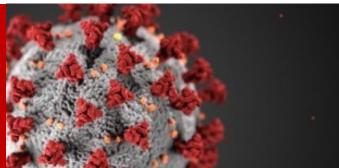
Steps to Take Before Moderna COVID-19 Vaccine Administration



■ 1. REGISTER/CHECK IN PATIENT ON ZOTEC (PHASE 1A) OR ACCENTURE

- Zotec (Phase 1A) or Accenture is used to register patient's appointment and vaccination record documentation
- Patient search must be completed in Zotec (Phase 1A) or Accenture prior to COVID-19 vaccine administration. The data will automatically be entered into CHIRP, our state immunization registry.
- Patient search in Zotec (Phase 1A) or Accenture is critical for second dose patients to ensure the same COVID-19 vaccine product is used for the second dose. Please refer to the Registering and Checking In the Patient section or www.in.gov/isdh/28690.htm for more information.



2. PROVIDE RECIPIENT EUA FACT SHEET

- Required under the National Childhood Vaccine Injury Act
- Must be given prior to administration of each dose of the vaccine
- Must provide the most current version. Please check the manufacturer website for the most recent version. The Moderna Provider and Recipient Fact Sheets can be found on www.modernatx.com.
- This also serves as an opportunity to educate the patient and address any questions or concerns patients may have. Refer to the **Patient** Education section or www.modernatx.com for more information.



3. SCREENING THE PATIENT

- The key to preventing the majority of serious adverse reactions is through patient screening
- Every person who administers the COVID-19 Vaccine should screen every patient for contraindications, previous allergies, and precautions prior to administering the vaccine dose
- The contraindications can be found on the Provider Fact Sheet. The Moderna COVID-19 Vaccine website is located at www.modernatx.com.
- Refer to the **Screening the Patient section** for more guidance.



Steps to Take During Moderna COVID-19 Vaccine Administration



1. CHECKING VIAL EXPIRATION DATE

- You can confirm the vial expiration date by looking up the lot number at https://www.modernatx.com/covid19vaccine-eua/providers/vial-lookup
- Refer to the Preparing and Thawing the Vaccine section for more information or visit www.modernatx.com



2. VACCINE PREPARATION (NO DILUTION REQUIRED)

- The Moderna COVID-19 Vaccine does not require dilution.
- Thaw vaccine at room temperature or in refrigerated conditions
- After thawing, let vial stand at room temperature for 15 minutes before administering. After thawing, do not refreeze the vial(s)
- Record the date and time for first use on the Moderna COVID-19
 Vaccine vial label. Discard vial after 6 hours
- Refer to the Preparing and Thawing the Vaccine section for more information on thawing and room temperature exposure time restrictions or www.modernatx.com



3. ADMINISTER INTRAMUSCULAR ROUTE

- Make sure staff are wearing appropriate PPE.
- Practice hand hygiene before administration, between patients, and when changing gloves (if worn), and any time your hands/gloves are soiled.
- Administer a single 0.5mL dose at first visit.
- Use appropriate gauge needle for body type (23-25 gauge).
- Administer intramuscularly in the deltoid muscle.
- Refer to the Administering the Vaccine section or www.modernatx.com for more information.



Image source: Intramuscular Injection Sites. Healthline. Medically reviewed by Deborah Weatherspoon, Ph.D., R.N., CRNA. Written by Jacquelyn Cafasso. Updated on March 7, 2019. Retrieved from https://www.healthline.com/health/intramuscular-injection

Steps to Take After Moderna COVID-19 Vaccine Administration



1. WAIT 15 MIN. & PROVIDE COVID-19 VACCINATION RECORD CARD

- Discard all used materials in appropriate waste receptacles.
- Monitor your patient after the vaccination for 15 minutes in a designated area to monitor for potential vaccine reaction.
- Using the COVID-19 vaccination record card provided in the ancillary kit, please record the time and day for second-dose appointment.
- The second dose is administered 21 days later and it must be <u>Moderna</u>
 COVID-19 vaccine; it is not interchangeable with other COVID-19
 vaccines.
- Refer to the **Closing the Loop section** for more information.



2. CLOSING THE LOOP

- V-safe is a new smartphone-based, after-vaccination health checker for people who receive COVID-19 Vaccines. Facilities are required to provide information on the V-safe program to vaccinated individuals and counsel them on the importance of enrolling.
- Vaccine Adverse Event Reporting System (VAERS) (www.vaers.hhs.gov) is an early warning system, co-managed by CDC and FDA, which monitors for potential vaccine safety problems; anyone can report possible vaccine side effects to VAERS. Click here for an informational video on VAERS.
- Refer to the **Closing the Loop section** for more information.



3. DOCUMENT VACCINATION IN ZOTEC (PHASE 1A) OR ACCENTURE

- Each COVID-19 dose administered must be entered into Zotec (Phase 1A) or Accenture at the time of vaccination.
- Vaccinations must be reported within 24 hours of administration.
- Refer to the Documenting the Vaccination Visit section or www.in.gov/

 isdh/28690.htm for more guidance.

