APPENDIX D
Vaccine Administration

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Vaccine Administration

Appropriate vaccine administration is critical to vaccine effectiveness. The recommended site, route and dosage for each vaccine are based on clinical trials, practical experience and theoretical considerations. The following information provides general guidelines for administration of vaccines for those who administer vaccines, as well as those in training, education and supervisory positions. This information should be used in conjunction with professional standards for medication administration, vaccine manufacturers’ product guidelines, CDC’s Advisory Committee on Immunization Practices (ACIP) General Recommendations on Immunization, the American Academy of Pediatrics’ (AAP) Report of the Committee on Infectious Diseases Red Book, and state/agency-related policies and procedures. An education plan that includes competency-based training on vaccine administration should be considered for all persons who administer vaccines to children or adults (refer to "Skills Checklist for Immunization" - page D16).

Preparation

Patient Preparation - Patients should be prepared for vaccination with consideration for their age and stage of development. Parents/guardians and patients should be encouraged to take an active role before, during and after the administration of vaccines (see "Be There for Your Child During Shots" poster at http://cdlhn.com/default.htm, search for IMM674S).

- Screening - All patients should be screened for contraindications and precautions for each scheduled vaccine. Many state immunization programs and other organizations have developed and make available standardized screening tools. Basic screening questions can be found in Chapter 2. Sample screening forms for children and adults are available from the Immunization Action Coalition (www.immunize.org).

- Vaccine Safety & Risk Communication - Parents/guardians and patients are exposed through the media to information about vaccines, some of which is inaccurate or misleading. Healthcare providers should be prepared to discuss the benefits and risks of vaccines using Vaccine Information Statements (VIS) and other reliable resources. Establishing an open dialogue provides a safe, trust-building environment in which individuals can freely evaluate information, discuss vaccine concerns and make informed decisions regarding immunization (see Chapter 4 and Appendices E and F).

- Atraumatic Care - Vaccine safety issues and the need for multiple injections have increased the concerns and anxiety associated with immunizations. Healthcare providers need to display confidence and establish an environment that promotes a sense of security and trust for the patient and family, utilizing a variety of techniques to minimize the stress and discomfort associated with receiving injections. This is particularly important when administering vaccines to children.

- Positioning & Comforting Restraint - The healthcare provider must accommodate for the patient's comfort, safety, age, activity level, and the site of administration when considering patient positioning and restraint. For a child, the parent/guardian should be encouraged to hold the child during administration. If the parent is uncomfortable, another person may assist or the patient may be
positioned safely on an examination table (refer to "Comforting Restraint for Immunizations" - page D23).

- **Pain Control** - Pain is a subjective phenomenon influenced by multiple factors, including an individual’s age, anxiety level, previous healthcare experiences, and culture. Consideration for these factors is important as the provider develops a planned approach to management of injection pain (see “Be There for Your Child During Shots” poster).

- **Topical Anesthetics** or a vapocoolant spray may be applied to decrease pain at the injection site. These products should be used only for the ages recommended and as directed by the product manufacturer.

- **Analgesic Agents** - A non-aspirin containing pain reliever may be considered to decrease discomfort and fever following vaccination. These products should be used only in age-appropriate doses.

- **Diversionary Techniques** - Age-appropriate non pharmacologic techniques may provide distraction from pain associated with injections. Diversion can be accomplished through a variety of techniques, some of which are outlined on the “Be There for Your Child During Shots” poster.

- **Dual Administrators** - Some providers favor the technique of two individuals simultaneously administering vaccines at separate sites. The premise is that this procedure may decrease anxiety from anticipation of the next injection(s). The effectiveness of this procedure in decreasing pain or stress associated with vaccine injections has not been evaluated.

**Infection Control** - Healthcare providers should follow Standard Precautions to minimize the risks of spreading disease during vaccine administration.

- **Handwashing** - The single, most effective disease prevention activity is good handwashing. Hands should be washed thoroughly with soap and water or cleansed with an alcohol-based waterless antiseptic between patients, before vaccine preparation or any time hands become soiled, e.g. diapering, cleaning excreta.

- **Gloving** - Gloves are not required to be worn when administering vaccines unless the person administering the vaccine is likely to come into contact with potentially infectious body fluids or has open lesions on the hands. It is important to remember that gloves cannot prevent needlestick injuries.

- **Needlestick Injuries** should be reported immediately to the site supervisor, with appropriate care and follow-up given as directed by state/local guidelines. Safety needles or needle-free injection devices should be used if available to reduce the risk of injury.

- **Equipment Disposal** - Used needles should not be detached from syringes, recapped or cut before disposal. All used syringe/needle devices should be placed in puncture proof containers to prevent accidental needlesticks and reuse. Empty or expired vaccine vials are considered medical waste and
Appendix D

Vaccine Preparation - Proper vaccine handling and preparation is critical in maintaining the integrity of the vaccine during transfer from the manufacturer’s vial to the syringe and ultimately to the patient.

- Equipment Selection

  - Syringe Selection - A separate needle and syringe should be used for each injection. A parenteral vaccine may be delivered in either a 1-mL or 3-mL syringe as long as the prescribed dosage is delivered. Syringe devices with sharps engineered sharps injury protection are available, recommended by OSHA, and required in many states to reduce the incidence of needle stick injuries and potential disease transmission. Personnel should be involved in evaluation and selection of these products. Staff should receive training with these device before using them in the clinical area.

  - Needle Selection - Vaccine must reach the desired tissue site for optimal immune response. Therefore, needle selection should be based upon the prescribed route, size of the individual, volume and viscosity of the vaccine, and injection technique. (See Subcutaneous & Intramuscular Injections, below.) Typically, vaccines are not highly viscous, and therefore a fine gauge needle (22-25 gauge) can be used.

  - Needle-Free Injection - A new generation of needle-free vaccine delivery devices has been developed in an effort to decrease the risks of needlestick injuries to healthcare workers and to prevent improper reuse of syringes and needles. For more information on needle-free injection technology, see the CDC website: www.cdc.gov/od/science/iso/vaxtech/nfit/.

- Inspecting Vaccine - Each vaccine vial should be carefully inspected for damage or contamination prior to use. The expiration date printed on the vial or box should be checked. Vaccine can be used through the last day of the month indicated by the expiration date unless otherwise stated on the package labeling. Expired vaccine should never be used.

- Reconstitution - Some vaccines are prepared in a lyophilized form that requires reconstitution, which should be done according to manufacturer guidelines. Diluent solutions vary; use only the specific diluent supplied for the vaccine. Once reconstituted, the vaccine must be either administered within the time guidelines provided by the manufacturer or discarded. Changing the needle after reconstitution of the vaccine is not necessary unless the needle has become contaminated or bent. Continue with standard medication preparation guidelines.

- Prefilling Syringes - CDC strongly discourages filling syringes in advance, because of the increased risk of administration errors. Once the vaccine is in the syringe it is difficult to identify the type or brand of vaccine. Other problems associated with this practice are vaccine wastage, and possible bacterial growth in vaccines that do not contain a preservative. Furthermore, medication administration guidelines state that the individual who administers a medication should be the one to draw up and prepare it. An alternative to prefilling syringes is to use filled syringes supplied by the vaccine manufacturer. Syringes other than those filled by the manufacturer are designed for immediate
administration, not for vaccine storage.

In certain circumstances, such as a large influenza clinic, more than one syringe can be filled. One person should prefill only a few syringes at a time, and the same person should administer them. Any syringes left at the end of the clinic day should be discarded.

Under no circumstances should MMR, varicella, or zoster vaccines ever be reconstituted and drawn prior to the immediate need for them. These live virus vaccines are unstable and begin to deteriorate as soon as they are reconstituted with diluent.

- **Labeling** - Once a vaccine is drawn into a syringe, the content should be indicated on the syringe. There are a variety of methods for identifying or labeling syringes (e.g. keep syringes with the appropriate vaccine vials, place the syringes in a labeled partitioned tray, or use color coded labels or preprinted labels).
- **Subcutaneous** (Sub-Q or SC) injections are administered into the fatty tissue found below the dermis and above muscle tissue.

- **Site** - Subcutaneous tissue can be found all over the body. The usual sites for vaccine administration are the thigh (for infants <12 months of age) and the upper outer triceps of the arm (for persons ≥12 months of age). If necessary, the upper outer triceps area can be used to administer subcutaneous injections to infants.
- **Needle Gauge & Length** - 5/8-inch, 23- to 25-gauge needle

- **Technique**

  - Follow standard medication administration guidelines for site assessment/selection and site preparation.

  - To avoid reaching the muscle, pinch up the fatty tissue, insert the needle at a 45° angle and inject the vaccine into the tissue.

  - Withdraw the needle and apply light pressure to the injection site for several seconds with a dry cotton ball or gauze.

Subcutaneous Administration Techniques
• **Intramuscular** (IM) injections are administered into muscle tissue below the dermis and subcutaneous tissue.

- **Site** - Although there are several IM injection sites on the body, the recommended IM sites for vaccine administration are the vastus lateralis muscle (anterolateral thigh) and the deltoid muscle (upper arm). The site depends on the age of the individual and the degree of muscle development.
- **Needle Gauge** - 22- to 25-gauge needle

- **Needle Length** - For all intramuscular injections, the needle should be long enough to reach the muscle mass and prevent vaccine from seeping into subcutaneous tissue, but not so long as to involve underlying nerves, blood vessels, or bone. The vaccinator should be familiar with the anatomy of the area into which the vaccine will be injected.

Decision on needle size and site of injection must be made for each person on the basis of the size of the muscle, the thickness of adipose tissue at the injection site, the volume of the material to be administered, injection technique, and the depth below the muscle surface into which the material is to be injected.

- **Infants (Younger Than 12 Months)**
  For the majority of infants, the anterolateral aspect of the thigh is the recommended site for injection because it provides a large muscle mass. The muscles of the buttock have not been used for administration of vaccines in infants and children because of concern about potential injury to the sciatic nerve, which is well documented after injection of antimicrobial agents into the buttock. If the gluteal muscle must be used, care should be taken to define the anatomic landmarks.*

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*If the gluteal muscle is chosen, injection should be administered lateral and superior to a line between the posterior superior iliac spine and the greater trochanter or in the ventrogluteal site, the center of a triangle bounded by the anterior superior iliac spine, the tubercle of the iliac crest, and the upper border of the greater trochanter.
Injection technique is the most important factor to ensure efficient intramuscular vaccine delivery. If the subcutaneous and muscle tissue are bunched to minimize the chance of striking bone, a 1-inch needle is required to ensure intramuscular administration in infants. For the majority of infants, a 1-inch, 22-25-gauge needle is sufficient to penetrate muscle in an infant's thigh. For newborns (first 28 days of life) and premature infants, a 5/8 inch needle usually is adequate if the skin is stretched flat between thumb and forefinger and the needle inserted at a 90-degree angle to the skin.

- **Toddlers and Older Children (12 Months through 10 Years)**
  The deltoid muscle should be used if the muscle mass is adequate. The needle size for deltoid site injections can range from 22 to 25 gauge and from 5/8 to 1 inch on the basis of the size of the muscle and the thickness of adipose tissue at the injection site. A 5/8-inch needle is adequate only for the deltoid muscle and only if the skin is stretched flat between thumb and forefinger and the needle inserted at a 90° angle to the skin. For toddlers, the anterolateral thigh can be used, but the needle should be at least 1 inch in length.

- **Adolescents and Adults (11 Years or Older)**
  For adults and adolescents, the deltoid muscle is recommended for routine intramuscular vaccinations. The anterolateral thigh also can be used. For men and women weighing less than 130 lbs (60 kg) a 5/8-1-inch needle is sufficient to ensure intramuscular injection. For women weighing 130-200 lbs (60-90 kg) and men 130-260 lbs (60-118kg), a 1-1½-inch needle is needed. For women weighing more than 200 lbs (90 kg) or men weighing more than 260 lbs (118 kg), a 1½-inch needle is required.

- **Technique**
  - Follow standard medication administration guidelines for site assessment/selection and site preparation.
  - To avoid injection into subcutaneous tissue, spread the skin of the selected vaccine administration site taut between the thumb and forefinger, isolating the muscle. Another technique, acceptable mostly for pediatric and geriatric patients, is to grasp the tissue and "bunch up" the muscle.
  - Insert the needle fully into the muscle at a 90° angle and inject the vaccine into the tissue.
  - Withdraw the needle and apply light pressure to the injection site for several seconds with a dry cotton ball or gauze.

- **Aspiration** - Aspiration is the process of pulling back on the plunger of the syringe prior to injection to ensure that the medication is not injected into a blood vessel. Although this practice is advocated by some experts, the procedure is not required because no large blood vessels exist at the recommended injection sites.
• **Multiple Vaccinations** - When administering multiple vaccines, NEVER mix vaccines in the same syringe unless approved for mixing by the Food and Drug Administration (FDA). If more than one vaccine must be administered in the same limb, the injection sites should be separated by 1-2 inches so that any local reactions can be differentiated. Vaccine doses range from 0.2 mL to 1 mL. The recommended maximum volume of medication for an IM site, varies among references and depends on the muscle mass of the individual. However, administering two IM vaccines into the same muscle would not exceed any suggested volume ranges for either the vastus lateralis or the deltoid muscle in any age group. The option to also administer a subcutaneous vaccine into the same limb, if necessary, is acceptable since a different tissue site is involved.

If a vaccine and an immune globulin preparation are administered simultaneously (e.g., Td/Tdap and tetanus immune globulin [TIG] or hepatitis B vaccine and hepatitis B immune globulin [HBIG]), a separate anatomic site should be used for each injection. The location of each injection should be documented in the patient’s medical record.

• **Nonstandard Administration** - Deviation from the recommended route, site and dosage of vaccine is strongly discouraged and can result in inadequate protection. In situations where nonstandard administration has occurred, refer to the ACIP General Recommendation on Immunization, *MMWR* 2006; 55 (RR-15), for specific guidance.
Special Situations

Bleeding Disorders - Individuals with a bleeding disorder or who are receiving anticoagulant therapy may develop hematomas in IM injection sites. Prior to administration of IM vaccines the patient or family should be instructed about the risk of hematoma formation from the injection. Additionally, a physician familiar with the patient’s bleeding disorder or therapy should be consulted regarding the safety of administration by this route. If the patient periodically receives hemophilia replacement factor or other similar therapy, IM vaccine administration should ideally be scheduled shortly after replacement therapy. A 23-gauge or finer needle should be used and firm pressure applied to the site for at least 2 minutes. The site should not be rubbed or massaged.

Latex Allergy - Administration of a vaccine supplied in a vial or syringe that contains natural rubber (refer to product information) should not be administered to an individual with a history of a severe (anaphylactic) allergy to latex, unless the benefit of vaccination clearly outweighs the risk of an allergic reaction. These situations are rare. Medical consultation and direction should be sought regarding vaccination. A local or contact sensitivity to latex is not a contraindication to vaccination.

Syncopal or Vasovagal Response ("fainting") may occur during vaccine administration, especially with adolescents and adults. Because individuals may fall and sustain injury as a result, the provider should have the patient sit during injection(s). A syncopal or vasovagal response is not common and is not an allergic reaction. However, if syncope develops, the provider should observe and administer supportive care until the patient is recovered.

Anaphylaxis (a life-threatening acute allergic reaction) - Each facility that administers vaccines should have a protocol, procedures and equipment to provide initial care for suspected anaphylaxis. Facility staff should be prepared to recognize and respond appropriately to this type of emergency situation. All staff should maintain current CPR certification. Emergency protocols, procedures and equipment/supplies should be reviewed periodically. For additional information on medical management of vaccine reactions in children, teens, and adults, see the 2006 ACIP General Recommendations on Immunization (p. 19), the 2006 AAP Red Book (pp. 64-66), and pages D28-D31 of this appendix. Although both fainting and allergic reactions are rare, vaccine providers should strongly consider observing patients for 15 minutes after they are vaccinated.

Documentation

All vaccines administered should be fully documented in the patient’s permanent medical record. Documentation should include:

1. Date of administration
2. Name or common abbreviation of vaccine
3. Vaccine lot number
4. Vaccine manufacturer
5. Administration site

6. Vaccine Information Statement (VIS) edition date (found in the lower right corner of the back of the VIS).

7. Name and address of vaccine administrator. This should be the address where the record is kept. If immunizations are given in a shopping mall, for example, the address would be the clinic where the permanent record will reside.

Facilities that administer vaccines are encouraged to participate in state/local immunization information systems. The patient or parent should be provided with an immunization record that includes the vaccines administered with dates of administration.

The California Department of Health Services' Immunization Branch has developed a complete package of resources on vaccine administration, available at http://www.eziz.org/pages/vaccineadmin.html
The Skills Checklist is a self-assessment tool for health care staff who administer immunizations. To complete it, review the competency areas below and the clinical skills, techniques, and procedures outlined for each of them. Score yourself in the Self-Assessment column. If you check Need to Improve you indicate further study, practice or change is needed. When you check Meets or Exceeds you indicate you believe you are performing at the expected level of competence, or higher.

**Supervisors:** Use the Skills Checklist to clarify responsibilities and expectations for staff who administer vaccines. When you use it for performance reviews, give staff the opportunity to score themselves in advance. Next observe their performance as they provide immunizations to several patients and score in the Supervisor Review columns. If improvement is needed, meet with them to develop a Plan of Action (over) that will help them achieve the level of competence you expect; circle desired actions or write in others. In 30 days, observe their performance again. When all competency areas meet expectations, file the Skills Checklist in their personnel folder. At the end of the probationary period and annually thereafter, observe them again and complete the Skills Checklist.

<table>
<thead>
<tr>
<th>Competency</th>
<th>Clinical Skills, Techniques, and Procedures</th>
<th>Self-Assessment</th>
<th>Supervisor Review</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>A. Patient/Parent Education</strong></td>
<td>1. Welcomes patient/family; establishes rapport; and answers any questions.</td>
<td>Need to Improve</td>
<td>Meets or Exceeds</td>
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<td></td>
<td>2. Explains what vaccines will be given and which type(s) of injection will be done.</td>
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<td>3. Accommodates language or literacy barriers and special needs of patient/parents to help make them feel comfortable and informed about the procedure.</td>
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<td>4. Verifies patient/parents received the Vaccine Information Statements for indicated vaccines and had time to read them and ask questions.</td>
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<td>5. Screens for contraindications. (MA: score NA— not applicable— if this is MD function.)</td>
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<td></td>
<td>6. Reviews comfort measures and after care instructions with patient/parents, inviting questions.</td>
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<td><strong>B. Medical Protocols</strong></td>
<td>1. Identifies the location of the medical protocols (i.e., immunization protocol, emergency protocol, reference material).</td>
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<td>2. Identifies the location of the epinephrine, its administration technique, and clinical situations where its use would be indicated.</td>
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<td>3. Maintains up-to-date CPR certification.</td>
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<td></td>
<td>4. Understands the need to report any needlestick injury and to maintain a sharps injury log.</td>
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<td><strong>C. Vaccine Handling</strong></td>
<td>1. Checks vial expiration date. (Double-checks vial label and contents prior to drawing up.</td>
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<td>2. Maintains aseptic technique throughout.</td>
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<td></td>
<td>3. Selects the correct needle size: 1”- 1 1/2” for IM (DTaP, Td, Hib, HepA, HepB, Pneumo Conj., Poliw. 1/2” for SC (MMR, Var), IPV and Pneumo Poly depends on route to be used.</td>
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<td>4. Shakes vaccine vial and/or reconstitutes and mixes using the diluent supplied. Inverts vial and draws up correct dose of vaccine. Rechecks vial label.</td>
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<td>5. Labels each filled syringe or uses labeled tray to keep them identified.</td>
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<td>6. Demonstrates knowledge of proper vaccine handling, e.g. protects MMR from light, log refrigerator temperature.</td>
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<tr>
<td>Competency</td>
<td>Clinical Skills, Techniques, and Procedures</td>
<td>Self-Assessment</td>
<td>Supervisor Review</td>
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<tr>
<td>D. Administering Immunizations</td>
<td>1. Rechecks the physician’s order or instructions against prepared syringes.</td>
<td>Need to Improve</td>
<td>Need to Improve</td>
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<tr>
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<td>2. Washes hands and if office policy puts on disposable gloves.</td>
<td>Meets or Exceeds</td>
<td>Meets or Exceeds</td>
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<td>3. Demonstrates knowledge of the appropriate route for each vaccine. (M for DTaP, Td, Hb, HepA, HepB, Pneumo Conj, PCV for MMR, Var; Either SC or IM for IPV and Pneumo Poly).</td>
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<td>4. Positions patient and/or restrains the child with parent’s help; locates anatomic landmarks specific for IM or SC.</td>
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<td>5. Prep the site with an alcohol wipe using a circular motion from the center to a 2 to 3” circle. Allows alcohol to dry.</td>
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<td>6. Controls the limb with the non-dominant hand; holds the needle an inch from the skin and inserts it quickly at the appropriate angle (45° for SC or 90° for IM).</td>
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<td>7. Injects vaccine using steady pressure; withdraws needle at angle of insertion.</td>
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<td></td>
<td>8. Applies gentle pressure to injection site for several seconds with a dry cotton ball.</td>
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<td></td>
<td>10. Encourages comfort measures before, during and after the procedure.</td>
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<tr>
<td>E. Records Procedures</td>
<td>1. Fully documents each immunization in patient’s chart: date, lot number, manufacturer, site, VIS date, name/initials.</td>
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<td></td>
<td>2. If applicable, demonstrates ability to use IZ registry or computer to call up patient record, assess what is due today, and update computer immunization history.</td>
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<td>3. Asks for and updates patient’s record of immunizations and reminds them to bring it to each visit.</td>
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</tbody>
</table>

**Plan of Action:** Circle desired next steps and write in the agreed deadline and date for the follow-up performance review.  

- a. Watch video on immunization techniques.  
- b. Review office protocols.  
- c. Review manuals, textbooks, wall charts or other guides.  
- d. Review package inserts.  
- e. Review vaccine handling guidelines or video.  
- f. Observe other staff with patients.  
- g. Practice injections.  
- h. Read Vaccine Information Statements.  
- i. Be mentored by someone who has these skills.  
- j. Role play with other staff interactions with parents and patients, including age-appropriate comfort measures.  
- k. Attend a skills training or other courses or training.  
- l. Attend health care customer satisfaction or cultural competency training.  
- m. Renew CPR certification.  

**Other:**

Employee Signature

Date

Plan of Action Deadline

Supervisor Signature

Date

Data of Next Performance Review
Giving All the Doses Under 12 Months

- Needle Lengths:
  IM = 1 inch  SC = 5/8 inch

- Using combination vaccines will decrease the number of injections

- IM injections are given in the infant’s thigh

- SC injections may be given in the arm or thigh

- Separate injection sites by 1-2 inches

- May consider a 5/8” needle for IM injections only in newborns less than 4 wks

Giving All the Doses 12 Months and Older

- Needle Lengths
  IM = 1 to 1.5 inches
  SC = 5/8 inch

- Separate injection sites by 1-2 inches

- Anterolateral thigh is the preferred site for multiple IM injections

- Deltoid (upper arm) is an option for IM in children ≥ 18 mo with adequate muscle mass

- Using combination vaccines will decrease the number of injections needed to keep a child up-to-date
Appendix D

GIVING ALL THE DOSES
11-12 Years of Age

- Needle Lengths
  IM = 1 to 1.5 in
  SC = 5/8 in

- Separate injection sites by 1-2 inches

- Professional judgment is appropriate when selecting needle length for use in all children, especially small infants or larger children.

- Assess for other recommended vaccines that may be needed:
  MMR  Polio
  hep B  Hep A
  influenza

- Syncope or fainting after vaccination may occur in adolescents & young adults, usually within 15 minutes of vaccination

- When giving vaccines to teens:
  Have the patient sit down while you are giving vaccine(s)
  Consider observing patients for 15-20 minutes after vaccination

NOTE:
Var should be administered to school age children and adolescents without:
- history of 2 doses of varicella vaccine
- a healthcare provider’s diagnosis of varicella disease or
  verification of history of typical varicella disease
- history of shingles

HPV4 is licensed for use in girls only 9-26 years of age
MMRV (ProQuad®) is licensed for children 12 months thru 12 years of age only

Alliance for Immunization in Michigan
2009 AIM Kit—Adolescent Immunization Section

December 6, 2007
COMFORTING RESTRAINT

FOR IMMUNIZATIONS

• The method:

This method involves the parent in embracing the child and controlling all four limbs. It avoids “holding down” or overpowering the child, but it helps you steady and control the limb of the injection site.

• For infants and toddlers:

Have parent hold the child on parent’s lap.

1. One of the child’s arms embraces the parent’s back and is held under the parent’s arm.

2. The other arm is controlled by the parent’s arm and hand. For infants, the parent can control both arms with one hand.

3. Both legs are anchored with the child’s feet held firmly between the parent’s thighs, and controlled by the parent’s other arm.

• For kindergarten and older children:

Hold the child on parent’s lap or have the child stand in front of the seated parent.

1. Parent’s arms embrace the child during the process.

2. Both legs are firmly between parent’s legs.

IMMUNIZATION TECHNIQUES
Safe Effective Caring
## Medical Management of Vaccine Reactions in Children and Teens

All vaccines have the potential to cause an adverse reaction. To minimize adverse reactions, patients should be carefully screened for precautions and contraindications before vaccine is administered. Even with careful screening, reactions can occur. These reactions can vary from trivial and inconvenient (e.g., soreness, itching) to severe and life threatening (e.g., anaphylaxis). If reactions occur, staff should be prepared with procedures for their management. The table below describes procedures to follow if various reactions occur.

<table>
<thead>
<tr>
<th>Reaction</th>
<th>Symptoms</th>
<th>Management</th>
</tr>
</thead>
<tbody>
<tr>
<td>Localized</td>
<td>Soreness, redness, itching, or swelling at the injection site</td>
<td>Apply a cold compress to the injection site. Consider giving an analgesic (pain reliever) or antipruritic (anti-itch) medication.</td>
</tr>
<tr>
<td></td>
<td>Slight bleeding</td>
<td>Apply an adhesive compress over the injection site.</td>
</tr>
<tr>
<td></td>
<td>Continuous bleeding</td>
<td>Place thick layer of gauze pads over site and maintain direct and firm pressure; raise the bleeding injection site (e.g., arm) above the level of the patient’s heart.</td>
</tr>
<tr>
<td>Psychological fright and syncopate</td>
<td>Fright before injection is given</td>
<td>Have patient sit or lie down for the vaccination.</td>
</tr>
<tr>
<td>(fainting)</td>
<td>Extreme paleness, sweating, coldness of the hands and feet, nausea, light-headedness, dizziness, weakness, or visual disturbances</td>
<td>Have patient lie flat or sit with head between knees for several minutes. Loosen any tight clothing and maintain an open airway. Apply cool, damp cloths to patient’s face and neck.</td>
</tr>
<tr>
<td></td>
<td>Fall, without loss of consciousness</td>
<td>Examine the patient to determine if injury is present before attempting to move the patient. Place patient flat on back with feet elevated.</td>
</tr>
<tr>
<td></td>
<td>Loss of consciousness</td>
<td>Check the patient to determine if injury is present before attempting to move the patient. Place patient flat on back with feet elevated. Call 911 if patient does not recover immediately.</td>
</tr>
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<td>Anaphylaxis</td>
<td>Sudden or gradual onset of generalized itching, erythema (redness), or urticaria (hives); angioedema (swelling of the lips, face, or throat); severe bronchospasm (wheezing); shortness of breath; shock; abdominal cramping; or cardiovascular collapse</td>
<td>See “Emergency Medical Protocol for Management of Anaphylactic Reactions in Children and Teens” on the next page for detailed steps to follow in treating anaphylaxis.</td>
</tr>
</tbody>
</table>

### Supplies Needed
- Aqueous epinephrine 1:1000 dilution, in ampules, vials of solution, or prefilled syringes, including epinephrine auto-injectors (e.g., EpiPen). If EpiPens are to be stocked, both EpiPen Jr. (0.15 mg) and adult EpiPens (0.30 mg) should be available.
- Diphenhydramine (Benadryl) injectable (50 mg/mL solution) and oral (12.5 mg/5 mL suspension) and 25 mg or 50 mg capsules or tablets.
- Syringes: 1-3 cc, 22-25g, 1", 1½", and 2" needles for epinephrine and diphenhydramine (Benadryl)
- Pediatric & adult airways (small, medium, and large)
- Sphygmomanometer (child, adult & extra-large cuffs) and stethoscope
- Pediatric & adult size pocket masks with one-way valve
- Alcohol swabs
- Tongue depressors
- Flashlight with extra batteries (for examination of mouth and throat)
- Wrist watch
- Tourniquet
- Cell phone or access to an on-site phone
Emergency Medical Protocol for Management of Anaphylactic Reactions in Children and Teens

**Signs and Symptoms of Anaphylactic Reaction**
Sudden or gradual onset of generalized itching, erythema (redness), or urticaria (hives); angioedema (swelling of the lips, face, or throat); bronchospasm (wheezing); shortness of breath; shock; abdominal cramping; or cardiovascular collapse.

**Treatment in Children and Teens**

a. If itching and swelling are confined to the injection site where the vaccination was given, observe patient closely for the development of generalized symptoms.

b. If symptoms are generalized, activate the emergency medical system (EMS; e.g., call 911) and notify the on-call physician. This should be done by a second person, while the primary nurse assesses the airway, breathing, circulation, and level of consciousness of the patient.

c. Administer aqueous epinephrine 1:1000 dilution (i.e., 1 mg/mL) intramuscularly; the standard dose is 0.01 mg/kg body weight, up to 0.3 mg maximum single dose in children and 0.5 mg maximum in adolescents (see chart below).

d. In addition, for anaphylaxis, administer diphenhydramine either orally or by intramuscular injection; the standard dose is 1 mg/kg body weight, up to 30 mg maximum dose in children and 100 mg maximum dose in adolescents (see chart below).

e. Monitor the patient closely until EMS arrives. Perform cardiopulmonary resuscitation (CPR), if necessary, and maintain airway. Keep patient in supine position (flat on back) unless he or she is having breathing difficulty. If breathing is difficult, patient’s head may be elevated, provided blood pressure is adequate to prevent loss of consciousness. If blood pressure is low, elevate legs. Monitor blood pressure and pulse every 5 minutes.

f. If EMS has not arrived and symptoms are still present, repeat dose of epinephrine every 10–20 minutes for up to 3 doses, depending on patient’s response.

g. Record all vital signs, medications administered to the patient, including the time, dosage, response, and the name of the medical personnel who administered the medication, and other relevant clinical information.

h. Notify the patient’s primary care physician.

**Suggested Dosing of Epinephrine and Diphenhydramine**

<table>
<thead>
<tr>
<th>Age Group</th>
<th>Weight * in kg</th>
<th>Weight (lbs)* in lbs</th>
<th>Epinephrine Dose 1 mg/mL injectable (1:1000 dilution) intramuscular</th>
<th>Diphenhydramine (Benadryl) 12.5 mg/5 mL liquid 25 and 50 mg capsules or tabs 50 mg/mL injectable</th>
</tr>
</thead>
<tbody>
<tr>
<td>1–6 mos</td>
<td>4–7 kg</td>
<td>9–15 lbs</td>
<td>0.05 mg (0.05 mL)</td>
<td>5 mg</td>
</tr>
<tr>
<td>7–18 mos</td>
<td>7–11 kg</td>
<td>15–24 lbs</td>
<td>0.1 mg (0.1 mL)</td>
<td>10 mg</td>
</tr>
<tr>
<td>19–36 mos</td>
<td>11–14 kg</td>
<td>24–31 lbs</td>
<td>0.15 mg (0.15 mL)</td>
<td>15 mg</td>
</tr>
<tr>
<td>37–48 mos</td>
<td>14–17 kg</td>
<td>31–37 lbs</td>
<td>0.15 mg (0.15 mL)</td>
<td>20 mg</td>
</tr>
<tr>
<td>49–59 mos</td>
<td>17–19 kg</td>
<td>37–42 lbs</td>
<td>0.2 mg (0.2 mL)</td>
<td>30 mg</td>
</tr>
<tr>
<td>5–7 yrs</td>
<td>19–23 kg</td>
<td>42–51 lbs</td>
<td>0.2 mg (0.2 mL)</td>
<td>40 mg</td>
</tr>
<tr>
<td>8–10 yrs</td>
<td>23–35 kg</td>
<td>51–77 lbs</td>
<td>0.3 mg (0.3 mL)</td>
<td></td>
</tr>
<tr>
<td>11–12 yrs</td>
<td>35–45 kg</td>
<td>77–99 lbs</td>
<td>0.4 mg (0.4 mL)</td>
<td></td>
</tr>
<tr>
<td>13 yrs &amp; older</td>
<td>45 kg</td>
<td>99 lbs</td>
<td>0.5 mg (0.5 ml)</td>
<td>50–100 mg</td>
</tr>
</tbody>
</table>

*Dosage by body weight is preferred.

These standing orders for the medical management of vaccine reactions in child and teenage patients shall remain in effect for patients of the ____________________________ until rescinded or until ________________ date ________________.

Medical Director’s signature ____________________________ Effective date ____________________________

Sources:
- American Pharmacists Association, Glueckstein, JD. Pharmacy-Based Immunization Delivery, 2002.

(www.immunize.org/csg/pdf/0802a.pdf • item #5082a (6/06))

Immunization Action Coalition  •  1573 Selby Ave.  •  St. Paul, MN 55104  •  (651) 647-9009  •  www.immunize.org  •  www.vaccineinformation.org
Medical Management of Vaccine Reactions in Adult Patients

All vaccines have the potential to cause an adverse reaction. In order to minimize adverse reactions, patients should be carefully screened for precautions and contraindications before vaccine is administered. Even with careful screening, reactions may occur. These reactions can vary from trivial and inconvenient (e.g., soreness, itching) to severe and life threatening (e.g., anaphylaxis). If reactions occur, staff should be prepared with procedures for their management. The table below describes procedures to follow if various reactions occur.

<table>
<thead>
<tr>
<th>Reaction</th>
<th>Symptoms</th>
<th>Management</th>
</tr>
</thead>
<tbody>
<tr>
<td>Localized</td>
<td>Soreness, redness, itching, or swelling at the injection site</td>
<td>Apply a cold compress to the injection site. Consider giving an analgesic (pain reliever) or antipruritic (anti-itch) medication.</td>
</tr>
<tr>
<td></td>
<td>Slight bleeding</td>
<td>Apply an adhesive compress over the injection site.</td>
</tr>
<tr>
<td></td>
<td>Continuous bleeding</td>
<td>Place thick layer of gauze pads over site and maintain direct and firm pressure; raise the bleeding injection site (e.g., arm) above the level of the patient’s heart.</td>
</tr>
<tr>
<td>Psychological fright and syncope (fainting)</td>
<td>Fright before injection is given</td>
<td>Have patient sit or lie down for the vaccination.</td>
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<tr>
<td></td>
<td>Extreme paleness, sweating, coldness of the hands and feet, nausea, light-headedness, dizziness, weakness, or visual disturbances</td>
<td>Have patient lie flat or sit with head between knees for several minutes. Loosen any tight clothing and maintain an open airway. Apply cool, damp cloths to patient’s face and neck.</td>
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<td>Sudden or gradual onset of generalized itching, erythema (redness), or urticaria (hives); angioedema (swelling of the lips, face, or throat); severe bronchospasm (wheezing); shortness of breath; shock; abdominal cramping; or cardiovascular collapse.</td>
<td>See “Emergency Medical Protocol for Management of Anaphylactic Reactions in Adults” on the next page for detailed steps to follow in treating anaphylaxis.</td>
</tr>
</tbody>
</table>

(continued on page 2)
Emergency Medical Protocol for Management of Anaphylactic Reactions in Adults

Supplies Needed

- Aqueous epinephrine 1:1000 (i.e., 1 mg/mL) dilution, in ampules, vials of solution, or prefilled syringes, including epinephrine autoinjectors (e.g., EpiPen). If EpiPens are stocked, at least three adult EpiPens (0.30 mg) should be available.
- Diphenhydramine (Benadryl) injectable (50 mg/mL solution) and 25 mg or 50 mg capsules or tablets and syrup (12.5 mg/5 mL suspension)
- Syringes: 1-3 cc, 22-25g, 1", 1½", and 2" needles for epinephrine and diphenhydramine (Benadryl)
- Wristwatch with second hand
- Adult airways (small, medium, and large)
- Sphygmomanometer (adult and extra-large cuffs) and stethoscope
- Adult size pocket mask with one-way valve
- Alcohol swabs
- Tourniquet
- Tongue depressors
- Flashlight with extra batteries (for examination of the mouth and throat)
- Cell phone or access to an on-site phone

Signs and Symptoms of Anaphylactic Reaction

Sudden or gradual onset of generalized itching, erythema (redness), or urticaria (hives); angioedema (swelling of the lips, face, or throat); bronchospasm (wheezing); shortness of breath; shock; abdominal cramping; or cardiovascular collapse.

Treatment in Adults

a. If itching and swelling are confined to the injection site where the vaccination was given, observe patient closely for the development of generalized symptoms.

b. If symptoms are generalized, activate the emergency medical system (EMS; e.g., call 911) and notify the on-call physician. This should be done by a second person, while the primary nurse assesses the airway, breathing, circulation, and level of consciousness of the patient.

c. Administer aqueous epinephrine 1:1000 dilution intramuscularly, 0.01 mL/kg/dose (adult dose ranges from 0.3 mL to 0.5 mL, with maximum single dose of 0.5 mL).

d. In addition, for systemic anaphylaxis, administer diphenhydramine either orally or by intramuscular injection; the standard dose is 1–2 mg/kg, up to 100 mg maximum single dose.

e. Monitor the patient closely until EMS arrives. Perform cardiopulmonary resuscitation (CPR), if necessary, and maintain airway. Keep patient in supine position (flat on back) unless he or she is having breathing difficulty. If breathing is difficult, patient’s head may be elevated, provided blood pressure is adequate to prevent loss of consciousness. If blood pressure is low, elevate legs. Monitor blood pressure and pulse every 5 minutes.

f. If EMS has not arrived and symptoms are still present, repeat dose of epinephrine every 10–20 minutes for up to 3 doses, depending on patient’s response.

g. Record all vital signs, medications administered to the patient, including the time, dosage, response, and the name of the medical personnel who administered the medication, and other relevant clinical information.

h. Notify the patient’s primary care physician.

Sources:

These standing orders for the medical management of vaccine reactions in adult patients shall remain in effect for patients of the ____________________________ until rescinded or until ______________.

name of clinic
date

Medical Director’s signature

Effective date

Immunization Action Coalition • www.immunize.org • www.vaccineinformation.org • www.immunize.org/catg.d/p31082.pdf