### Purpose
- To reduce morbidity and mortality from coronavirus disease 2019 (COVID-19) by vaccinating persons who meet the criteria established by the Centers for Disease Control and Prevention’s Advisory Committee on Immunization Practices (ACIP).

### Policy
- Where authorized under state law, standing orders enable eligible nurses and other healthcare professionals (e.g., pharmacists) to assess and vaccinate persons who meet the criteria in the "Procedure" section below without the need for clinician examination or direct order from the attending provider at the time of the interaction.

### Procedure
Assess persons 12 years of age and older for vaccination with Pfizer-BioNTech COVID-19 Vaccine based on the following criteria:

Persons who **ARE NOT** moderately or severely immunocompromised:
- If the recipient has never received a COVID-19 vaccine, administer 1 dose of Pfizer-BioNTech COVID-19 Vaccine.
- If the recipient has received 1 previous dose of:
  - Pfizer-BioNTech COVID-19 Vaccine, administer the second dose at least 3 to 8 weeks after the first dose.
  - If the first-dose vaccine product cannot be determined or is no longer available, administer Pfizer-BioNTech COVID-19 Vaccine at least 3 to 8 weeks after the first dose.
- While a 3-week interval remains optimal for moderately to severely immunocompromised persons, adults ages 65 years and older, and others who need rapid protection because of community transmission or risk of disease, an 8-week interval may be optimal for some people, including males 12-39 years of age because of the small risk of myocarditis associated with mRNA COVID-19 vaccines. Vaccine effectiveness may also be increased with an interval longer than 3 weeks.
- If the recipient has received 2 previous doses of Pfizer-BioNTech COVID-19 Vaccine, administer a booster dose at least 5 months after dose 2.
- If the recipient has received 3 previous doses of Pfizer-BioNTech COVID-19 Vaccine (2 primary series doses and a booster), a second booster dose may be administered to persons 50 years of age or older at least 4 months after the most recent dose.
- If the recipient has received 1 dose of Janssen COVID-19 Vaccine, administer a booster dose at least 2 months after the single primary series dose (mRNA preferred).

### Additional Clinical Considerations
- Persons who have received HCT or CAR-T-cell therapy:
  - Revaccinate persons who received doses of COVID-19 vaccine prior to receiving or during HCT or CAR-T-cell therapy with a primary series at least 3 months (12 weeks) after transplant or CAR-T-cell therapy.
  - For persons who received a COVID-19 vaccine:
    - Outside of the United States

### Table: Vaccine Diluent Dosage (amount)/ Route

<table>
<thead>
<tr>
<th>Vaccine</th>
<th>Diluent</th>
<th>Dosage (amount)/ Route</th>
</tr>
</thead>
<tbody>
<tr>
<td>12 years of age and older (purple cap)</td>
<td>1.8 mL of 0.9% sodium chloride (normal saline, preservative-free) diluent</td>
<td>0.3 mL/IM injection</td>
</tr>
</tbody>
</table>

*Inform recipients, especially males 12 through 29 years of age and their parents/legal representative (when relevant) of the possibility of myocarditis or pericarditis following receipt of mRNA COVID-19 vaccines and the need to seek care if symptoms of myocarditis or pericarditis develop after vaccination. Educational materials are available at [https://www.cdc.gov/coronavirus/2019-ncov/vaccines/safety/myocarditis.html](https://www.cdc.gov/coronavirus/2019-ncov/vaccines/safety/myocarditis.html)

- Pfizer-BioNTech COVID-19 Vaccine may be coadministered with other vaccines without regard to timing, including simultaneous administration.
- For recommendations for COVID-19 vaccination and SARS-CoV-2 infection guidance, including after receiving passive antibody products, can be found at: [https://www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html#CoV-19-vaccination](https://www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html#CoV-19-vaccination)

### Screen for Contraindications and Precautions

- **Contraindications:**
  - History of a:
    - Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a component of the COVID-19 vaccine
    - Known diagnosed allergy to a component of the vaccine (see [https://www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html#Appendix-C](https://www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html#Appendix-C) for a list of vaccine components)

### Needle Gauge/Length/Injection Site

<table>
<thead>
<tr>
<th>Sex and Weight of Patient</th>
<th>Needle Gauge/Length</th>
<th>Injection Site $^5$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Female or male fewer than 130 lbs</td>
<td>22-25 ½”–1”</td>
<td>Deltoid muscle of arm</td>
</tr>
<tr>
<td>Female or male 130–152 lbs</td>
<td>22-25 1”</td>
<td>Deltoid muscle of arm</td>
</tr>
<tr>
<td>Female 152–200 lbs</td>
<td>22-25 1–1½”</td>
<td>Deltoid muscle of arm</td>
</tr>
<tr>
<td>Male 152–260 lbs</td>
<td>22-25 1–1½”</td>
<td>Deltoid muscle of arm</td>
</tr>
<tr>
<td>Female 200+ lbs</td>
<td>22-25 1½”</td>
<td>Deltoid muscle of arm</td>
</tr>
<tr>
<td>Male 260+ lbs</td>
<td>22-25 1½”</td>
<td>Deltoid muscle of arm</td>
</tr>
</tbody>
</table>

- Provide all recipients and/or parents/legal guardians with a copy of the current Fact Sheet for Recipients and Caregivers.
- Prepare to administer the vaccine. Choose the correct needle gauge, needle length, and injection site for persons:
  - 12 through 18 years of age:
    - Needle gauge/length: 22-25 gauge, 1-inch
    - Site: Deltoid muscle of arm.
  - 19 years of age and older: See chart.
  - Mix Pfizer-BioNTech COVID-19 Vaccine with 0.9% sodium chloride (normal saline, preservative-free) diluent according to the manufacturer’s instructions. Follow manufacturer’s guidance for storing/handling mixed vaccine.
- Administer 0.3 mL Pfizer-BioNTech COVID-19 Vaccine by intramuscular (IM) injection.
- Document vaccination.
- COVID-19 vaccination providers must document vaccine administration in their medical record systems within 24 hours of administration and use their best efforts to report administration data to the relevant system (e.g., immunization information system) for the jurisdiction as soon as practicable and no later than 72 hours after administration.

$^1$ An immediate allergic reaction is defined as any hypersensitivity-related signs or symptoms such as urticaria, angioedema, respiratory distress (e.g., wheezing, stridor), or anaphylaxis that occur within 4 hours following exposure to a vaccine or medication.

$^3$ Consider consultation with an allergist-immunologist to help determine if the patient can safely receive vaccination. Healthcare providers and health departments may also request a consultation from the Clinical Immunization Safety Assessment COVIDvax Project ([https://www.cdc.gov/vaccinesafety/ensuringsafety/monitoring/cisa/index.html](https://www.cdc.gov/vaccinesafety/ensuringsafety/monitoring/cisa/index.html)). Vaccination of these individuals should only be done in an appropriate setting under the supervision of a healthcare provider experienced in the management of severe allergic reactions.

- People with a contraindication to mRNA COVID-19 vaccines (including due to a known PEG allergy) have a precaution to Janssen COVID-19 vaccination. People who have received an mRNA COVID-19 vaccine dose should wait at least 28 days to receive Janssen COVID-19 Vaccine.

$^4$ Alternately, the anterolateral thigh can be used. A 1.5-inch needle may be used if administering vaccine in this site.

$^5$ Some experts recommend a 5/8-inch needle for men and women who weigh less than 130 pounds. If used, skin must be stretched tightly (do not bunch subcutaneous tissue).
Pfizer-BioNTech COVID-19 Vaccine
Standing Orders for Administering Vaccine

- Document each recipient's vaccine administration information:
  » Medical record: The vaccine and the date it was administered, manufacturer, lot number, vaccination site and route, name and title of the person administering the vaccine.
  » Vaccination record card: Date of vaccination, product name/manufacturer, lot number, and name/location of the administering clinic or healthcare professional. Give to the vaccine recipient.
  » Immunization information system (IIS): Report the vaccination to the appropriate state/local IIS.

- Additional preparation and administration information is available on the manufacturer's website at www.cvdvaccine.com.

- Be prepared to manage medical emergencies.
  - Vaccination providers should observe patients after vaccination to monitor for the occurrence of immediate adverse reactions, including syncope:
    » 30 minutes: persons with a history of:
      • A contraindication to another type of COVID-19 vaccine product.
      • Immediate (within 4 hours of exposure) non-severe allergic reaction to a COVID-19 vaccine.
      • Immediate allergic reaction of any severity to a non-COVID-19 vaccine or injectable therapies.
      • Anaphylaxis due to any cause.
    » 15 minutes: All other persons
  - Syncope may occur in association with injectable vaccines, in particular among adolescents. Procedures should be in place to avoid falling injuries and manage syncopal reactions.
  - Have a written protocol to manage medical emergencies following vaccination, as well as equipment and medications, including at least 3 doses of epinephrine, H1 antihistamine, blood pressure monitor, and timing device to assess pulse.
  - Healthcare personnel who are trained and qualified to recognize the signs and symptoms of anaphylaxis as well as administer intramuscular epinephrine should be available at the vaccination location at all times.

- For more information, please see:
  » CDC's General Best Practice Guidelines for Immunization, “Preventing and Managing Adverse Reactions,” at https://www.cdc.gov/vaccines/hcp/acip-recs/general-recs/adverse-reactions.html
  » Immunization Action Coalition's “Medical Management of Vaccine Reactions in Adults in a Community Setting” at https://www.immunize.org/catg.d/p3082.pdf

- Report adverse events to the Vaccine Adverse Event Reporting System (VAERS).
  - While this vaccine is under Emergency Use Authorization (EUA) (https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization), healthcare professionals are required to report to VAERS:
    » Vaccine administration errors (whether associated with an adverse event [AE] or not)
    » Serious AEs (irrespective of attribution to vaccination)
    » Cases of COVID-19 that result in hospitalization or death
    » Any additional AEs and revised safety requirements per the Food and Drug Administration's (https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization) conditions for use of an authorized vaccine throughout the duration of the EUA
  - Healthcare professionals are encouraged to report to VAERS (https://vaers.hhs.gov):
    » Clinically important adverse events that occur after vaccination, even if you are not sure whether the vaccine caused the adverse event

Note: For more information/guidance, please contact the immunization program at your state or local health department or the appropriate state body (e.g., state board of medical/nursing/pharmacy practice).

Standing Orders Authorization

This policy and procedure shall remain in effect for all patients of the ________________________________ effective___________________ until rescinded or until ________________.

Medical director (or other authorized practitioner) ________________________________ / ________________________________ / ________________________________.

Adapted with appreciation from the Immunization Action Coalition (IAC) standing orders