lost work days per illness
- Risk factors include child or employee in child care facility and travel
- Children often asymptomatic but can infect others

Hepatitis A Vaccine

ACIP recommends 2 doses for all children:

- 12 through 23 months of age and
- Children 2 through 18 years who have not previously received the vaccine, can receive the vaccine at subsequent visits.

ACIP recommends the vaccine for any adult who requests vaccination And for adults at high risk for acquiring hepatitis A infection, including:

- Chronic liver disease
- Those traveling or working in countries with high or intermediate rates of infection
- Men Who Have Sex with Men
- Users of Injecting and Non-Injecting Drugs
- Contact with adoptees from countries with high rates of hepatitis A if contact will be within 60 days of arrival in U.S.**

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https://www.cdc.gov/vaccines/hcp/aciprecs/vacc-specific/hepa.html

Hepatitis **B**

Transmission:

- 1. Percutaneous or mucosal exposure to blood or body fluids including contaminated surfaces
- 2. Exposure by sexual contact
- 3. Perinatal infection from HBsAg + mother

ACIP recommends hepatitis B vaccine for:

- All newborns, to be administered <u>"within 24 hours</u>" of birth, using <u>single</u> antigen vaccine; Dose 2 at 1-2 mos. of age and Dose 3 at 6-18 mos. of age.
- All children and adolescents less than 19 years of age who did not complete the series as an infant.
- All adults aged 19-59 years should receive Hep B vaccine
- Hepatitis B vaccine is recommended for adults age 60 years or older with risk factors for hepatitis B virus infection
- **People age 60 years or older without** known risk factors for hepatitis B virus infection **may** also complete a HepB vaccine series.

https://www.cdc.gov/vaccines/hcp/acip-recs/vaccspecific/hepb.html

Meningococcal Disease (caused by N. meningitidis)

- Usually presents as meningitis, bacteremia or both
 - Transmitted through direct contact with respiratory tract secretions from patients and asymptomatic carriers
 - Nasopharyngeal carriage rate is highest in adolescents and young adults in the U.S.
 - Incidence of meningococcal disease declined during 2020– 2021, but increased in 2022
 - Recent outbreaks in the US (people experiencing homelessness, men who have sex with men)
 - New strains emerging in the US Predominantly affecting racial and ethnic minority groups – Unclear how this will change overall epidemiology
 - More complete 2021 and 2022 data are needed
 - More years of data needed to understand post-COVID-19 epidemiology

https://www.cdc.gov/vaccines/schedules/

https://www.cdc.gov/vaccines/pubs/pinkbook/mening.html

Trends in Meningococcal Disease Incidence by Serogroup – United States, 2006–2022*



Source: NNDSS data with additional serogroup data from Active Bacterial Core surveillance (ABCs) and state health departments *2021 and 2022 data are preliminary

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https://www.cdc.gov/vaccines/acip/meetings/downloads/sli des-2023-02/slides-02-23/Mening-02-Rubis-508.pdf

Signs and Symptoms of Meningococcal Disease

- Symptoms of meningitis
 - Sudden onset of fever
 - Headache
 - Stiff neck
 - Photophobia
 - Nausea and vomiting
- Symptoms of meningococcemia
 - All of the above are possible
 - Cold hands and feet
 - Pruritic rash

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- Risk factors
 - Persistent complement component deficiencies
 - Asplenia,
 - HIV infection
 - Exposure during an outbreak; Travel/residence in a country where disease is endemic/epidemic
 - Household crowding, smoking,
 - Unvaccinated college freshmen in dorms (particularly serogroup B)
 - Military recruits





https://www.cdc.gov/mmwr/volumes/69/rr/rr6909a1.htm; MMWR, Sept 2020, Vol 69, RR 9

Quadrivalent Meningococcal Conjugate Vaccine (MCV4) (Men A,C,W, Y)

Menactra[™] licensed for 9 mos. through 55 years Menveo® licensed for ages 2 mos. through 55 years MenQuadfi® licensed for ages ≥ 2 yrs. of age

ACIP recommends for adolescents:

- Dose 1---age 11-12 years preferred
- Booster dose---age 16 years
- If 1st dose is received ≥16 years of age, a 2nd dose is not needed, unless they become at increased risk for meningococcal disease
- Effective July 1, 2021, for the 2021-2022 school year, a meningococcal conjugate (MCV4/MenACWY) booster was required for all high school students entering the 11th grade and who are 16 years of age or older.
- First-year college students who live in residential housing (if not previously vaccinated at age 16 years or older) or military recruits

https://www.cdc.gov/vaccines/schedules/

https://dph.georgia.gov/public-health-regulations/regulationsrule-making and https://dph.georgia.gov/immunization-section and 53

https://www.cdc.gov/vaccines/vpd/mening/hcp/index.html

Meningococcal Vaccines for High Risk Persons 6 weeks – 55 years*

Menactra[™] licensed for 9 mos. through 55 years Menveo® licensed for ages 2 mos. through 55 years MenQuadfi® licensed for ages ≥ 2 yrs. of age

Recommended for persons 2 months through 55 years**:

- human immunodeficiency virus (HIV)***
- Persistent complement component deficiency, complement inhibitor
- functional or anatomic asplenia (sickle cell disease)
- microbiologists exposed to isolates of *N. meningitidis*
- part of a community outbreak due to vaccine serogroups
- persons traveling internationally to regions with endemic meningococcal disease

For persons in any of these categories, consult the current ACIP Immunization Schedules for specific dosages and guidelines

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*https://www.cdc.gov/mmwr/volumes/69/rr/rr6909a1.htm

Serogroup B Meningococcal Vaccine

Bexsero® licensed for ages 10 through 25 years (2 dose) Trumenba® licensed for ages 10 through 25 years (2 or 3 dose)

ACIP recommends serogroup B meningococcal vaccine for*:

- Persons with persistent complement component deficiencies
- Persons with anatomic or functional asplenia
- Persons receiving complement inhibitor
- Microbiologists routinely exposed to isolates of *Neisseria meningitidis*

Persons considered at greater risk because of a serogroup B meningococcal disease outbreak**
 Based on shared clinical decision making:

 A Men B vaccine series <u>may</u> be administered to adolescents and young adults 16 through 23 years of age to provide short-term protection against most strains of Men B. Preferred age is 16-18 years.

Serogroup B Meningococcal Vaccine Administration

Bexsero® licensed for ages 10 through 25 years (2 dose) Trumenba® licensed for ages 10 through 25 years (2 dose or 3 dose) The 2 vaccine products are not interchangeable.

MenB-FHbp (Trumenba[®])

- 2 dose schedule administered at 0, 6 months
- Given to healthy adolescents who are not at increased risk for meningococcal disease
- 3 dose schedule administered at 0, 1-2, 6 months
- Given to persons at increased risk for meningococcal disease and for use during serogroup B outbreaks

MenB-4C (Bexsero®)

- 2 dose schedule 0, 1-2 months
- Given to healthy adolescents who are not at increased risk for meningococcal disease
- Given to persons at increased risk for meningococcal disease and for use during
 https://www.cdc.gov/mmwr/volumes/69/rr/rr6909a1.htm and serogroup B outbreaks https://www.cdc.gov/vaccines/schedules/hcp/imz/child-adolescent.html#note-

Meningococcal Vaccine Booster Recommendations*

For persons at continued risk

- Meningococcal quadrivalent vaccine for persons who remain at increased risk
- Persons ≥10 years of age who previously received a MenB vaccine series
- See *MMWR: Tables 2-11 https://www.cdc.gov/mmwr/volumes/69/rr/rr6909a1.htm#B1_down for further details.

https://www.cdc.gov/vaccines/hcp/aciprecs/vacc-specific/mening.html

RotaTeq® (Merck) and Rotarix® (GSK)*

- RV 5, RotaTeq®: 3 doses; ages 2, 4, 6 months
- RV 1,Rotarix®: 2 doses; ages 2 and 4 months
- ACIP recommendation:
- 2-3 doses depending on brand
- Administer either vaccine as directed below:
 - Minimum age for first dose: 6 weeks
 - Maximum age for first dose: 14 weeks 6 days. Do not start the series on or after age 15 weeks, 0 days
 - Minimum interval between doses: 4 weeks
 - Maximum age for any dose: 8 months 0 days
- If any dose is Rotateq®, 3 doses are required
- Use RotaTeq® if allergy to latex

https://www.cdc.gov/vaccines/vpd/rotavirus/index.html and

https://www.cdc.gov/vaccines/vpd/rotavirus/hcp/recommendations.htm

l and https://www.cdc.gov/vaccines/schedules/hcp/imz/child-

Types of Human Papilloma Virus (HPV)*

(More Than 200 Types Identified)



HPV Vaccine*

Gardasil 9[®] (9vHPV) <u>HPV types 6, 11, 16, 18, 31, 33, 45, 52, 58</u>

ACIP recommends HPV vaccine starting at age 11 or 12 years for:

- All males and females through 26 years of age
- Catch-up vaccination for persons through age 26 who are not adequately vaccinated

Gardasil 9 is now also licensed for all persons 9 through 45 yrs. of age**

- Use the 3-dose schedule for persons 15-45 years of age
- Based on shared clinical decision making, the series <u>may</u> be given to persons ages 27-45.

Reasons to Immunize Against HPV at age 11-12 Years

- Higher antibody level attained when given to pre-teens rather than to older adolescents or women
- At this age, more likely to be administered before onset of sexual activity
- HPV can be transmitted by other skin-to-skin contact, not just sexual intercourse
- There is no link between vaccine and riskier sexual behavior
- Even those who abstain from sex until marriage can be infected by their marital partner
- Individuals need to complete the series for full protection
- This is an anti-cancer vaccine, and......

Over 90% of HPV cancers are preventable through HPV vaccination.

Bottom line: NOT receiving a healthcare provider's recommendation for HPV vaccine was <u>one of the main</u> reasons parents reported for <u>not</u> vaccinating their adolescent children.**

Presented by Anne Schuchat, MD, RADM US Public Health Service, Asst. Surgeon General, Director NCIRD at Immunize Georgia Conference, Atlanta, GA, 9-11-14

Increasing HPV Vaccination Rates Among Adolescents: Challenges and Opportunities. PolicyLab: The Children's Hospital of Philadelphia, 2016. http://www.immunize.org/askexperts/experts_hpv.asp

COVID-19 vaccination schedule for most people

Ages 6 months–4 years								
COVID-19 vaccination history	Bivalent vaccine	Number of bivalent doses indicated Dosage (mL/ug)	Vaccine vial cap and label colors	Interval betwee	en doses*			
Unvaccinated	Moderna or	Age 5 years						
	Pfizer BioNTech†	COVID-19 vaccination h	istory Biv	valent vaccine	Number of bivalent doses indicated	Dosage (mL/ug)	Vaccine vial cap and label colors	Interval between doses*
		Unvaccinated		Moderna	2	0.25 mL/25 ug	Dark blue cap;	Dose 1 and Dose 2:
1 dose monovalent Moderna	Moderna			or			gray label border	4–8 weeks
			Pfi	izer BioNTech	1	0.2 mL/10 ug	Orange	
2 doses monovalent Moderna	Moderna	1 dose monovalent Moderna		Moderna <i>or</i>	1	0.25 mL/25 ug	Dark blue cap; gray label border	4–8 weeks after monovalent dose
2 doses monovalent Moderna and 1	NA; previously received 1		Pfi	izer BioNTech	1	0.2 mL/10 ug	Orange	At least 8 weeks after monovalent dose

Ages 6–11 years

Bivalent vaccine	Number o doses in
Moderna	1
or	
Pfizer BioNTech	1
Moderna	1
or	
Pfizer BioNTech	1
NA; previously received 1 bivalent vaccine dose	N
NA; previously received 1 bivalent vaccine dose	N
	Bivalent vaccine Moderna or Pfizer BioNTech Moderna or Pfizer BioNTech NA; previously received 1 bivalent vaccine dose NA; previously received 1 bivalent vaccine dose

Ages 12 years and older

COVID-19 vaccination history	Bivalent vaccine	Number of bivalent doses indicated	Dosage (mL/ug)	Vaccine vial cap and label colors	Interval between doses*
Unvaccinated	Moderna <i>or</i>	1	0.5 mL/50 ug	Dark blue cap; gray label border	_
	Pfizer BioNTech	1	0.3 mL/30 ug	Gray	_
1 or more doses monovalent mRNA (no doses bivalent mRNA)	Moderna or	1	0.5 mL/50 ug	Dark blue cap; gray label border	At least 8 weeks after last monovalent dose
	Pfizer BioNTech	1	0.3 mL/30 ug	Gray	At least 8 weeks after last monovalent dose
Ever received 1 dose bivalent mRNA (regardless of monovalent vaccine history)	NA; previously received 1 bivalent vaccine dose	NA	NA	NA	NA

Moderna

or

0.2 mL/10 ug

Dark pink cap;

yellow label

At least 8 weeks after

last monovalent dose

People ages 65 years and older have the option to receive 1 additional bivalent mRNA vaccine dose at least 4 months after the first dose of a bivalent mRNA vaccine. If Moderna is used, administer 0.5 mL/50 ug (dark blue cap and label with a gray border); if Pfizer-BioNTech is used, administer 0.3 mL/30 ug (gray cap and label with a gray border).

https://www.cdc.gov/vaccines/covid-19/clinical-considerations/interimconsiderations-us.html#not-immunocompromised

05/4/2023

Table 2. Recommended COVID-19 vaccination schedule for people who are moderately or severely immunocompromised Ages 6-11 years COVID-19 vaccination history, May 2023 Ages 6-11 years

mRNA COVID-19 vaccines

Ages 6 months–4 years

	Bivalent	Number of bivalent		Vaccine vial cap and	
COVID-19 vaccination history	vaccine	doses indicated*	Dosage (mL/ug)	label colors	Interval between dose

Uni Ages 5 years

COVID-19 vaccination history	Bivalent vaccine	Number of bivalent doses indicated*	Dosage (mL/ug)	Vaccine vial cap and label colors	Interval betweer	1 dose
Unvaccinated	Moderna [†] <i>or</i> Pfizer	3	0.25 mL/25 ug	Blue cap; gray label border	Dose 1 and Dc 4 weeks Dose 2 and Dose 3 4 weeks	2 doses
	BioNTech	3	0.2 mL/10 ug	Orange	Dose 1 and Dc 3 weeks Dose 2 and dose 3 4 weeks	3 doses
1 dose monovalent Moderna	Moderna [†]	2	0.25 mL/25 ug	Blue cap; gray label border	Dose 1: 4 weeks monovalent c Dose 1 and Dc At least 4 we	3 doses
2 doses monovalent Moderna	Moderna†	1	0.25 mL/25 ug	Blue cap; gray label border	At least 4 weeks a monovalent c	dose bi 1 dose
3 doses monovalent Moderna	Moderna <i>or</i>	1	0.25 mL/25 ug	Blue cap; gray label border	At least 8 weeks a monovalent c	
	Pfizer BioNTech	1	0.2 mL/10 ug	Orange	At least 8 weeks afte monovalent dos	er last se
3 doses monovalent Moderna and 1 dose bivalent mRNA	-	See footnote	-	_	_	
1 dose monovalent Pfizer-BioNTech	Pfizer- BioNTech	2	0.2 mL/10 ug	Orange	Dose 1: 3 weeks a monovalent dos Dose 1 and Dose At least 4 week	fter Se 2: s

COVID-	19 Vaccine	schedule for	people	moderately	or
severely	y immunoco	mpromised			

https://www.cdc.gov/vaccines/covid-19/clinical-considerations/interimconsideନିର୍ଣ୍ଣିଶନ୍ତିୟିନ୍ତି.html#immunocompromised

COVID-19 vaccination history	Bivalent vaccine	Number of bivalent doses indicated*	Dosage (mL/ug)	Vaccine vial cap and label colors	Interval between doses
Unvaccinated	Moderna [†] <i>or</i> Pfizer-	3	0.25 mL/25 ug	Blue cap; gray label border	Dose 1 and Dose 2: 4 weeks Dose 2 and Dose 3: At least 4 weeks
	BioNTech [‡]	3	0.2 mL/10 ug	Orange	Dose 1 and Dose 2: 3 weeks Dose 2 and dose 3: At least 4 weeks
1 dose monovalent Moderna	Moderna [†]	2	0.25 mL/25 ug	Blue cap; gray label border	Dose 1: 4 weeks after monovalent dose Dose 1 and Dose 2: At least 4 weeks

Ages 12 years and older

COVID-19 vaccination history	Bivalent vaccine	Number of bivalent doses indicated*	Dosage (mL/ug)	Vaccine vial cap and label colors	Interval between doses
Unvaccinated	Moderna [†] <i>or</i> Pfizer	3	0.5 mL/50 ug	Blue cap; gray label border	Dose 1 and Dose 2: 4 weeks Dose 2 and Dose 3: At least 4 weeks
	BioNTech [‡]	3	0.3 mL/30 ug	Gray	Dose 1 and Dose 2: 3 weeks Dose 2 and dose 3: At least 4 weeks
1 dose monovalent Moderna	Moderna [†]	2	0.5 mL/50 ug	Blue cap; gray label border	Dose 1: 4 weeks after monovalent dose Dose 1 and Dose 2: At least 4 weeks
2 doses monovalent Moderna	Moderna†	1	0.5 mL/50 ug	Blue cap; gray label border	At least 4 weeks after last monovalent dose
3 doses monovalent Moderna	Moderna <i>or</i>	1	0.5 mL/50 ug	Blue cap; gray label border	At least 8 weeks after last monovalent dose
	Pfizer- BioNTech	1	0.3 mL/30 ug	Gray	At least 8 weeks after last monovalent dose
3 doses monovalent Moderna and 1 dose bivalent mRNA	_	See footnote	_	_	_

What are the main updates?

- The new guidance allows:
 - older adults and
 - immunocompromised adults (adults with a weakened immune system)
 to get a second dose of the updated bivalent vaccine.
- The FDA and CDC made this recommendation because older adults and people with weakened immune systems are at higher risk for severe COVID-19, and data show that the effectiveness of COVID-19 vaccines wanes over time. An additional dose of the updated vaccine offers these two groups of individuals extra protection from getting seriously ill with COVID-19



What are the main updates (2)?

 Also moving forward, the monovalent Moderna and Pfizer-BioNTech COVID-19 vaccines (the first COVID-19 vaccines that protect against the original COVID-19 virus strain only) are no longer authorized for use in the United States.



Vaccination of Children and Adolescents

- CDC recommends that people ages 6 months and older receive at least 1 bivalent mRNA COVID-19 vaccine.
- At the time of initial vaccination, depending on vaccine product, children ages 6 months–4 years are recommended to receive 2 or 3 bivalent mRNA vaccine doses; children age 5 years are recommended to receive 1 or 2 bivalent mRNA vaccine doses
- Please utilize the schedule based on age and receipt of prior doses.

COVID-19 vaccination schedule changes for people moderately or severely immunocompromised

•At the time of initial vaccination, people ages 6 months and older are recommended to receive 3 bivalent mRNA doses

•People ages 6 months and older who previously received only monovalent doses are recommended to receive 1 or 2 bivalent mRNA vaccine doses, depending on age and vaccine product

•People who previously received a bivalent mRNA vaccine dose(s) have the option to receive 1 or more additional bivalent mRNA doses

https://www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html#table-01

Critical Elements for Immunization Services



The 7 Rights of Vaccine Administration

Right Patient ✓ Right Vaccine or Diluent ✓ Right *Time** ✓ Right **Dosage** Right Route, Needle Length, Technique Right Site for route indicated ✓ Right **Documentation**

* Correct age, appropriate interval, and administer before vaccine or diluent expires



Ref: Epidemiology and Prevention of Vaccine-Preventable Diseases. 13th Edition, 2015.

Appropriate Vaccine Storage & Handling is <u>Very Important</u>

- Store all vaccines as recommended by manufacturer
- Monitor and record temperatures of refrigerator and freezer twice daily
- Take immediate action for all out-of-range temps
- Implement a vaccine emergency system
- Maintain temperature log records for 3 years
- DO NOT STORE ANYTHING ELSE in the refrigerator!

Check Expiration Date of Vaccines and Diluents



Note: Some multidose vials have a beyond use date (BUD) that becomes effective once the vial is entered with a needle. This date may vary from the expiration date printed on the vial. Consult package insert, but be sure to indicate this BUD date change on the vial.

Sites for Vaccine Administration

Intramuscular (IM)

DTaP, Tdap, Hib, Td, Hep A, Hep B, PCV13, IIV, MCV4, HPV , Herpes Zoster, COVID-19

Subcutaneous (SQ, SC, or sub-Q)

MMR (GSK product Priorix), MPSV4, Herpes Zoster (1 dose vaccine)

Either intramuscular or subcutaneous IPV, PPSV23, MMR, MMRV, Varicella

<u>Intranasal</u> LAIV










Vaccine Administration Best practices – Route, Dose, Site, Needle Size

Administering Vaccines: Dose, Route, Site, and Needle Size

Vaccine		Dose	Route	Injection Site and Nee	edle Size			
COVID-19	Pfizer-BioNTech •age 5 to <12 yrs: 0.2 mL pe •age ≥12 yrs: 0.3 mL adult/a primary and booster doses	Tech 12 yrs: 0.2 mL pediatric formulation ("orange cap") rs: 0.3 mL adult/adolescent formulation for nd booster doses		IM Subcutaneous (Subcut Use a 23–25 gauge needle to the person's age and bc		i injection Choose the injection site that is appropriate dy mass.		
	Moderna; ≥18 yrs: 0.5 mL p Janssen: ≥18 yrs: 0.5 mL for	imary series*; 0.25 mL booster primary & booster doses		AGE	NEEDLE LENGTH	INJECTION SITE		
Diphtheria, 1 (DTaP, DT, T	Tetanus, Pertussis dap, Td)	0.5 mL	ІМ	Infants (1–12 mos)	5/8"	Fatty tissue over anterolat- eral thigh muscle		
Haemophilu	s influenzae type b (Hib)	0.5 mL	IM	Children 12 mos or older		Fatty tissue over anterolat- eral thigh muscle or fatty		
	(1.1	≤18 yrs: 0.5 mL		adolescents, and adults	5/8"			
Hepatitis A	(нера)	≥19 yrs: 1.0 mL	IM	Intramuscular (IM) injection				
Hepatitis B Persons 11–15 yrs	(HepB) may be given Recombivax HB	Engerix-B; Recombivax HB ≤19 yrs: 0.5 mL ≥20 yrs: 1.0 mL	IM	Use a 22–25 gauge needle. Choose the injection site and needle length that is appropriate to the person's age and body mass.				
(Merck) 1.0 mL adult formulation on a 2-dose schedule.		Heplisav-B ≥18 yrs: 0.5 mL		AGE	NEEDLE LENGTH	INJECTION SITE		
Human papillomavirus (HPV)		0.5 mL	IM	Newborns (1st 28 days)	5/8"1	Anterolateral thigh muscle		
		0.2 ml (0.1 ml in each	Intra-	Infants (1-12 mos)	1"	Anterolateral thigh muscle		
Influenza, liv	e attenuated (LAIV)	nostril)	nasal	Taddlara (1. 2. vaara)	1–11⁄4"	Anterolateral thigh muscle ²		
		Afluria: 0.25 ml	spray	l loddiers (1–2 years)	5⁄8—1"1	Deltoid muscle of arm		
Influenza in	activated (IIV): for ages	Eluzone: 0.25 or 0.5 ml		Children	5⁄8—1"1	Deltoid muscle of arm ²		
6–35 month	s	Fluariz Elucebraz Elutaval:	IM	(3–10 years)	1–1¼"	Anterolateral thigh muscle		
		0.5 mL		Adolescents and teens	5/8—1" ¹	Deltoid muscle of arm ²		
Influenza, in	activated (IIV), ≥3 yrs;	0.5 mL		(11–18 years)	1–11/2"	Anterolateral thigh muscle		
recombinant	t (RIV), ≥18 yrs; HD-IIV) >65 yrs	EluZone HD: 0.7 ml	IM	Adults 19 years or older				
inginuose (110-114) 203 yrs			i					

Measles, Mumps, Rubella (MMR)	0.5 mL	Subcu	
Meningococcal serogroups A, C, W, Y (MenACWY)	0.5 mL	IM	
Meningococcal serogroup B (MenB)	0.5 mL	IM	
Pneumococcal conjugate (PCV)	0.5 mL	IM	
Pneumococcal polysaccharide (PPSV)	0.5 mL	IM or Subcu	
Polio, inactivated (IPV)	0.5 mL	IM or Subcu	
Potenting (DV)	Rotarix: 1.0 mL	Oral	
Rotavirus (RV)	Rotateq: 2.0 mL	Orai	
Varicella (VAR)	0.5 mL	Subcu	
Zoster (Zos)	Shingrix: 0.5 [†] mL	IM	
Combination Vaccines			
DTaP-HepB-IPV (Pediarix) DTaP-IPV/Hib (Pentacel) DTaP-IPV (Kinrix; Quadracel) DTaP-IPV-Hib-HepB (Vaxelis)	0.5 mL	IM	
MMRV (ProQuad)	≤12 yrs: 0.5 mL	Subcu	
HepA-HepB (Twinrix)	≥18 yrs: 1.0 mL	IM	
* If immunocompromised, Moderna 0.5 mL fo 3-dose primary series, then 0.25 mL for boost dose. [†] The Shingrix vial might contain more than 0.5 mL. Do not administer more than 0.5 mL	r Intranasal (NAS) administration of Flumist (LAIV) vaccine		

Female or male <130 lbs	5/8-1"1	Deltoid muscle of arm
Female or male 130–152 lbs	1"	Deltoid muscle of arm
Female 153–200 lbs Male 153–260 lbs	1–11⁄2"	Deltoid muscle of arm
Female 200+ lbs Male 260+ lbs	11⁄2"	Deltoid muscle of arm
Female or male, any weight	11⁄2"	Anterolateral thigh muscle

¹ A 5/8° needle may be used in newborns, preterm infants, and patients weighing less than 130 lbs (<60 kg) for IM injection in the deltoid muscle only if the skin stretched tight, the subcutaneous tissue is not bunched, and the injection is made at a 90-degree angle to the skin.
² Preferred site

Intr

NOTE: Always refer to the package insert included with each biologic for complete vaccine administration information. CDC's Advisory Committee on Immunization Practices (ACIP) recommendations for the particular vaccine should be reviewed as well. Access the ACIP recommendations at www.immunize.org/acip.

amuscular (IM) injection	Subcutaneous (Subcut) injection				
90° angle	45° angle				
cutaneous tissue muscle	skin subcutaneous tissue muscle				

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 www.immunize.org/catg.d/p3085.pdf . Item #P3085 (11/21)

https://www.immunize.org/catg.d/p3085.pdf

How to administer IM and SC vaccine injections

NEEDLE SIZE

5%"* (22-25 gauge)

How to Administer Intramuscular and Subcutaneous Vaccine Injections Administration by the Intramuscular (IM) Route

INIECTION SITE

Anterolateral thigh muscle

PATIENT AGE

Newborn (0-28 days)

Administer these vaccines via IM route

- Diphtheria-tetanus-pertussis (DTaP, Tdap)
- Diphtheria-tetanus (DT, Td)
- = Haemophilus influenzae type b (Hib)
- = Hepatitis A (HepA)
- Hepatitis B (HepB)
- Human papillomavirus (HPV)
 Inactivated influenza (IIV)
- Meningococcal serogroups A,C,W, Y (MenACWY)
- Meningococcal serogroup B (MenB)
 Pneumococcal conjugate (PCV13)
 Zoster, recombinant (RZV)

Administer inactivated polio (IPV) and pneumococcal polysaccharide (PPSV23) vaccines either IM or

subcutaneously (Subcut).

Infant (1-12 mos) 1" (22–25 gauge) Anterolateral thigh muscle Anterolateral thigh muscle 1-11/4" (22-25 gauge) Toddler (1–2 years) Alternate site: Deltoid muscle of arm if 5/8*-1" (22-25 gauge) muscle mass is adequate 5/8*-1" (22-25 gauge) Deltoid muscle (upper arm) Children (3-10 years) Alternate site: Anterolateral thigh 1-11/4" (22-25 gauge) muscle Deltoid muscle (upper arm) 5/8[†]-1" (22-25 gauge) Children and adults Alternate site: Anterolateral thigh (11 years and older) 1-11/2" (22-25 gauge) muscle

* A 3/2" needle usually is adequate for neonates (first 28 days of life), preterm infants, and children ages 1 through 18 years if the skin is stretched flat between the thumb and forefinger and the needle is inserted at a 90° angle to the skin.

† A 5%" needle may be used in patients weighing less than 130 lbs (<60 kg) for IM injection in the deltoid muscle only if the skin is stretched flat between the thumb and forefinger and the needle is inserted at a 90° angle to the skin; a 1" needle is sufficient in patients weighing 130–152 lbs (60–70 kg); a 1–1½" needle is recommended in women weighing 153–200 lbs (70–90 kg) and men weighing 153–260 lbs (70–118 kg); a 1½" needle is recommended in women weighing more than 200 lbs (91 kg) or men weighing more than 260 lbs (118 kg).



Needle insertion

Use a needle long enough to reach deep into the muscle.

Insert needle at a 90° angle to the skin with a quick thrust.

(Before administering an injection of vaccine, it is not necessary to aspirate, i.e., to pull back on the syringe plunger after needle insertion.[¶])

Multiple injections given in the same extremity should be separated by a minimum of 1", if possible.

CDC. "General Best Practices Guidelines for Immunization: Best Practices Guidance of the ACIP" at https://www.cdc.gov/vaccines/ hcp/acip-recs/general-recs/downloads/ general-recs.pdf



Intramuscular (IM) injection site for infants and toddlers

Insert needle at a 90° angle into the

anterolateral thigh muscle.

IM injection site

(shaded area)

Intramuscular (IM) injection site for children and adults



Give in the central and thickest portion of the deltoid muscle – above the level of the armpit and approximately 2–3 fingerbreadths (~2") below the acromion process. See the diagram. To avoid causing an injury, do not inject too high (near the acromion process) or too low.

CONTINUED ON THE NEXT PAGE

https://www.immunize.org/catg.d/p2020.pdf

5/6/2023

SIRVA Shoulder anatomy



<u>SIRVA</u> = <u>Shoulder Injury</u> <u>Related to Vaccine</u> <u>Administration</u>

TIPS TO AVOID THIS INJURY

- Landmark the site---don't "eyeball" it
- If possible, be seated to vaccinate a seated pt.
- Expose the shoulder completely
- Roll the sleeve up---don't pull the shirt over the neck
 - REMEMBER! 2-3 FINGERS DOWN FROM THE ACROMION

 \bullet

https://www.hhs.gov/sites/default/files/Nair_Special%20Highlight_SIRVA%20remediated.pdf

Training Tools: Skills Checklist for Vaccine Administration

Skills Checklist for Vaccine Administration

During the COVID-19 pandemic, the CDC recommends additional infection control measures for vaccination (see www.cdc.gov/vaccines/pandemic-

The Skills Checklist is a self-assessment tool for healthcare staff who administer vaccines to several patients, and score in the Supervisor administer immunizations. To complete it, review the competency areas below and the clinical skills, techniques and procedures outlined for each area. Score yourself in the Self-Assessment column. If you check Needs to Improve, you indicate further study, practice, others, or change is needed. When you check Meets or Exceeds, you indicate you believe you are performing at the expected level of competence, The video "Immunization Techniques: Best Practices with Infants,

or higher. Supervisors: Use the Skills Checklist to clarify responsibilities and correctly. (View at www.youtube.com/watch?v=WsZ6NEiilfl or order expectations for staff who administer vaccines. When you use it to online at www.immunize.org/dvd.) Another helpful resource is assist with performance reviews, give staff the opportunity to score CDC's Vaccine Administration eLearn course, available at www.cdc. themselves in advance. Next, observe their performance as they

Review columns. If improvement is needed, meet with them to develop a Plan of Action (see bottom of page 3) to help them achieve the level of competence you expect: circle desired actions or write in

> Children, and Adults" helps ensure that staff administer vaccines gov/vaccines/hcp/admin/resource-library.html.

> > Administering

Immunization

guidance/index.html).								
				Supervisor Review				
COMPETENCY	CLINICAL SKILLS, TECHNIQUES, AND PROCEDURES	NEEDS TO IMPROVE	MEETS OR EXCEEDS	NEEDS TO IMPROVE	MEETS OR EXCEEDS	PLAN OF ACTION		
A	 Welcomes patient/family and establishes rapport. 							
Patient/Parent	 Explains what vaccines will be given and which type(s) of injection(s) will be done. 							
Lucation	 Answers questions and accommodates language or literacy barriers and special needs of patient/parents to help make them feel comfortable and informed about the procedure. 							
	 Verifies patient/parents received Vaccine Information Statements (VISs) for indicated vaccines and has had time to read them and ask questions. 							
	Screens for contraindications (if within employee's scope of work).			Skills Che	cklist for Vacc	ine Administration (continu	ued)	
	 Reviews comfort measures and aftercare instructions with patient/parents, and invites questions. 						,	
0	1. Identifies the location of the medical protocols (e.g., immunization							
Medical and	protocol, emergency protocol, reporting adverse events to the Vaccine Adverse Event Reporting system [VAERS], reference material).		COMPETENCY CLINICAL SKILI		LS, TECHNIQUES, AND PROCEDURES			
Office Protocols	 Identifies the location of epinephrine, its administration technique, and clinical situations where its use would be indicated. 			G Vaccine Preparation		1. Performs proper hand hygiene prior to preparing vaccine.		
	3. Maintains up-to-date CPR certification.		<u> </u>			When removing vaccine from the refrigerator or freezer, looks at the storage unit's temperature to make sure it is in proper range.		
	 Understands the need to report any needlestick injury and to maintain a sharps injury log. 					 Checks vial expiration date. Double-checks vial label and contents pri to drawing up. 		ole-checks vial label and contents prior
	 Demonstrates knowledge of proper vaccine handling (e.g., maintains and monitors vaccine at recommended temperature and protects from light). 		4. Prepares and draws is not adjacent to a		 Prepares and draws up is not adjacent to areas 	ws up vaccines in a designated clean medication area th areas where potentially contaminated items are place		

CONTINUED ON THE NEXT PAGE

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KINS CHECKIST IOF Vac	cine Administration (continued)					page 50
		Self-Ass	essment		Superviso	or Review
COMPETENCY	COMPETENCY CLINICAL SKILLS, TECHNIQUES, AND PROCEDURES		MEETS OR EXCEEDS	NEEDS TO IMPROVE	MEETS OR EXCEEDS	PLAN OF ACTION
D Administering	 Controls the limb with the non-dominant hand; holds the needle an inch from the skin and inserts it quickly at the appropriate angle (90° for IM or 45° for Subcut). 					
Immunizations	9. Injects vaccine using steady pressure; withdraws needle at angle of insertion.					
(continued)	10. Applies gentle pressure to injection site for several seconds (using, e.g., gauze pad, bandaid).					
	11. Uses strategies to reduce anxiety and pain associated with injections.					
	12. Properly disposes of needle and syringe in "sharps" container.					
	13. Properly disposes of vaccine vials.					
8	 Fully documents each vaccination in patient chart: date, lot number, manufacturer, site, VIS date, name/initials. 					
Records Procedures	 If applicable, demonstrates ability to use state/local immunization registry or computer to call up patient record, assess what is due today, and update computerized immunization history. 					
	 Asks for and updates patient's vaccination record and reminds them to bring it to each visit. 					

Plan of Action Circle desired next steps and write in the resource-library.html. agreed deadline for b. Review office protocols. completion, as well as date for the follow-up nerformance review d. Review package inserts. lines or video f. Observe other staff with patients.

a. Watch video on immunization techniques and g. Practice injections review CDC's Vaccine Administration eLearn, h. Read Vaccine Information Statements available at www.cdc.gov/vaccines/hcp/admin/ i. Be mentored by someone who appropriate immunization ski j. Role play (with other staff) int c. Review manuals, textbooks, wall charts, or parents and patients, includir other guides (e.g., Key Vaccination Resources comfort measures. for Healthcare Professionals at k. Attend a skills training or othe www.immunize.org/catg.d/p2005.pdf courses/training I. Attend healthcare customer s e. Review vaccine storage and handling guide cultural competency training. m. Renew CPR certification.

File the	Skills	Checklist in	the	employee's personnel	
folder					

has demonstrated ills.	
eractions with g age appropriate	PLAN OF ACTION DEADLINE
er appropriate	DATE OF NEXT PERFORMANCE REV
atisfaction or	EMPLOYEE SIGNATURE
	SUPERVISOR SIGNATURE

IMMUNIZATION ACTION COALITION Saint Paul, Minnesota • 651-647-9009 • www.immunize.org • www.vaccineinformation.org www.immunize.org/catg.d/p7010.pdf • Item #P7010 (2/21)

Other

https://www.immunize.or g/catg.d/p7010.pdf

CONTINUED ON THE NEXT PAGE

tissue over triceps).

5. Selects the correct needle size for IM and Subcut based on patient age

6. Maintains aseptic technique throughout, including cleaning the rubbe

7. Prepares vaccine according to manufacturer instructions. Inverts vial and

the expiration date on the equipment (syringes and needles) if present. 9. Labels each filled syringe or uses labeled tray to keep them identified. 1. Verifies identity of patient. Rechecks the provider's order or instructions

2. Utilizes proper hand hygiene with every patient and, if it is office policy, put

on disposable gloves. (If using gloves, changes gloves for every patient.) 3. Demonstrates knowledge of the appropriate route for each vaccine

4. Positions patient and/or restrains the child with parent's help. 5. Correctly identifies the injection site (e.g., deltoid, vastus lateralis, fatty

6. Locates anatomic landmarks specific for IM or Subcut injections. 7. Preps the site with an alcohol wipe, using a circular motion from the center to a 2" to 3" circle. Allows alcohol to dry.

and/or weight, site, and recommended injection technique

septum (stopper) of the vial with alcohol prior to piercing it.

draws up correct dose of vaccine. Rechecks vial label 8. Prepares a new sterile syringe and sterile needle for each injection. Check

against the vial and the prepared syringes.

Self-Assessment NEEDS TO MEETS OR NEEDS T

MPROVE EXCEEDS

IMPROV

General Best Practice Guidelines for Immunization*

(formerly General Recommendations on Immunization)

- Timing and Spacing
- Contraindications and Precautions
- Prevention and Management of Adverse Reactions
- Vaccine Administration
- Storage and Handling of Immunobiologics
- Altered Immunocompetence
- Special Situations
- Vaccination Records
- Vaccination Programs
- Vaccine Information Sources
- Errata available at https://www.cdc.gov/vaccines/hcp/acip-recs/generalrecs/general-recs-errata.html

5/6/2023

*Kroger AT, Duchin J, Vázquez M. General Best Practice Guidelines for Immunization. Best Practices Guidance of the Advisory Committee on Immunization Practices (ACIP). [www.cdc.gov/vaccines/hcp/acip-recs/general-recs/downloads/general-recs.pdf].



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- Information sheets produced by the CDC
- Explanation of risks and benefits of a vaccine
- Federal law requires that a VIS be handed to patient/parent before each dose of vaccine is given
- Must be provided for any vaccine that is covered by the Vaccine Injury Compensation Program
- Available through Immunization Action Coalition (IAC) at www.immunize.org

Always Document...

Accept only written documentation of prior immunizations

After vaccine administration <u>document</u>:

- ✓ Publication date of VIS & Date VIS given
- ✓ Date, site, route, antigen(s), manufacturer, lot #
- Person administering vaccine, practice name and address
- ✓ Vaccine refusals with a signed "Refusal to Vaccinate Form"



Exemptions From School/Day Care Requirements

Medical Exemption O.C.G.A. §20-2-771(d)

- Used when a physical disability or medical condition contraindicates a particular vaccine.
- Requires an <u>annual review.</u>
- The medical exemption is documented in GRITS.

Religious Exemption O.C.G.A. §20-2-771(e)

- Parent or guardian must be directed to http://dph.georgia.gov/immunization-section to obtain an Affidavit of Religious Objection to Immunization form.
- This form must be signed and notarized and provided to the school.
- Must be kept on file at school/facility in lieu of an immunization certificate.
- Affidavit does not expire.

5/6/2023

Georgia does NOT have a philosophical exemption.



A 'Birth to Death' Immunization Registry

- Providers administering vaccines in Georgia must provide appropriate information to GRITS.
- GRITS personnel can work with your EHR/EMR vendor to create an interface between your system and GRITS.
- Use GRITS to generate reminders on medical records and/or notify patients when vaccines are needed.
- Assess your immunization rates using GRITS to improve patient care, HEDIS scores, and identify problem areas.

Call the GRITS Training Coordinator (404) 463-0807 or e-mail : https://dph.georgia.gov/georgia-immunization-registry-grits

Recommended Healthcare Personnel Vaccinations

- Hepatitis B (exposure risk) check immunity
- Influenza (annual)
- Measles, Mumps, Rubella (MMR)
- Varicella (Chickenpox)
- Tetanus, Diphtheria, Pertussis (Tdap)
- Meningococcal (recommended for microbiologists who are routinely exposed to isolates of N. meningitidis).
- COVID-19 vaccine

5/6/2023 Are YOU up to date?

Healthcare Personnel Vaccination Recommendations¹

VACCINES AND RECOMMENDATIONS IN BRIEF

- Hepatitis B If previously unvaccinated, give a 2-dose (Heplisav-B) or 3-dose (Engerix-B or Recombivax HB) series. Give intramuscularly (IM). For HCP who perform tasks that may involve exposure to blood or body fluids, obtain anti-HBs serologic testing 1-2 months after dose #2 (for Heplisav-B) or dose #3 (for Engerix-B or Recombivax HB).
- Influenza Give 1 dose of influenza vaccine annually. Inactivated injectable vaccine is given IM. Live attenuated influenza vaccine (LAIV) is given intranasally.
- MMR For healthcare personnel (HCP) born in 1957 or later without serologic evidence of immunity or prior vaccination, give 2 doses of MMR, 4 weeks apart. For HCP born prior to 1957, see below. Give subcutaneously (Subcut).
- Varicella (chickenpox) For HCP who have no serologic proof of immunity, prior vaccination, or diagnosis or verification of a history of varicella or herpes zoster (shingles) by a healthcare provider, give 2 doses of varicella vaccine, 4 weeks apart. Give Subcut
- Tetanus, diphtheria, pertussis Give 1 dose of Tdap as soon as feasible to all HCP who have not received Tdap previously and to pregnant HCP with each pregnancy (see below). Give Td or Tdap boosters every 10 years thereafter. Give IM.
- Meningococcal Give both MenACWY and MenB to microbiologists who are routinely exposed to isolates of Neisseria meningitidis. As long as risk continues: boost with MenB after 1 year, then every 2-3 years thereafter; boost with MenACWY every 5 years, Give MenACWY and MenB IM.

Hepatitis A, typhoid, and polio vaccines are not routinely recommended for HCP who may have on-the-job exposure to fecal material

Hepatitis B

Unvaccinated healthcare personnel (HCP) and/ or those who cannot document previous vaccination should receive either a 2-dose series of Heplisav-B at 0 and 1 month or a 3-dose series of either Engerix-B or Recombivax HB at 0, 1, and 6 months. HCP who perform tasks that may involve exposure to blood or body fluids should be tested for hepatitis B surface antibody (anti-HBs) 1-2 months after dose #2 of Heplisav-B or dose #3 of Engerix-B or Recombivax HB to document immunity.

 If anti-HBs is at least 10 mIU/mL (positive), the vaccinee is immune. No further serologic testing or vaccination is recommended.

 If anti-HBs is less than 10 mIU/mL (negative), the vaccinee is not protected from hepatitis B virus (HBV) infection, and should receive another 2-dose or 3-dose series of HepB vaccine on the routine schedule, followed by anti-HBs testing 1-2 months later. A vaccinee whose anti-HBs remains less than 10 mIU/ mL after 2 complete series is considered a "non-responder."

For non-responders: HCP who are non-responders should be considered susceptible to HBV and should be counseled regarding precautions to prevent HBV infection and the need to obtain HBIG prophylaxis for any known or probable parenteral exposure to hepatitis B surface antigen (HBsAg)-positive blood or blood with unknown HBsAg status. It is also possible that nonresponders are people who are HBsAg positive. HBsAg testing is recommended. HCP found

to be HBsAg positive should be counseled and medically evaluated. For HCP with documentation of a complete

2-dose (Heplisav-B) or 3-dose (Engerix-B or Recombivax HB) vaccine series but no documentation of anti-HBs of at least 10 mIU/mL (e.g., those vaccinated in childhood): HCP who are at risk for occupational blood or body fluid exposure might undergo anti-HBs testing upon hire or matriculation. See references 2 and 3 for details.

students in these professions, and volunteers,

Influenza All HCP, including physicians, nurses, paramedics,

emergency medical technicians, employees of nursing homes and chronic care facilities,

> should receive annual vaccination against influenza. Live attenuated influenza vaccine (LAIV) may be given only to non-pregnant healthy HCP age 49 years and younger. Inactivated injectable influenza vaccine (IIV) is preferred over LAIV for HCP who are in close contact with severely immunosuppressed patients (e.g., stem cell transplant recipients) when they require protective isolation.

> > Measles, Mumps, Rubella (MMR) HCP who work in medical facilities should be

immune to measles, mumps, and rubella. HCP born in 1957 or later can be considered immune to measles, mumps, or rubella only if they have documentation of (a) laboratory confirmation of disease or immunity or (b) appropriate vaccination against measles, mumps, and rubella (i.e., 2 doses of live

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It is recommended that all HCP be immune to varicella. Evidence of immunity in HCP includes documentation of 2 doses of varicella vaccine given at least 28 days apart, laboratory evidence of immunity, laboratory confirmation of disease. or diagnosis or verification of a history of varicella or herpes zoster (shingles) by a healthcare provider

outbreak of rubella.

Varicella

Tetanus/Diphtheria/Pertussis (Td/Tdap)

measles and mumps vaccines given on or after

the first birthday and separated by 28 days or

vaccine). HCP with 2 documented doses of MMR are not recommended to be serologically

tested for immunity; but if they are tested and

results are negative or equivocal for measles,

mumps, and/or rubella, these HCP should be

considered to have presumptive evidence of immunity to measles, mumps, and/or rubella

and are not in need of additional MMR doses.

Although birth before 1957 generally is con-

vaccine should be considered for unvacci-

nated HCP born before 1957 who do not have

laboratory evidence of disease or immunity to

measles and/or mumps. One dose of MMR vaccine should be considered for HCP with no

laboratory evidence of disease or immunity

to rubella. For these same HCP who do not

have evidence of immunity, 2 doses of MMR

of measles or mumps and 1 dose during an

vaccine are recommended during an outbreak

sidered acceptable evidence of measles. mumps, and rubella immunity, 2 doses of MMR

more, and at least 1 dose of live rubella

All HCPs who have not or are unsure if they have previously received a dose of Tdap should receive a dose of Tdap as soon as feasible, without regard to the interval since the previous dose of Td. Pregnant HCP should be revaccinated during each pregnancy. All HCPs should then receive Td or Tdap boosters every 10 years thereafter.

Meningococcal

Vaccination with MenACWY and MenB is recommended for microbiologists who are routinely exposed to isolates of N. meningitidis The two vaccines may be given concomitantly but at different anatomic sites, if feasible,

REFERENCES

1 CDC. Immunization of Health-Care Personnel: Recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR, 2011; 60(RR-7).

2 CDC. Prevention of Hepatitis B Virus Infection in the Unit ed States, Recommendations of the Advisory Committee on Immunization Practices. MMWR, 2018; 67(RR1):1-30 3 IAC. Pre-exposure Management for Healthcare Personnel with a Documented Hepatitis B Vaccine Series Who Have Not Had Post-vaccination Serologic Testing, Accessed at www.immunize.org/catg.d/p2108.pdf.

For additional specific ACIP recommendations, visit CDC's website at www.cdc.gov/vaccines/hcp/acip-recs/vaccspecific/index.html or visit IAC's website at www.immunize.org/acip

Stay Current!



 Sign up for listserv sites which provide timely information pertinent to your practice <u>www.immunize.org/resources/emailnews.asp</u>

- AAP Newsletter
- CDC immunization websites (32 in all)
- CHOP Parents Pack Newsletter
- IAC Express, Needle Tips and Vaccinate Adults
- Websites specific to particular vaccines



EVERYONE IS A PART OF THE TEAM THAT CAN

MAKE SURE PATIENTS RECEIVE THE

IMMUNIZATIONS THEY NEED!

Questions?

Contacts for more immunization information and resources!

National Center for Immunization and Respiratory Diseases, CDC

E-mail	NIPInfo@cdc.gov
Hotline	800.CDC.INFO
Website	http://www.cdc.gov/vaccines

Georgia Immunization Program

E-mail	DPH-Immunization@dph.ga.gov
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- Hotline 404-657-3158
- Website http://dph.georgia.gov/immunization-section

Immunization Action Coalition

E-mail	admin@immunize.org
Phone	651.647.9009
Website	www.immunize.org