CLINICAL PROCEDURE EMS COVID19 VACCINE ADMINISTRATION PFIZER mRNA BNT162b2

Indications

This medicinal product has been given Emergency Use Authorization by the FDA for active immunization in individuals 16 years of age and older to prevent COVID-19 caused by SARS-CoV-2 virus

Contraindications - Mission Specific

- Age < 16 years
- Current Illness (Current Infection)
- Hx of severe allergic reaction to a previous dose of this vaccine or any vaccine ingredients
- Current pregnancy or chance of becoming pregnant (Refer patient to their PMD)
- Breastfeeding (Refer patient to their PMD)
- Any other vaccination within the last 14 days (eg influenza etc) (Have patient return after the 14 day window)
- Testing positive for COVID-19 in the last 2 weeks
- Any of the following symptoms in the last 10 days: fever (>100.4F), chills, cough, shortness of breath, difficulty breathing, fatigue, muscle or body aches, headache, new altered sense of taste or smell, sore throat, congestion or runny nose, nausea, vomiting, or diarrhea

Cautions

- History of severe allergies or reactions to any medications, foods, vaccines, or latex → Monitor closely after administration (30 minutes minimum)
- Immunocompromised or on a medication that affects the immune system → Inform patient vaccine might not provide as strong an immune protection
- Bleeding disorder or taking blood thinners \rightarrow Risk of hematoma at injection site
- Has received a first dose of another COVID-19 Vaccine → Ensure same manufacturer as previous dose

Procedure

- Prepare patient and supplies:
 - Ensure appropriate monitoring equipment and treatment supplies are available to manage any adverse reactions (e.g. Anaphylaxis)
 - o Ensure correct patient identification
 - Verify "Covid-19 Screening and Consent Form" has been completed
 - Ensure "Notice of Privacy Practices" and "EUA Fact Sheet for Recipients and Caregivers" have been provided
 - Re-confirm patient meets indications and has no contraindications
- Thaw and prepare dose (if not already done):





<u>Administer Vaccine Dose:</u>



Monitor for adverse reactions (e.g. anaphylaxis) for <u>minimum 15 minutes</u> and initiate immediate treatment (below) as needed

• If signs of severe allergic reaction/anaphylaxis (dyspnea, stridor, severe urticaria, tachycardia, hypotension, or Altered Mental Status) activate emergency response system and initiate treatment if available:

- Epinephrine 0.3 mg (1mg/mL concentration) intramuscular (may use epinephrine auto-injector if available)
- Perform Airway Management as required per local EMS protocols
- Establish IV/IO access and initiate cardiac monitoring
- Diphenhydramine 50 mg IV/IO or intramuscular
- Methylprednisolone sodium succinate 125 mg IV/IO
- Albuterol 2.5 mg nebulized if wheezing/dyspnea, may repeat x 1
- Obtain 12-lead ECG after any epinephrine administration
- Initiate transport per local EMS protocols
- Consult OLMC for additional epinephrine/epinephrine drip as needed
- Report any adverse reactions

<u>Documentation</u>: Use provided forms to document vaccine manufacturer, injection site, lot number and expiration date.

Complications

- Allergic/anaphylactic Reaction
- Bleeding, local site pain, infection
- Common side effects (fever, headache, chills, muscle aches, fatigue)

References

- PFIZER-BIONTECH COVID-19 VACCINE (BNT162, PF-07302048) VACCINES AND RELATED BIOLOGICAL PRODUCTS ADVISORY COMMITTEE BRIEFING DOCUMENT MEETING DATE: 10 December 2020
- <u>https://www.cdc.gov/vaccines/hcp/acip-recs/vacc-specific/covid-19/clinicalconsiderations.html</u>
- <u>https://www.cdc.gov/vaccines/hcp/vis/index.html</u>
- CDC Vaccine Storage and Handling Toolkit November 2020