

Ensuring Proper COVID-19Vaccine Administration

April 8, 2021

Housekeeping

How to Ask Questions

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icon found at the bottom part of your screen

- A box will open where you can type in questions, comments, indicate sound problems, etc.
- Use this throughout the webinar to ask questions

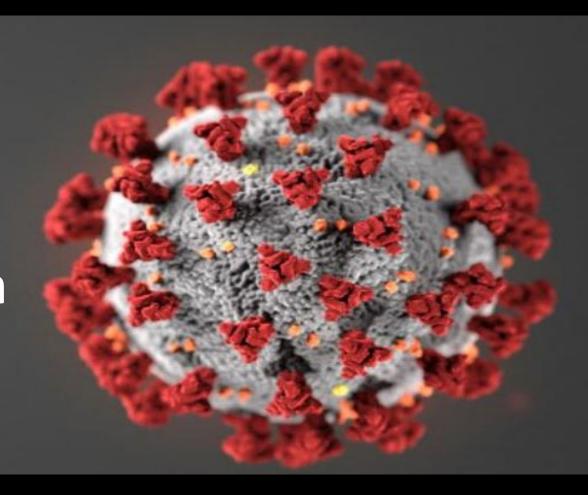
Slides & Recording

 This webinar is being recorded and a link as well as slides will be emailed out through our listserv as well as posted on our website at: www.michigan.gov/COVIDvaccine → Provider **Guidance and Education**

Topics Covered

- COVID-19 Vaccine Preparation/Administration
 - Pfizer-BioNTech
 - Moderna
 - Janssen
- COVID-19 Vaccine Administration Summary
- Administration Equipment and Anatomical Site
- COVID-19 Vaccine Administration Errors and Deviations
- Additional Vaccine Administration Resources

COVID-19 Vaccine Preparation/Administration



Vaccines & Immunizations

CDC









↑ COVID-19 Vaccination Product Info by US Vaccine Clinical Care Provider Requirements and Support Training and Education Vaccine Recipient Education Health Departments Planning & Partnerships

Vaccine Effectiveness Research

Vaccination Toolkits

COVID-19 Vaccination

Clinical Resources for Each COVID-19 Vaccine

Find information for COVID-19 vaccination administration, storage and handing, reporting, and patient education for each specific vaccine

Product Information by US Vaccine





Vaccines for COVID-19: Information for You and Your Family
Individuals looking for information about COVID-19 vaccines should visit the COVID-19 website.

U.S. COVID-19 Vaccine Product Information

Find a suite of information and materials that are needed for each specific COVID-19 vaccine that cover administration, storage and handling, safety, and reporting.

Pfizer-BioNTech

Moderna

Janssen/J&J

Requirements, Trainings, and Resources

Vaccine Storage and Handling Toolkit

COVID-19 Vaccine Quick Reference Guide for Healthcare Professionals

Training and Education

https://www.cdc.gov/vaccines/covid-19/info-by-product/index.html

Pfizer-BioNTech COVID

Vaccine Preparation and Administra



Vaccine Preparation and Administration Summary



Administer the Vaccine

Assess recipient status: Screen for contraindications and precautions. Review vaccination history.



Choose the correct equipment, including the correct needle size.

Review medical considerations.

 Use a new, sterile needle and syringe for each injection. Use low dead-volume syringes/needles to extract 6 doses from a single vial. If sufficient low-dead volume syringes are not available, withdraw vaccine using a combination of low dead-volume syringes and non-low dead-volume syringes per vial (e.g., 4 low dead-volume syringes and 2 non-low dead-volume syringes).



0.3 mL

Cleanse the stopper on the vial of mixed vaccine with a new, sterile alcohol prep pad. Withdraw 0.3 mL of mixed vaccine into the syringe.

- Regardless of the type of syringe used, ensure the amount of vaccine in the syringe equals 0.3 mL
- If the amount of vaccine remaining in the vial cannot provide a full 0.3 mL dose, discard the vial and contents.
- Do NOT combine vaccine from multiple vials to obtain a dose.



Scheduling Doses

Vaccination History†‡	And
0 doses	-
1 dose (Pfizer COVID-19 Vaccine)	It has been at least 21 days sinc dose 1
	It has not been at least 21 days from dose 1
2 doses (Pfizer COVID-19 Vaccine) at least 21 days apart ⁶	
2 doses (1 product unknown) at least 28 days apart [‡]	-

†mRNA COVID-19 vaccines should not be administered at the same time as other v or after the administration of mRNA COVID-19 vaccine, However, mRNA COVID-19 where the benefits of vaccination are deemed to outweigh the potential unknown as part of wound management, rabies vaccination for post-exposure prophylaxis, delays in mRNA COVID-19 vaccination.

Every effort should be made to determine wh vaccine product cannot be determined or is n

Administer the second dose as close to ti the series does not need to be restarted. Do

Pfizer-BioNTech COVID-19 Vaccine



General Information

Vaccine: COVID-19 vaccine (Pfizer)

Diluent: 0.9% sodium chloride (normal saline, preservative-free) Use a new vial every time.

Multidose vial: 6 doses per vial

Dosage: 0.3 mL

Vaccine MUST be mixed with diluent before administration.

» Age Indications

16 years of age and older

>> Schedule

2-dose series separated by 21 days

A series started with COVID-19 vaccine (Pfizer) should be completed with this product.

» Administration

Intramuscular (IM) injection in the deltoid muscle





» Thawing Frozen Vaccine

- Frozen vaccine must be thawed before using.
- Thaw vaccine in the refrigerator or at room temperature:
- Amount of time needed to thaw vaccine varies based on temperature and number of vials.
 - Do NOT refreeze thawed vaccine

Refrigerator: Between 2°C and 8°C (36°E and 46°E) Unpunctured vials may be sto How to Thaw, Prepare, and Administer the Pfizer-BioNTech Vaccine 120 hours (5 days).

Unpunctured vials cannot be more than 2 hours (including How to Thaw the Vaccine

Prepare the Vaccine

Follow aseptic technique. Perf before vaccine preparation, be when changing gloves (if wor hands become soiled."

Room temperature (for imm

Allow vaccine to come to room can be held at room temperatu before mixina.

Before mixing, check the expira vaccine and diluent, NEVER use diluent. The expiration dates fo the vaccine are located on the

How to Prepare the Vaccine

Remove vaccine from the freez

With the vaccine at room tempe vial 10 times. Do not shake the shaken, contact the manufacture white to off-white in color and n particles. Do not use if liquid is d

Using a new, sterile alcohol pre wipe off the stoppers of the dil vials. Using a 21-gauge (or narr withdraw 1.8 mL of 0.9% sodi (normal saline, preservative-fre syringe. Discard diluent vial and any remaining diluent every time. Do NOT use bacteriostatic

How to Administer the Vaccine

Scheduling Doses

COVID-19 Vaccine (Pfizer) Administration Resources

Pfizer COVID-19 Vaccine Standing Orders 12

Preparation and Administration Summary 🍱

Preparation Infographic [8] (new)

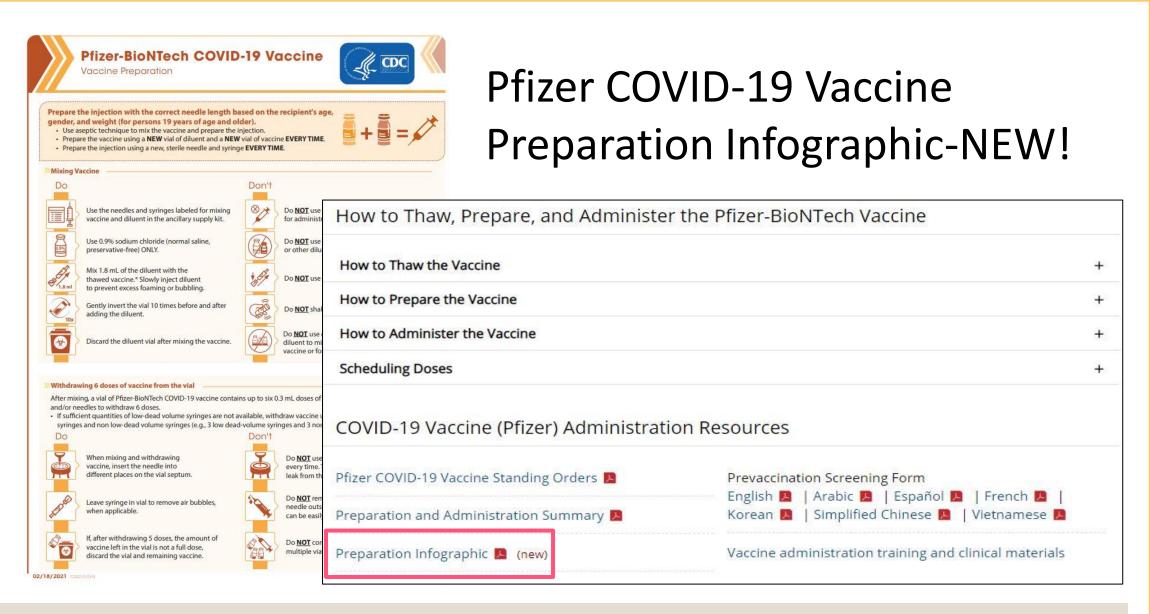
Prevaccination Screening Form

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Vaccine administration training and clinical materials

Do not return to freezer storage.

https://www.cdc.gov/vaccines/covid-19/info-by-product/pfizer/index.html



Maximizing Doses of Pfizer-BioNTech COVID-19 Vaccine



Consistently maximizing doses per vial of COVID-19 vaccines is essential to administering vaccines to more people. Selection of syringe and needle as well as technique for preparing doses to optimize vial pressure is key to maximizing doses for each Pfizer-BioNTech COVID-19 Vaccine vial. The USP COVID-19 Vaccine Handling Toolkit: Operational Considerations for Healthcare Practitioners, offers considerations to ensure complete doses are withdrawn and additional safe practices. Visit www.usp.org/covid-vaccine-handling to learn more.

Syringe and Needle Type

Maximize doses withdrawn from vials (at least 6 doses) by utilizing low-dead volume (LDV) syringes/needles whenever possible. Practice settings that may not have adequate quantities of LDV syringes can maximize dose by utilizing a combination of LDV and non-LDV syringes by utilizing a combination of LDV and non-LDV syringes. The ratio of LDV to non-LDV syringes shad 3 non-LDV syringes to do non-LDV syringes to do non-LDV syringes to do non-LDV syringes should be dependent on the type of syringe and needle used. Additional considerations to support dose optimization include:

- Use 1-inch needles, 21-gauge or narrower (e.g., 25-gauge), to withdraw vaccine.
- Use 1 mL syringes with 0.01 mL markings for accurate dose withdrawal.
- Use needle that is fixed to syringe,

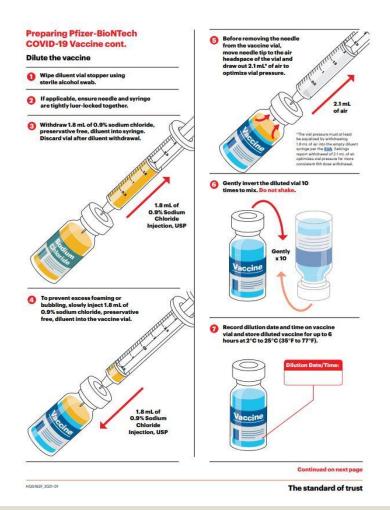
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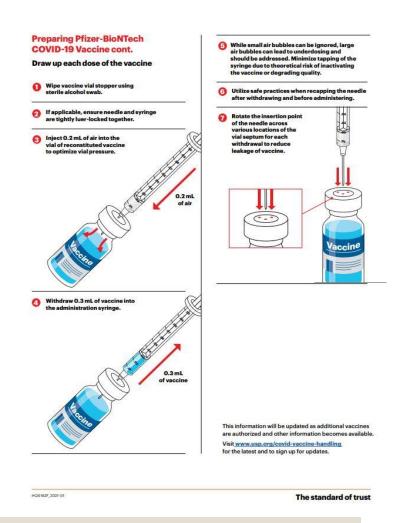
 Use the correct needle gauge and length for the recipient patient based on age, gender and weight recommendations based on <u>CDC's Vaccine</u>
 Administration: Needle Gauge and Length guide.

Preparing Pfizer-BioNTech COVID-19 Vaccine The Pfizer-BioNTech COVID-19 Vaccine resource webpage provides preparation instruction that should be reviewed to ensure quality vaccine preparation. The following are additional considerations for withdrawing doses including optimizing vial pressure to ensure maximizing doses for each Pfizer-BioNTech COVID-19 Vaccine vial. Follow aseptic technique throughout vaccine preparation. **Prepare for Dilution** A Pfizer-BioNTech COVID-19 vaccine vial must reach room temperature before dilution and be diluted within 2 hours of removal from frozen or refrigerated storage. Inspect liquid to ensure it is a white to offwhite suspension which may contain white to off-white opaque amorphous particles Invert vaccine vial gently 10 times. Do not shake.

Continued on next page

The standard of trust





https://www.usp.org/sites/default/files/usp/document/our-impact/covid-19/maximizing-doses-of-the-pfizer-biontech-covid19-vaccine.pdf

Pfizer-BioNTech Preparation/Administration Key Points

- Use aseptic technique to mix the vaccine and prepare the injection
 - Vaccine does not contain a preservative
- EVERY TIME
 - Use a NEW diluent vial and a NEW vaccine vial when preparing the vaccine
 - Use a NEW sterile needle and syringe when preparing the injection
- Check expiration dates on vaccine vial and diluent-Never use expired vaccine or diluent
- When mixing diluent with the vaccine
 - ONLY use the needles and syringes LABELED for mixing from the ancillary supply kit
 - Do NOT use the needles and syringes designated for administration

Pfizer-BioNTech Preparation/Administration Key Points Cont.

- Allow vaccine to come to room temperature
 - Vials can be held at room temperature for up to 2 hours before mixing
 - Amount of time needed to thaw vaccine varies based on temperature and number of vials
- Gently invert vial 10 times before and after adding diluent
 - If vial is shaken contact the manufacturer
- Mix 1.8 mL of diluent with thawed vaccine-Do NOT use all the diluent in the vial
 - Slowly inject diluent to prevent excess foaming or bubbling
 - Do NOT use bacteriostatic normal saline or other diluents to mix the vaccine
- Leave syringe in vial to remove air bubbles, when applicable

Pfizer-BioNTech Preparation/Administration Key Points Cont.

- When mixing and withdrawing vaccine, insert needle into different places on vial septum
- Keep mixed vaccine between 2°C and 25°C (36°F and 77°F) and administer within 6 hours.
 Discard any unused vaccine after 6 hours
 - Note the date/time the vaccine was mixed on the vial
- After withdrawing 5 doses, if amount of vaccine left in vial is not a full dose (0.3mL), discard the vial and remaining vaccine
 - Do NOT combine remaining vaccine from multiple vials to obtain a full dose
- Administer vaccine immediately by intramuscular (IM) injection in the deltoid muscle

U.S. COVID-19 Vaccine Product Information

Find a suite of information and materials that are needed for each specific COVID-19 vaccine that cover administration, storage and handling, safety, and reporting.

Pfizer-BioNTech

Moderna

Janssen/J&J

Requirements, Trainings, and Resources

Vaccine Storage and Handling Toolkit

COVID-19 Vaccine Quick Reference Guide for Healthcare Professionals

Training and Education

https://www.cdc.gov/vaccines/covid-19/info-by-product/index.html

Moderna COVID-19 Va Vaccine Preparation and Administra



Prepare and Administer the Vaccine (continued)

Ensure the needle and syringe are secured tightly together to prev during preparation and administration.

Withdraw 0.5 mL of vaccine into the syringe." Ensure the prepared syr

- · Discard vial when there is not enough vaccine to obtain a complete dose.
- Do NOT combine residual vaccine from multiple vial obtain a dose.

Note the date and time the vial was first punctured. Keep the vaccing (36°F and 77°F) for up to 6 hours. Discard any unused vaccine after

Bring the dose of vaccine from the designated preparation area in for administration.

Ensure staff has the correct PPE before administering vaccine and imp vaccine recipients (if tolerated).

Administer the vaccine immediately by intramuscular (IM) injection in

Observe recipients after vaccination for an immediate adverse reaction

- 30 minutes: Persons with a:
- History of an immediate allergic reaction of any severity to a vacc
- Contraindication to Janssen COVID-19 Vaccine who receive Mod
- History of anaphylaxis due to any cause

It is not necessary to change needles between drawing vaccine from a vial and inje-

Scheduling Doses

Vaccination History ^{†§}	And	
0 doses	→	
1 dose (Moderna COVID-19 Vaccine)	It has been at least 28 days since dose 1	
	It has not been at least 28 days since dose 1	
2 doses (Moderna COVID-19 Vaccine) at least 28 days apart [‡]	→	
2 doses (1 product unknown) at least 28 days apart [§]	→	

mRNA COVID-19 vaccines should not be administered at the same time as other vaccines or after the administration of mRNA COVID-19 vaccine. However, mRNA COVID-19 an the benefits of vaccination are deemed to outweigh the potential unknown risks of v wound management, rabies vaccination for post-exposure prophylaxis, measles or hi mRNA COVID-19 vaccination.

Every effort should be made to determine which determined or is no longer available, any availab

*Administer the second dose as close to the the series does not need to be restarted. Dos

Moderna COVID-19 Vaccine

Vaccine Preparation and Administration Summary



General Information

Vaccine: COVID-19 vaccine (Moderna) Multidose vial: 10 doses per vial Dosage: 0.5 mL Do NOT mix with a diluent.

» Age Indications 18 years of age and older

» Schedule

2-dose series separated by 1 month (28 days). A series started with COVID-19 vaccine (Moderna) should be completed with this product.

Administration Intramuscular (IM) injection in the deltoid muscle

>> Thawing Frozen Vaccine

- Frozen vaccine must be thawed before using.
- Thaw vaccine in the refrigerator or at room temperature:
- Refrigerator: Between 2°C and 8°C (36°F and 46°F). Unpunctured vials may be stored in the refrigerator for up to 30 days.
- Room temperature: Between 8°C and Unpunctured vials may be held at roon up to 12 hours.

Expiration Date

To determine the expiration date, scan the lookup option, enter the lot number, and www.modernatx.com/covid19vaccine-eu pfizer/downloads/expiration-tracker.pdf)

Prepare and Administer the Vacci

Assess recipient status:

Screen for contraindications and precautions.

Follow aseptic technique. Perform hand gloves (if worn), and any time hands become

Vaccine must be thawed before using. If 15 minutes

Unpunctured vials: Check the expiration Punctured vials: Check the beyond-use

With the vial upright, gently swirl the vac Note: Gently swirl the vaccine before with

Examine the vaccine. It should be white to use if liquid contains other particulate ma

Using a new, sterile alcohol prep pad, clea

- Amount of time needed to thaw vaccine varies based on temperature and number of vials.
- In the refrigerator: Approximately 2 hours and 30 minutes
- Room temperature: Approximately 1 hour
- Do NOT refreeze thawed vaccine.



How to Thaw, Prepare, and Administer the Moderna Vaccine

How to Thaw the Vaccine

How to Prepare the Vaccine

How to Administer the Vaccine

Scheduling Doses

COVID-19 Vaccine (Moderna) Administration Resources

Moderna COVID-19 Vaccine Standing Orders 🔼

Preparation and Administration Summary 2

Vaccine Expiration Date Tracking Tool 18

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Vaccine administration training and clinical materials

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Choose the correct equipment, including the correct needle size. Use a new, sterile needle and syringe for each injection.



https://www.cdc.gov/vaccines/covid-19/info-by-product/moderna/index.html

Moderna Preparation/Administration Key Points

- Use aseptic technique to prepare the injection
 - Vaccine does not contain a preservative
- DO NOT mix with a diluent
- Vaccine must be thawed before using. If removing the vial from the refrigerator, let it stand at room temperature for 15 minutes
 - Amount of time needed to thaw vaccine varies based on temperature and number of vials
- Check expiration date of vaccine and diluent-Never use expired vaccine or diluent
 - This includes the Beyond Use Date for punctured vials
- Choose the correct equipment, including the correct needle size
- Use a new, sterile needle and syringe for each injection
- Ensure the needle and syringe are tightly together to prevent vaccine from leaking

Moderna Preparation/Administration Key Points Cont.

- With vial upright, gently swirl the vaccine-Do NOT shake
 - If vial is shaken, contact the manufacturer
 - Gently swirl the vaccine before withdrawing subsequent doses
- Withdraw 0.5 mL of vaccine into the syringe
 - Discard vial when there is not enough vaccine to obtain a complete dose
 - Do NOT combine residual vaccine from multiple vials to obtain a dose
 - Regardless of the type of syringe used, ensure the amount of vaccine in the syringe equals
 0.5 mL
- Keep the vaccine between 2°C and 25°C (36°F and 77°F) for up to 12 hours. Discard any unused vaccine after 12 hours
 - Note the date and time the vial was first punctured
- Administer vaccine immediately by intramuscular (IM) injection in the deltoid muscle

U.S. COVID-19 Vaccine Product Information

Find a suite of information and materials that are needed for each specific COVID-19 vaccine that cover administration, storage and handling, safety, and reporting.

Pfizer-BioNTech

Moderna

Janssen/J&J

Requirements, Trainings, and Resources

Vaccine Storage and Handling Toolkit

COVID-19 Vaccine Quick Reference Guide for Healthcare Professionals

Training and Education

https://www.cdc.gov/vaccines/covid-19/info-by-product/index.html

Janssen COVID-19 Vac (Johnson & Johnson)

Vaccine Preparation and Administra

Contraindications and Precautions

Contraindications:

- Severe allergic reaction (e.g., anaphylaxis) to a component of Janssen COVID-19 Vaccine
- Immediate allergic reaction of any severity or known (diagnosed) allergy to a component of the vaccine (see Table 1 in this document for a list of ingredients in COVID-19 vaccines)

Note: Persons who have a contraindication to Janssen COVID-19 Vaccine may be able to receive an mRNA COVID-19 vaccine (see footnote).

Precautions:

- History of an immediate allergic reaction* to any other vaccine. or injectable therapy (i.e., intramuscular, intravenous, or subcutaneous vaccines or therapies)
- This includes persons with a reaction to a vaccine or injectable therapy that contains multiple components, one of which is polysorbate or another vaccine component, but for whom it is unknown which component elicited the immediate allergic reaction.
- People with a contraindication to either mRNA COVID-19 vaccine have a precaution to Janssen COVID-19 Vaccine.1
- Moderate or severe acute illness

For more information, please see Interim Clinical Considerations for Use of COVID-19 Vaccines Currently Authorized in the United States at www.cdc.gov/vaccines/covid-19/info-by-product/ clinical-considerations.html

Management of Anaphylaxis

Be prepared to manage medical emergencies.

- 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 Interim Considerations: Preparing for the Potential Management of Anaphylaxis after COVID-19 Vaccination at https://www.cdc.gov/vaccines/covid-19/ clinical-considerations/managing-anaphylaxis.html.

Document the Vaccination

COVID-19 vaccination providers must document vaccine administration in their medical record systems within 24 hours of administration. They should aim to report administration data to the relevant system for the jurisc information system) as soon as prahours after administration.

Janssen COVID-19 Vaccine (Johnson & Johnson)

Vaccine Preparation and Administration Summary



General Information

Vaccine: Janssen COVID-19 Vaccine (Johnson & Johnson) Multidose vial: 5 doses per vial

Do NOT mix with a diluent. Discard vial when there is not enough vaccine to obtain a complete dose. Do NOT combine residual vaccine from multiple vials to obtain a dose.

Age Indications

18 years of age and older

Schedule

Do not use the Janssen COVID-19 Vaccine as part of any other COVID-19 vaccine series.

Administration

Intramuscular (IM) injection in the deltoid muscle





Vaccine administration training and clinical materials

Expiration Date

Dosage: 0.5 mL

The expiration date is NOT printed on the vaccine vial or carton. To determine the expiration date:

- Scan the QR code located on the outer carton, or
- Call 1-800-565-4008, or
- Go to www.vaxcheck.ini.

Write date on carton. As the expiration date approaches, check the expiration date again. Do not discard vaccine until ensuring the expiration date has passed. Use CDC's expiration date tracking tool to document expiration date changes.

Prepare and Administer the Va

Assess recipient status:

- Screen for contraindications and precau
- Review vaccination history.
- Review medical considerations.

Follow aseptic technique. Perform hand hy vaccine preparation, between patients, whe gloves (if worn), and any time hands become

Unpunctured vials: Check the expiration d use expired vaccine.

Punctured vials: Check the beyond-use tim use vaccine after the beyond-use time.

With the vial upright, gently swirl the vacc seconds. Do NOT shake. If the vial is shake the manufacturer. Note: Gently swirl the va before withdrawing subsequent doses.

Examine the vaccine. It should be a colorle slightly yellow, clear to very opalescent sus Do not use if liquid contains particulate ma is discolored.

Using a new, sterile alcohol prep pad, clea stopper of the multidose vaccine vial.

Choose the correct equipment, including th needle size. Use a new, sterile needle and sy each injection.

How to Prepare and Administer the Janssen Vaccine

How to Prepare the Vaccine

How to Administer the Vaccine

Scheduling

lanssen COVID-19 Vaccine Administration Resources

Janssen COVID-19 Vaccine Standing Orders 🔼

Preparation and Administration Summary 12

Prevaccination Screening Form

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15 minutes: All other persons

https://www.cdc.gov/vaccines/covid-19/info-by-product/janssen/index.html

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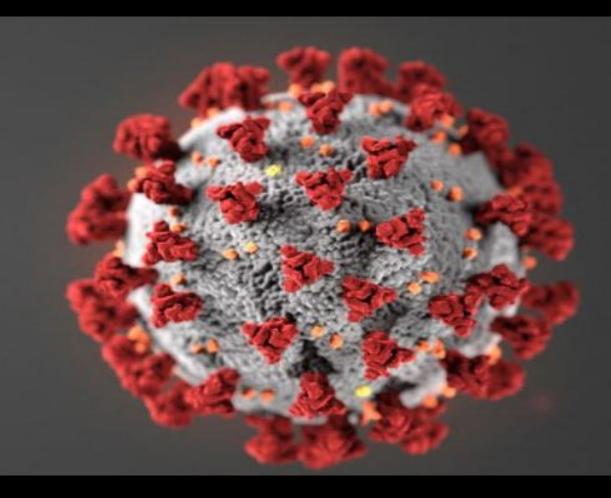
Janssen Preparation/Administration Key Points

- Use aseptic technique to prepare the injection
 - Vaccine does not contain a preservative
- DO NOT mix with a diluent
- Check expiration date of vaccine and diluent-Never use expired vaccine or diluent
 - This includes the Beyond Use Date for punctured vials
- Choose the correct equipment, including the correct needle size
- Use a new, sterile needle and syringe for each injection
- With vial upright, gently swirl the vaccine-Do NOT shake
 - If vial is shaken, contact the manufacturer
 - Gently swirl the vaccine before withdrawing subsequent doses

Janssen Preparation/Administration Key Points Cont.

- Ensure the needle and syringe are tightly together to prevent vaccine from leaking
- Withdraw 0.5 mL of vaccine into the syringe
 - Discard vial when there is not enough vaccine to obtain a complete dose
 - Do NOT combine residual vaccine from multiple vials to obtain a dose
 - Regardless of the type of syringe used, ensure the amount of vaccine in the syringe equals 0.5 mL
- Keep vaccine between 2°C and 8°C (36°F and 46°F) for up to 6 hours or at room temperature (up to 25°C or 77°F) for up to 2 hours. Discard if not used within this time
 - Note the date and time the vial was first punctured
- Administer vaccine immediately by intramuscular (IM) injection in deltoid muscle

COVID-19 Vaccine Administration Summary



Recap of Administration of COVID-19 Vaccines

 COVID-19 vaccines are administered intramuscularly as either a two-dose or one-dose series

Vaccine	Age Group	Dose Volume	Number Doses/Series	Interval Between Doses
Pfizer-BioNTech	16 years and older	0.3 mL	2	3 weeks (21 days)
Moderna	18 years and older	0.5 mL	2	1 month (28 days)
Janssen	18 years and older	0.5 mL	1	N/A

COVID-19 Vaccine

Quick Reference Guide for Healthcare Professionals

	Pfizer	Moderna	
Route	Intramuscular (IM) injection	Intramuscular (IM)	
Site	Deltoid	Deltoid	
Thawing Frozen	Between: 2°C and 8°C (36°F and 46°F) or	Between: 2°C and 8°C (36°F a	
Vaccine	Room temperature up to 25°C (77°F)	8°C to 25°C (46°F t	
	Do NOT refreeze thawed vaccine.	Do NOT refreeze t	
Mixing Vaccine	Mix vaccine with 1.8 mL of 0.9% sodium chloride (preservative-free, normal saline)	Do NOT mix with	
	Contraindications • Severe allergic reaction (e.g., anaphylaxis) after a previous COVID-19 vaccine • Immediate allergic reaction of any severity to a previous component of the vaccine		
	Note: Persons who have a contraindication to an mRNA be able to receive the Janssen COVID-19 vaccine (see for		
	Persons who have a contraindication to Janssen COVID- vaccine (see footnote).±		
Contraindications/	Precautions		
Precautions	 History of an immediate allergic reaction† to any other intravenous, or subcutaneous vaccines or therapies) 		
	» This includes people with a reaction to a vaccine or components, one of which is a vaccine component, elicited the immediate allergic reaction.		
	People with a contraindication to mRNA COVID-19 vacand vice versa. (see footnote).*		
	Moderate to severe acute illness		
	See Interim Clinical Considerations for Use of mRNA COV States <u>www.cdc.gov/vaccines/covid-19/info-by-product</u>		
Post-Vaccination Observation	30 minutes: Persons with a history of injectable therapy or a history of and 15 minutes: All other persons		
	Injection site: pain, swelling,	Injection site: pai	
Most common adverse events	redness Systemic: fatigue, headache, muscle pain, chills, fever, joint pain	Systemic: fatigue, pain, chills, fever, r	

For the purpose of this guidance, an immediate allergic reaction is defined as any hypersensitivity-related sidistress (e.g., wheezing, stridor), or anaphylaxis that occur within 4 hours following exposure to a vaccine or Consider consultation with an allergist-immunologist to help determine if the patient can safely receive vaccinat a consultation from the Clinical Immunization Safety Assessment COVID vax Project https://www.cdc.gov/vaccine these individuals should only be done in an appropriate setting under the supervision of a healthcare provider experience of the provider of the provi



Quick Reference Guide for Healthcare Professionals



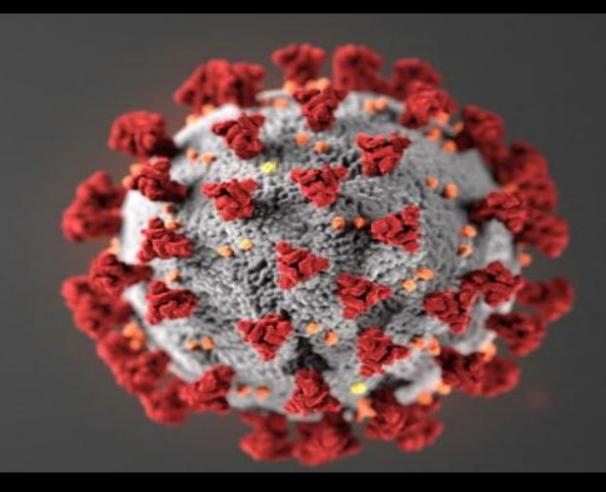
The table below provides basic information on the proper storage, preparation, and administration of the currently authorized COVID-19 vaccine products in the United States. For additional information and detailed clinical guidance go to the manufacturer's and CDC's webpages listed.

		Pfizer	Moderna	Janssen
G	EUA	www.fda.gov/emergency- preparedness-and-response/ coronavirus-disease-2019-covid-19/ pfizer-biontech-covid-19-vaccine	www.fda.gov/emergency- preparedness-and-response/ coronavirus-disease-2019-covid-19/ moderna-covid-19-vaccine	www.fda.gov/emergency- preparedness-and-response/ coronavirus-disease-2019- covid-19/janssen-covid-19-vaccine
N E	CDC Vaccine Information	www.cdc.gov/vaccines/covid-19/ info-by-product/pfizer/index.html	www.cdc.gov/vaccines/covid-19/ info-by-product/moderna/index. html	www.cdc.gov/vaccines/ covid-19/info-by-product/ janssen/index.html
R A L	Manufacturer Contact information	Website: www.cvdvaccine.com Medical information: 800-438-1985 Customer service: 800-879-3477	Website: www.modernatx.com Medical Information: 866-663-3762	Website: www.vaxcheck.inj. Medical information: 1-800-565-4008
	How supplied	Multidose vial: 6 doses	Multidose vial: 10 doses	Multidose vial: 5 doses
	Diluent	0.9% sodium chloride (preservative-free, normal saline) provided in the ancillary kit. Do NOT use other diluent.	None	None
STORAGE & HANDLI	Storage Temperatures: Before Puncture	Between: -80°C and -60°C (-112°F and -76°F) until the expiration date -25°C and -15°C (-13°F and 5°F) for up to 2 weeks 2°C and 8°C (36°F and 46°F) for up to 120 hours (5 days)	Between: -25°C and -15°C (-13°F and 5°F) until the expiration date 2°C and 8°C (36°F and 46°F) for up to 30 days	Between: 2°C and 8°C (36°F and 46°F) until the expiration date.
	Storage Temperatures: After puncture	Between: 2°C to 25°C (36°F to 77°F) for up to 6 hours. Discard any unused vaccine after 6 hours.	Between: 2°C and 25°C [36°F and 77°F] for up to 6 hours. Discard any unused vaccine after 6 hours.	Between: 2°C and 8°C (36°F and 46°F) for up to 6 hours. 9°C and 25°C (47°F and 77°F) for up to 2 hours. Discard any unused vaccine after these time frames.
N	Transport Temperatures: Before Puncture	Between: -80°C and -60°C (-112°F and -76°F) -25°C and -15°C (-13°F and 5°F) 2°C and 8°C (36°F and 46°F)	Between: -25°C and -15°C (-13°F and 5°F) 2°C and 8°C (36°F and 46°F) for up to 12 cumulative hours.	Between: 2°C and 8°C (36°F and 46°F)
	After Puncture	Between: 2°C to 25°C (36°F to 77°F) for up to 6 hours.	Between: 2°C and 25°C (36°F and 77°F) for up to 6 hours.	Between: 2°C and 8°C (36°F and 46°F) for up to 6 hours
	Type of Vaccine	mRNA	mRNA	Viral vector
	Age Indications	16 years of age and older	18 years of age and older	18 years of age and older
	Schedule	2-doses, separated by 21 days. Both doses must be Pfizer- BioNTech vaccine	2 doses, separated by 28 days. Both doses should be Moderna vaccine	1 dose only
	Dosage	0.3 mL	0.5 mL	0.5 mL
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COVID-19 Quick Reference Guide

https://www.cdc.gov/vaccines/covid-19/downloads/covid19-vaccine-quick-reference-guide-2pages.pdf

Administration Equipment and Anatomical Site





Vaccine Administration: Needle Gauge and Length

Vaccines must reach the desired tissue to provide an optimal immune response and reduce the likelihood of injection-site reactions. Needle selection should be based on the:

· Route

Age

 Gender and weight for adults

Injection site

(19 years and older)

The following table outlines recommended needle gauges and lengths. In addition, clinical judgment should be used when selecting needles to administer injectable vaccines.

Route	Age	Needle gauge and length	Injection site
Subcutaneous injection	All ages	23–25-gauge 5/8 inch (16 mm)	Thigh for infants younger than 12 months of age ¹ ; upper outer triceps area for persons 12 months of age and older
	Neonate, 28 days and younger	22–25-gauge 5/8 inch (16 mm²)	Vastus lateralis muscle of anterolateral thigh
	Infants, 1–12 months	22–25-gauge 1 inch (25 mm)	Vastus lateralis muscle of anterolateral thigh
	Toddlers, 1–2 years	22–25-gauge 1–1.25 inches (25–32 mm)	Vastus lateralis muscle of anterolateral thigh ³
		22–25-gauge 5/8 ² –1 inch (16–25 mm)	Deltoid muscle of arm
Intramuscular	Children, 3–10 years	22–25-gauge 5/8 ² –1 inch (16–25 mm)	Deltoid muscle of arm ³
injection		22–25-gauge 1–1.25 inches (25–32 mm)	Vastus lateralis muscle of anterolateral thigh
	Children, 11–18 years	22-25-gauge 5/8 ² -1 inch (16-25 mm)	Deltoid muscle of arm ^{3,5}
	Adults, 19 years and older 130 lbs (60 kg) or less 130–152 lbs (60–70 kg) Men, 152–260 lbs (70–118 kg) Women, 152–200 lbs (70–90 kg) Men, 260 lbs (118 kg) or more Women, 200 lbs (90 kg) or more	22–25-gauge 1 inch (25 mm ⁴) 1 inch (25 mm) 1–1.5 inches (25–38 mm) 1–5 inches (35–38 mm) 1.5 inches (38 mm) 1.5 inches (38 mm)	Deltoid muscle of arm ^{3,5}

May be administered into the upper outer triceps area if necessary

Needle Gauge and Length

https://www.cdc.gov/vaccines/hcp/admin/downloads/vaccine-administration-needle-length.pdf

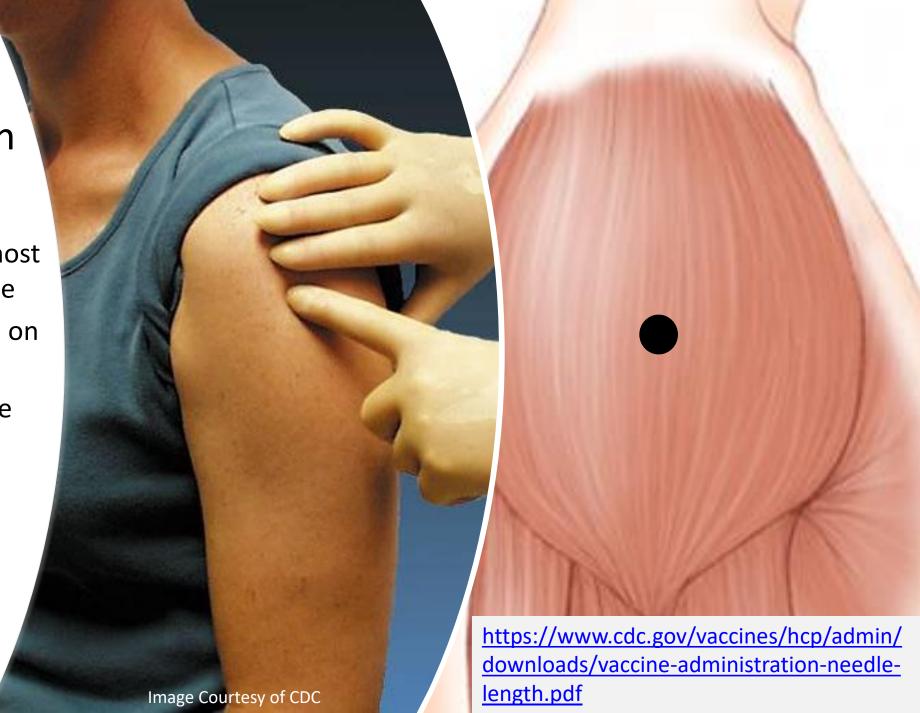
² If the skin is stretched tightly and subcutaneous tissues are not bunched

²Preferred site

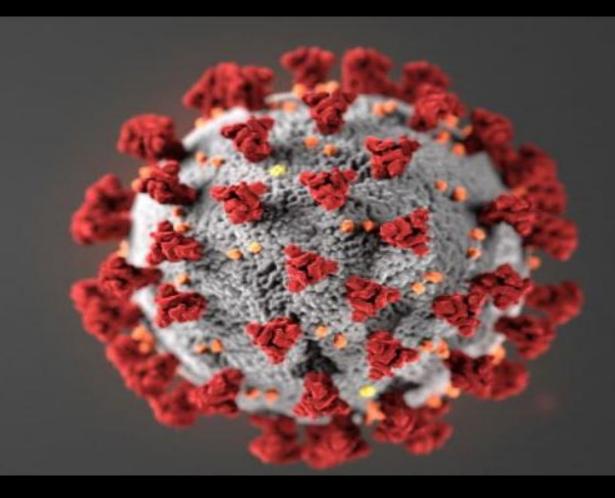
^{*}Some experts recommend a 5/8-inch needle for men and women weighing less than 60 kg, if used, skin must be stretched tightly and subcutaneous tissues must not be bunched



- Administer in thickest, most central part of the muscle
- Use needle length based on patient's age and weight
- Insert the needle into the muscle at a 90° angle
- Administer alone
 (14 days before or after other vaccines)



COVID-19 Vaccine Administration Errors and Deviations



COVID-19 Va

Administration Errors c

COVID-19 Vaccine

Administration Errors and Deviations



Interim recommendations for COVID-

Vaccines	Type	Ad
mRNA vaccines only (Pfizer-	Intervals	• S t f t
BioNTech and Moderna)	Mixed series	• 1: • 1: • 0:
Pfizer- BioNTech only	Diluent	• C
		• N
		• 11
		• li

Pfizer-BioNTech and Moderna vaccines only: *If the dose given in error is the first dose, a second dos this dose is the second dose, the series is complete, and "Do not administer the second dose until the person be extended to include additional age groups), even if this #If the dose given in error is the first dose, the second d from the date of receipt of the valid dose (not the date [†]If the administration error resulted in a higher-than-au However, if local or systemic side effects following vacci reactions, or are ongoing at the time of the second dose Although CDC provides considerations for a mixed seri reporting (as a mixed series is not authorized under the A vaccine administration error is any preventable event that may cause or lead to inappropriate use of vaccine or patient harm. This table provides resources for preventing and reporting COVID-19 vaccine administration errors, as well as actions to take after an error has occurred. For completeness, it includes additional scenarios that deviate from CDC recommendations for vaccine intervals but are not considered administration errors.

For all vaccine administration errors:

- Inform the recipient of the vaccine administration error.
- Consult with the state immunization program and/or immunization information system (IIS) to determine how the dose should be entered into the IIS, both as an administered dose and to account for inventory.
- Providers are required to report all COVID-19 vaccine administration errors—even those not associated with an adverse event—to VAERS.
- . Determine how the error occurred and implement strategies to prevent it from happening again.

Interim recommendations for COVID-19 vaccine administration errors and deviations

Vaccines	Туре	Administration error/deviation	Interim recommendation
	Site/route	Incorrect site (i.e., site other than the deltoid muscle [preferred site] or anterolateral thigh [alternate site])	Do not repeat dose.* Inform the recipient of the potential for local and systemic adverse events.
		Incorrect route (e.g., subcutaneous)	 Do not repeat dose.* Inform the recipient of the potential for local and systemic adverse events.
All currently	Age	Unauthorized age group	If received dose at age less than 16 years, do not give any additional dose at this time." If age 16 to 17 years and a vaccine other than Pfizer-BioNTech was inadvertently administered: If Moderna vaccine administered as the first dose, may administer Moderna vaccine as the second dos (as off-label use, because Moderna vaccine is not authorized in this age group). If Janssen vaccine administered, do not repeat dose with Pfizer-BioNTech vaccine.
authorized	Dosage	Higher-than-authorized dose volume administered	Do not repeat dose.*†
(Pfizer- BioNTech Moderna, and Janssen COVID-19		Lower-than-authorized dose volume administered (e.g., leaked out, equipment failure, recipient pulled away)	 If more than half of the dose was administered, do nor repeat dose.⁸ If less than half of the dose was administered or the proportion of the dose cannot be estimated, administer the authorized dose immediately (no minimum interval) in the opposite arm.⁸
vaccines) Inactive ingredients	Storage and handling	Dose administered after improper storage and handling (e.g., temperature excursion, more than allowed time after first vial puncture)	 Contact the manufacturer for guidance. If the manufacturer provides information supporting that the dose should be repeated, the repeated dose may be given immediately (no minimum interval) in the opposite arm.
		Dose administered past the expiration/beyond-use date	 Contact the manufacturer for guidance. If the manufacturer provides information supporting that the dose should be repeated, the repeated dose may be given immediately (no minimum interval) in the opposite arm.
	Coadministration	Dose administered within 14 days before or after another (i.e., non-COVID-19) vaccine	 Do not repeat COVID-19 vaccine* or other vaccine(s) doses. This deviation from CDC guidance does not require VAERS reporting.
		Dose administered within 90 days of monoclonal antibodies or convalescent plasma for	 Do not repeat COVID-19 vaccine dose. If person has already received one mRNA COVID-19 vaccine dose, defer administration of second dose for 90 days

Resource for preventing and reporting COVID-19 vaccine administration errors, as well as a simple table outlining actions to take after an error has occurred. Errors addressed include the following:

- Incorrect route or site
- Incorrect age
- Incorrect dosing interval
- Incorrect dose (high or low)
- Administration after improper storage conditions
- Administration after the expiration/beyond use date
- Diluent errors (wrong diluent, too much or too little) with the Pfizer-BioNTech vaccine

https://www.cdc.gov/vaccines/covid-19/downloads/covid19-vaccine-errors-deviations.pdf

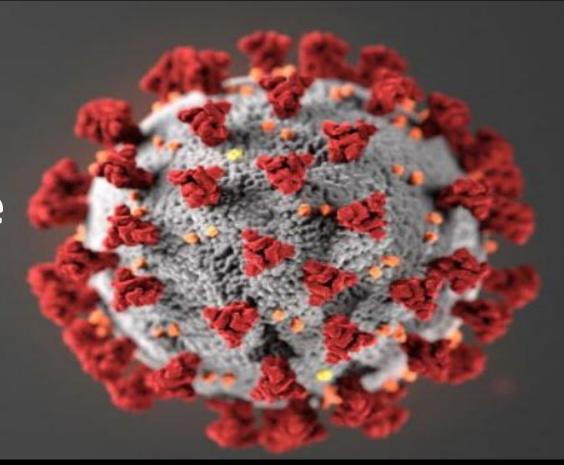
- Do Not Repeat the Dose
- If the dose given in error is the first dose, a second dose should be administered at the recommended interval (21 days [Pfizer-BioNTech] or 28 days [Moderna]).
- If this dose is the second dose, the series is complete, and no additional doses are needed
- If the administration error resulted in a higher-thanauthorized vaccine dose, in general the second dose may still be administered at the recommended interval. However, if local or systemic side effects following vaccination are clinically concerning (outside of the expected side effect profile), lead to serious adverse reactions, or are ongoing at the time of the second dose, the decision to administer the second dose may be assessed on a case-by-case basis
- Report to VAERS at https://vaers.hhs.gov/

What happens when someone administers 1.0mL of Moderna vaccine instead of 0.5mL?

- If more than half of the dose was administered, do not repeat dose
 - If the dose given in error is the first dose, a second dose should be administered at the recommended interval (21 days [Pfizer-BioNTech] or 28 days [Moderna]). If this dose is the second dose, the series is complete, and no additional doses are needed
- If less than half of the dose was administered or the proportion of the dose cannot be estimated, administer the authorized dose immediately (no minimum interval)
 - If the dose given in error is the first dose, the second dose should be administered at the recommended interval (21 days [Pfizer-BioNTech] or 28 days [Moderna]) from the date of receipt of the valid dose (not the date of receipt of the erroneous dose) in the opposite arm
- Report to VAERS at https://vaers.hhs.gov/

When administering Pfizer, a significant amount of vaccine leaked out. What should we do?

Additional COVID-19 Vaccine Administration Resources



Resource Library

Resource Library

Note: The materials listed on this page might be more current than vaccine administration information in previously published CDC documents, including the 13th edition of *Epidemiology and Prevention of Vaccine-Preventable Diseases* (the Pink Book). Always follow the most up-to-date guidelines in the Vaccine Storage and Handling Toolkit or more recently dated materials.

On This Page Web-based Training Courses Job Aids References Web Button

Web-based Training Courses

Vaccine Administration e-Learn

A self-paced vaccine administration course that provides comprehensive training using videos, job aids, and other resources.

You Call the Shots

An interactive, web-based immunization training course that includes the latest guidelines and recommendations in vaccine practice.



Title: Comfort and Restraint Techniques

Short Description: This training demonstrates comfort and restraint techniques. Determine the best position for the patient based on comfort, age, activity level, administration site, and safety. Instruct the parent on how to help the infant or child stay still so you can administer the vaccine(s) safely.

Title: Assemble a Manufacturer-filled Syringe

Short Description: This training addresses how to assemble a manufacturer-filled syringe, available for a variety of vaccines. CDC recommends that providers only prepare vaccines just prior to administration. Always prepare vaccines in a designated area that is not near any area where potentially contaminated items are placed.

Title: Single-Dose Vial

Short Description: This training addresses he dose and should be administered one time to vaccine just prior to administration.

https://www.cdc.gov/vaccines/hcp/admin/resource-library.html

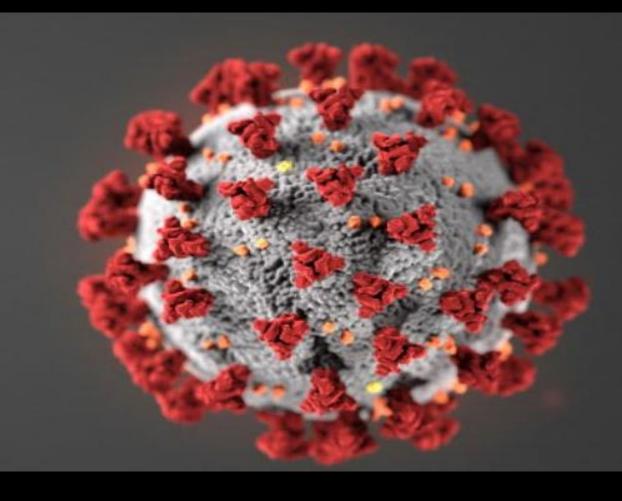
CDC's Training Materials

- Web-based training courses
 - Vaccine Administration e-Learn
 - You Call the Shots Modules
- Videos
 - Comfort and Restraint Technique
 - Assemble a Manufacturer-filled Syringe
 - Single-Dose Vial
 - Subcutaneous Injection
 - Intramuscular Injection
- Job Aids
 - Vaccine Administration: Needle Gauge and Length
 - Vaccine Administration: Intramuscular (different age groups)
 - Vaccine Administration: Preventing Vaccine Administration Errors
- References

Thank You!

Next "Noontime Knowledge" Update: April 22, 2021 at 12:00p.m. Topic: TBD

Please watch your email for an updated link and topic!



www.michigan.gov/COVIDvaccine -> Provider Guidance and Education