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# Road to Recovery: Update



Information current as of 3.04.2021

## Seniors and Staff Vaccine Doses to Date

54,202



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## Current Consonus Stats:



Clinics Held

727

Total  
Vaccines to  
Date

54,202

Booster  
Doses

26,265

Resident  
Consent

89%

Staff Consent

74%

Done with  
3<sup>rd</sup> Clinic

237



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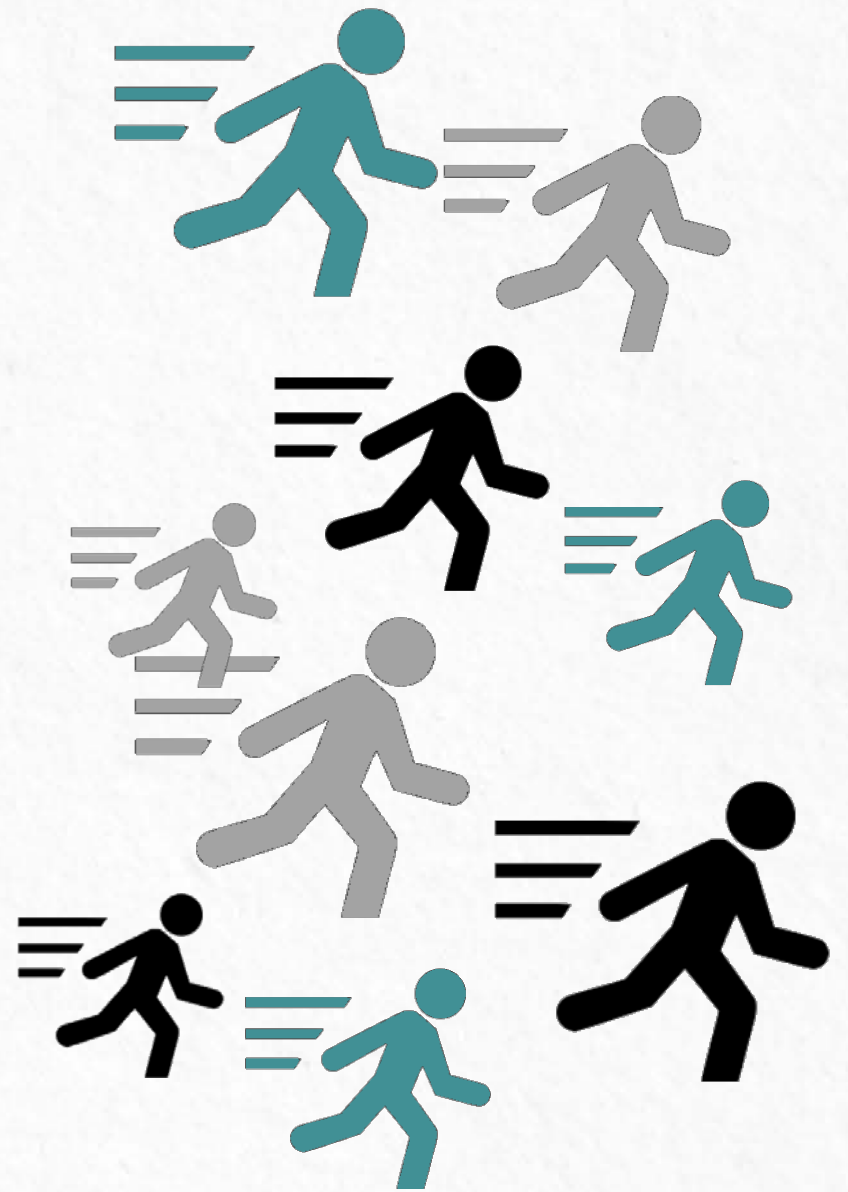




## Current State

### What we Know.....

- Important to keep going!
- Drop in case rate
- Reopening plan
- 5,000 Residents / 8,000 Staff left to vaccinate
- Consonus has 500 new recipients/week
- Pfizer is still available
- Moderna 2 weeks out
- The federal government needs to activate the "Retail" program




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# Phase 2 Federal “Retail” Program: Vaccine Continuity for LTC:



Centers for Disease Control and Prevention  
CDC 24/7: Saving Lives, Protecting People™





Vaccines site 


[Advanced Search](#)

[A-Z Index](#)

Vaccines & Immunizations

CDC > COVID-19 Vaccination > Planning & Partnerships > Long-Term Care Pharmacy Partnerships

 COVID-19 Vaccination

Product Info by US Vaccine +

Clinical Care +

Provider Requirements and Support

Training and Education

Recipient Education +

Planning & Partnerships -

COVID-19 Vaccination Program Operational Guidance

Federal Retail Pharmacy Program +

Long-Term Care Pharmacy Partnerships -

Ensuring Access in Long-term Care Facilities

FAQs about Long-Term Care Pharmacy Partnerships

Vaccine Effectiveness Research

Vaccination Toolkits +

## Ensuring Access to COVID-19 Vaccine in Long-Term Care Facilities

Long-term care facilities (LTCFs) provide a range of services, including medical and personal care, to adults who are unable to live independently. These types of facilities include, but are not limited to, skilled nursing facilities, nursing homes, assisted living facilities, and other congregate living settings where most people receiving care/supervision are older than 65 years of age. The communal nature of LTCFs and the population served (generally older adults often with underlying medical conditions) puts facility residents at increased risk of SARS-CoV-2 infection and severe illness from COVID-19.

The [Pharmacy Partnership for Long-Term Care \(LTC\) Program](#) is facilitating on-site vaccination of residents and staff at more than 70,000 enrolled LTCFs. Pharmacy partners participating in the program are conducting three on-site vaccination clinics for every enrolled facility. Most clinics provided by this program will likely be completed by the end of March 2021. As the Pharmacy Partnership for LTC program comes to an end, it is important to ensure LTCFs have continued access to COVID-19 vaccine for new or unvaccinated residents and staff.

## Ways for Long-term Care Facilities to Continue Receiving COVID-19 Vaccine

LTCF administrators and clinical leadership should consider the following options when making COVID-19 vaccination plans for their facilities.

LTCFs can receive COVID-19 vaccine from an LTC pharmacy that is enrolled as a COVID-19 vaccine provider with their state or territory.	+
LTCFs can receive COVID-19 vaccine from a long-term care (LTC) pharmacy* that is enrolled as a COVID-19 vaccine provider through the Federal Retail Pharmacy Program for COVID-19 Vaccination.	+
LTCFs can receive COVID-19 vaccine by enrolling directly with their state or territory as a COVID-19 vaccine provider.	+

<https://www.cdc.gov/vaccines/covid-19/long-term-care/pharmacy-partnerships-access.html>

# Vaccine Continuity Program for LTC

- Change From Clinics to Dispensing
- Moderna Product  
(Pfizer remains available)
- Federal program, applies to all States  
(requires state level activation)
- No Sign Up Required
- CDC Provider Agreement Remains
- Pharmacy Manages Reporting



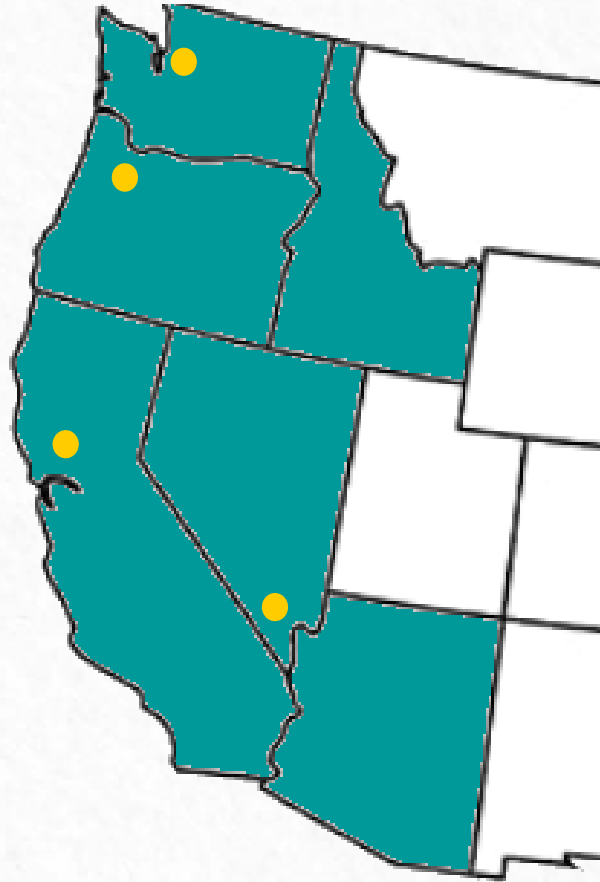
## State Update:

### Oregon / Washington

- Activation expected
- State initiatives for standing orders

### California

- Working with CDC for state activation



### Idaho/ Arizona

- National advocacy across state line

### Nevada

- Product allocation



# The Goal:



Maintain  $\geq 75\%$  full vaccination protection within LTC facilities.



Decrease community positivity rates.

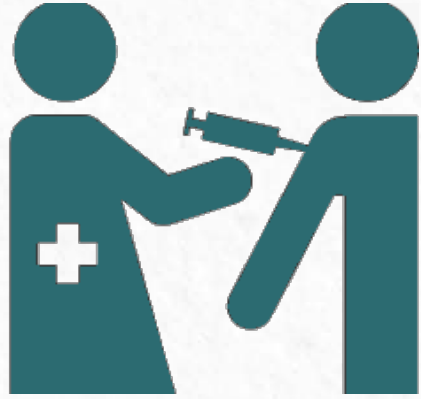


Reopen! Bring community and families back together.



Maximize product, minimizing waste

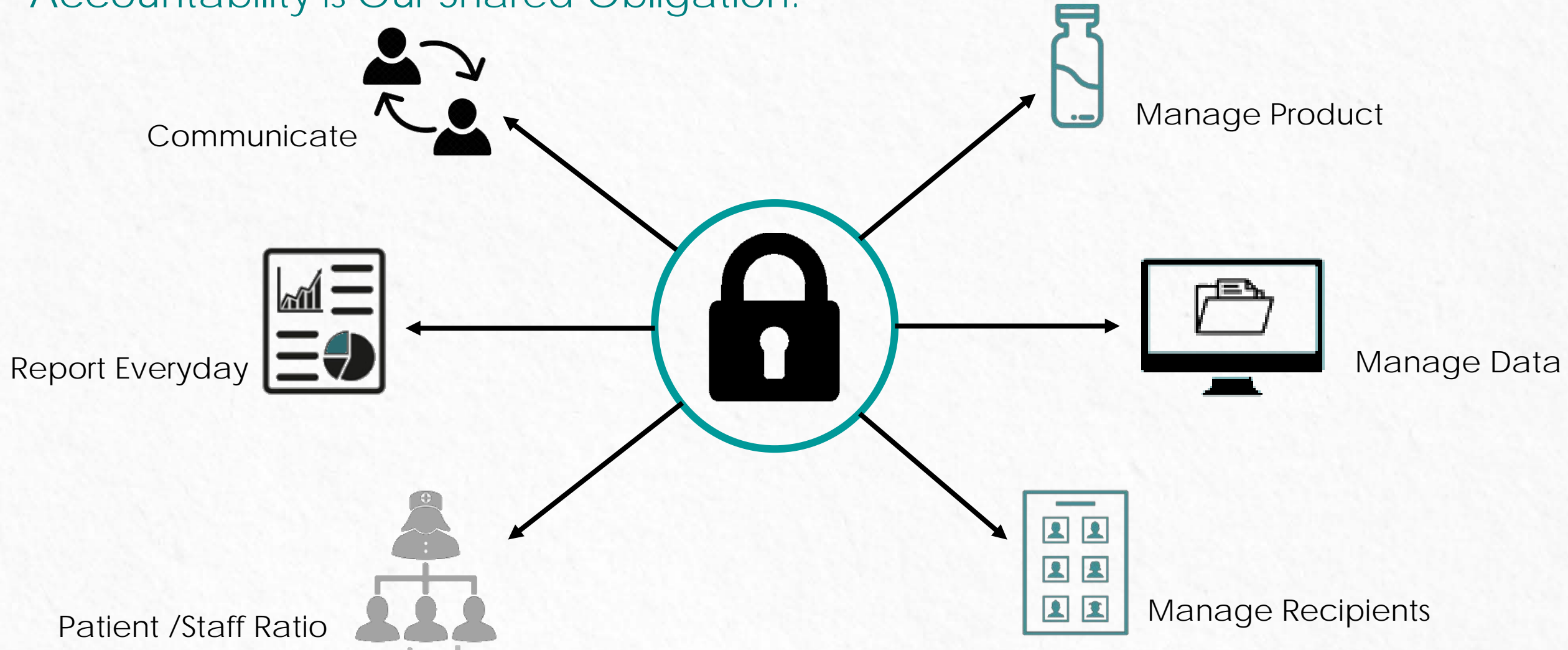
## Policy Considerations:



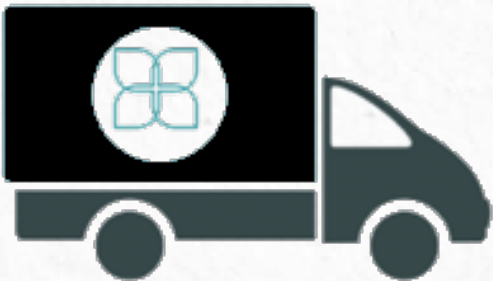
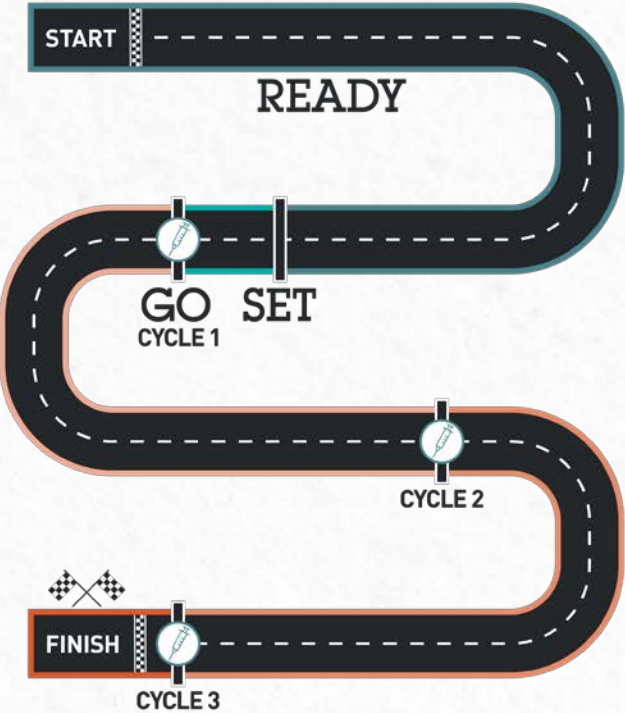
Leveraging  
Emergency Use  
Authorization for:

- Vaccinators
- Emergency Supply
- Funding

# Accountability is Our Shared Obligation:

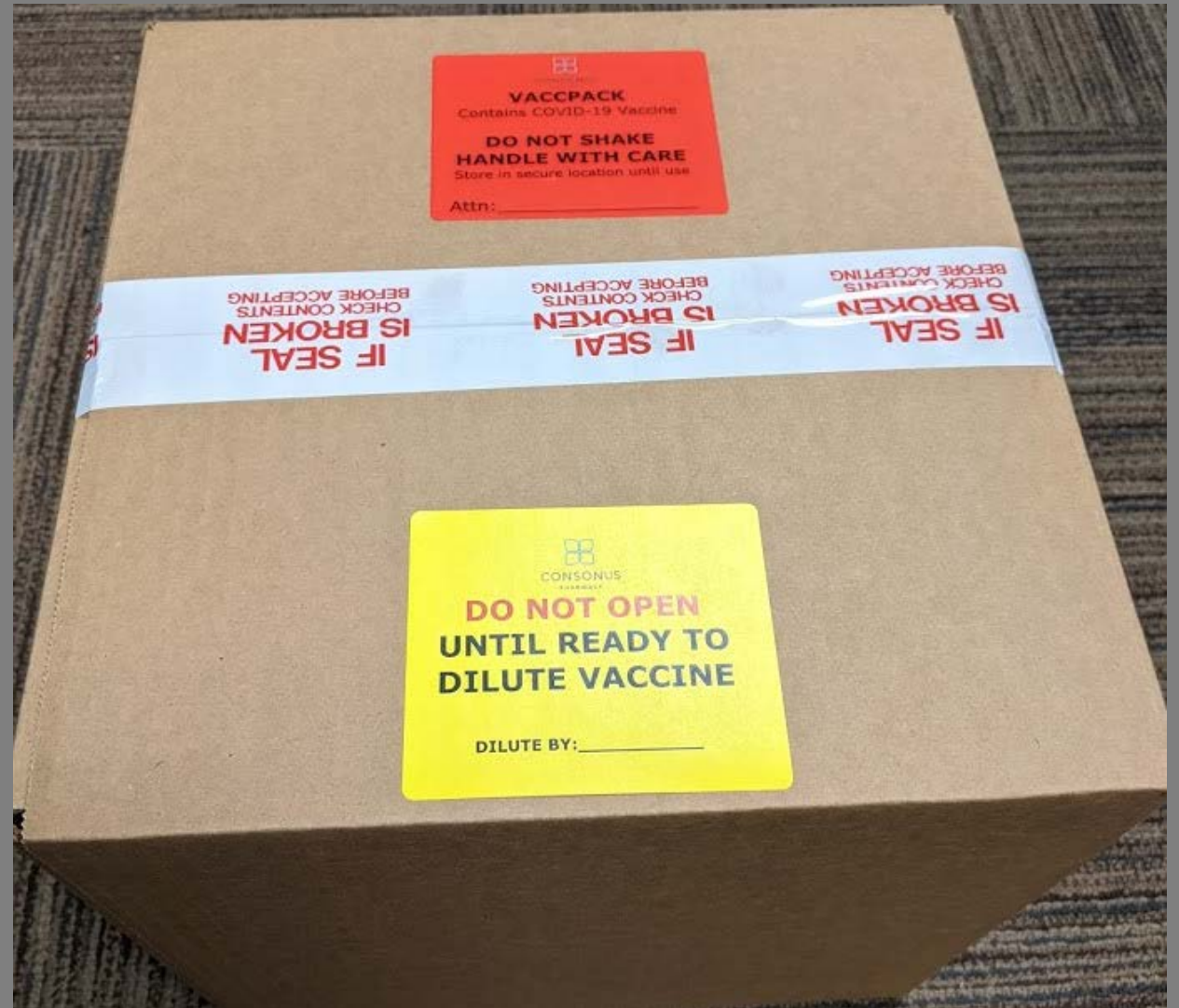


# Moving from Clinics to Continuity:





# Introducing VaccPack



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Activate:

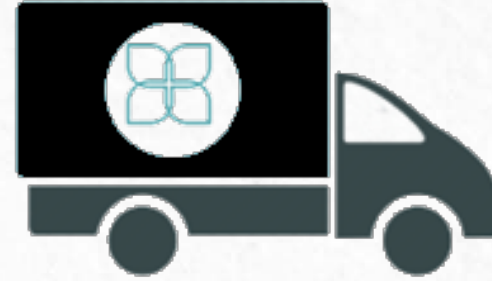


## Activate

- Agreement
- Identify FC
- Identify Vaccinator
- FC Website Training
- Vaccinator Training
- Life Enrichment List
- Standing Orders
- Physician Order



## Order



## Receive



## Vaccinate



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# Clinical Standing Orders for Administration

## Moderna COVID-19 Vaccine

Standing Orders for Administering Vaccine  
to Persons 18 Years of Age and Older



**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).

### » 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 18 years of age and older for vaccination with Moderna COVID-19 Vaccine based on the following criteria:

- No complete 2-dose COVID-19 vaccination history, regardless of brand. If 2 doses of a same-brand or mixed-brand series have been administered, no additional doses are recommended.
  - If the recipient has received 1 previous dose of Moderna COVID-19 Vaccine, a second dose of the same brand should be administered.
- If the vaccine product previously given cannot be determined or is no longer available, any mRNA COVID-19 vaccine product may be administered at least 28 days after the first dose.
- This vaccine is administered in a 2-dose series. Separate doses by at least 28 days, preferably within 6 weeks.\*

- Moderna COVID-19 Vaccine should not be administered at the same time as other vaccines. Separate Moderna COVID-19 Vaccine from other vaccines by 14 days before or after the administration of Moderna COVID-19 Vaccine.<sup>†</sup>

- Moderna COVID-19 Vaccine should be deferred for at least 90 days for persons who received passive antibody therapy (monoclonal antibodies or convalescent plasma) as part of COVID-19 treatment.

- If the recipient has a history of dermal filler use, advise them to contact their healthcare provider for evaluation if they develop swelling at or near the dermal filler site following vaccination.

- Screen for contraindications and precautions.

#### o Contraindications:

- Severe allergic reaction (e.g., anaphylaxis) to a previous dose or component of either mRNA COVID-19 vaccine.
- Immediate allergic reaction<sup>‡</sup> of any severity to a previous dose or component of an mRNA COVID-19 vaccine (including polyethylene glycol [PEG]). See Table 1 of vaccine components on page 3.
- Immediate allergic reaction of any severity to polysorbate (due to potential cross-reactive hypersensitivity with the vaccine ingredient PEG).

#### o Precautions:

- History of an immediate allergic reaction to any other vaccine or therapies [excluding subcutaneous immunotherapy of allergies, i.e., "allergy shots"] not related to a component of mRNA COVID-19 vaccines or polysorbate).
- Moderate to severe acute illness.

- Provide all recipients with a copy of the current federal Emergency Use Authorization (EUA) Fact Sheet for Recipients and Caregivers.
- Prepare to administer the vaccine.

Sex and Weight of Patient	Needle Gauge	Needle Length	Injection Site <sup>†</sup>
Female or male fewer than 130 lbs	22–25	¾"–1"	Deltoid muscle of arm
Female or male 130–152 lbs	22–25	1"	Deltoid muscle of arm
Female 152–200 lbs	22–25	1–1½"	Deltoid muscle of arm
Male 153–260 lbs	22–25	1–1½"	Deltoid muscle of arm
Female 200+ lbs	22–25	1½"	Deltoid muscle of arm
Male 260+ lbs	22–25	1½"	Deltoid muscle of arm

\*Administer the second dose as close to the recommended interval (28 days) as possible. If the second dose is not administered within 6 weeks of the first dose, the series does not need to be restarted. Doses inadvertently administered less than 28 days apart do not need to be repeated.

†However, mRNA COVID-19 and other vaccines may be administered within a shorter period in situations where the benefits of vaccination are deemed to outweigh the potential unknown risks of vaccine coadministration (e.g., tetanus toxoid-containing vaccination as part of wound management, rabies vaccination for post-exposure prophylaxis, measles or hepatitis A vaccination during an outbreak) or to avoid barriers or delays to mRNA COVID-19 vaccination.

‡For the purpose of this guidance, 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.

†Alternatively, the anterolateral thigh also can be used.

†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).

02/11/2021 CS321571.H

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## Pfizer-BioNTech COVID-19 Vaccine

Standing Orders for Administering Vaccine  
to Persons 16 Years of Age and Older



**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).

### » 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 16 years of age and older for vaccination with Pfizer-BioNTech COVID-19 Vaccine based on the following criteria:

- No complete 2-dose COVID-19 vaccination history, regardless of brand. If 2 doses of a same-brand or mixed-brand series have been administered, no additional doses are recommended.
  - If the recipient has received 1 previous dose of Pfizer-BioNTech COVID-19 Vaccine, the second dose of the same brand should be administered.
- If the vaccine product previously given cannot be determined or is no longer available, any mRNA COVID-19 vaccine product may be administered at least 28 days after the first dose.
- This vaccine is administered in a 2-dose series. Separate doses by at least 28 days, preferably within 6 weeks.\*

- Pfizer-BioNTech COVID-19 Vaccine should not be administered at the same time as other vaccines. Separate Pfizer-BioNTech COVID-19 Vaccine from other vaccines by 14 days before or after the administration of Pfizer-BioNTech COVID-19 Vaccine.<sup>†</sup>

- Pfizer-BioNTech COVID-19 Vaccine should be deferred for at least 90 days for persons who received passive antibody therapy (monoclonal antibodies or convalescent plasma) as part of COVID-19 treatment.

- Screen for contraindications and precautions.

#### o Contraindications:

- Severe allergic reaction (e.g., anaphylaxis) to a previous dose or component of either mRNA COVID-19 vaccine.
- Immediate allergic reaction<sup>‡</sup> of any severity to a previous dose or component of an mRNA COVID-19 vaccine (including polyethylene glycol [PEG]). See Table 1 of vaccine components on page 3.
- Immediate allergic reaction of any severity to polysorbate (due to potential cross-reactive hypersensitivity with the vaccine ingredient PEG).

#### o Precautions:

- History of an immediate allergic reaction to any other vaccine or therapies [excluding subcutaneous immunotherapy of allergies, i.e., "allergy shots"] not related to a component of mRNA COVID-19 vaccines or polysorbate).
- Moderate to severe acute illness.

- Provide all recipients with a copy of the current federal Emergency Use Authorization (EUA) Fact Sheet for Recipients and Caregivers.
- Prepare to administer the vaccine.

Sex and Weight of Patient	Needle Gauge	Needle Length	Injection Site <sup>†</sup>
Female or male fewer than 130 lbs	22–25	¾"–1"	Deltoid muscle of arm
Female or male 130–152 lbs	22–25	1"	Deltoid muscle of arm
Female 152–200 lbs	22–25	1–1½"	Deltoid muscle of arm
Male 153–260 lbs	22–25	1–1½"	Deltoid muscle of arm
Female 200+ lbs	22–25	1½"	Deltoid muscle of arm
Male 260+ lbs	22–25	1½"	Deltoid muscle of arm

\*Administer the second dose as close to the recommended interval (21 days) as possible. If the second dose is not administered within 6 weeks of the first dose, the series does not need to be restarted. Doses inadvertently administered less than 21 days apart do not need to be repeated.

†However, mRNA COVID-19 and other vaccines may be administered within a shorter period in situations where the benefits of vaccination are deemed to outweigh the potential unknown risks of vaccine coadministration (e.g., tetanus toxoid-containing vaccination as part of wound management, rabies vaccination for post-exposure prophylaxis, measles or hepatitis A vaccination during an outbreak) or to avoid barriers or delays to mRNA COVID-19 vaccination.

‡For the purpose of this guidance, 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.

†Alternatively, the anterolateral thigh also can be used.

†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).

02/11/2021 CS321570.H

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## Requirement to Give Vaccine:



- Epinephrine in prefilled autoinjector
- Antihistamine (Benadryl)
- Access to call 911



- First Aid Kit



- Stethoscope



- Blood pressure cuff



- Timing device for pulse



- Oxygen



- Pulse Oximeter



# Vaccine Allergic Reaction Kit:



- Epinephrine in prefilled autoinjector
- Antihistamine (Benadryl)
- Access to call 911



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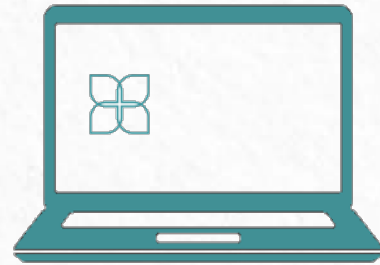
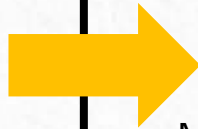


## Moving from Clinics to Continuity:



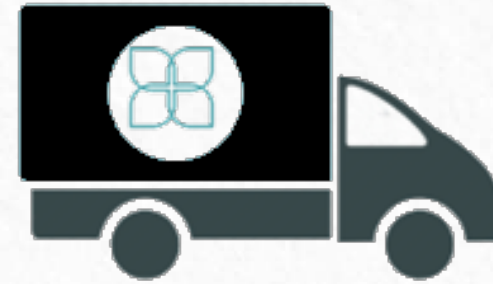
Activate

- Agreement
- Identify FC
- Identify Vaccinator
- FC Website Training
- Vaccinator Training
- Physician Order
- Standing Orders
- Life Enrichment List



Order

- Maintain patient's roster
- e-Consent
- EUA
- 2 Days in Advance
- Rx Verification



Receive



Vaccinate



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# Ordering the VaccPack:



My Patients

Upcoming Clinics

Vaccine Inventory

Manage Accounts

My Clinics

Help

Order Pfizer

Inventory Report Out

Patient Corrections

My Patients at

Total Number of Residents

22

Obtained (82%)

Refused (0%)

Needed (18%)

Stop - Do Not Give (0%)

Total Number of Staff

3

Obtained (67%)

Refused (0%)

Needed (33%)

Stop - Do Not Give (0%)

Vaccination Progress

27

1st Dose (11%)

None (44%)

Booster (44%)

Hope Village Patients

Search...

FILTER

EXPORT PATIENTS

ADD PATIENT

!	First Name	Last Name	Patient Type	First Dose	Booster Dose	Scheduled	Consent Status
	Another	Patient3	Resident	12/20/2020	12/23/2020	Yes	Obtained
	Cool	Guy	Resident	02/02/2021	02/26/2021	Yes	Obtained

# Ordering the VaccPack:



## Pfizer BioNTech VACCINE ORDER FORM

The survey will take approximately 4 minutes to complete.

\* Required

1. Servicing Pharmacy \*

- ☐ Consonus Pharmacy
- ☐ Community Pharmacy

2. Scheduled Date for Vaccinations: \*

Please input date in format of M/d/yyyy



3. I attest that Vaccine Administrators and I have read, viewed, and understand the training documents and videos shown under the "Training Materials" tab on the Consonus/Community Vaccine Administration Site (CVAS) which discuss the storage and handling, preparation and dilution, and administration of the Pfizer-BioNTech COVID-19 vaccine: \*

Link to Training Materials:

<https://confluence.marquiscompanies.com/display/KB/Vaccine+Prep+and+Administration+Library>

- ☐ I attest Vaccine Administrators and I have reviewed and understand the above training materials



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# Ordering the VaccPack:

Quantity:

- 1 vial = 6 doses
- Pfizer
- Maximum Order 4 vials

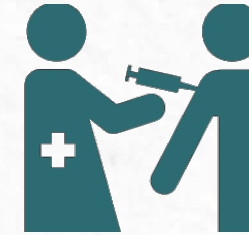
Roster Verified by Consonus



# Ordering the VaccPack: Key Points



Training Complete



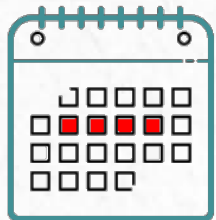
Vaccinator Scheduled



Order 2 days in Advance



Know State Phase Criteria



Shipped Monday – Friday Only

- Your clinics should be scheduled Tuesday -Saturday



Allergic Reaction Kit

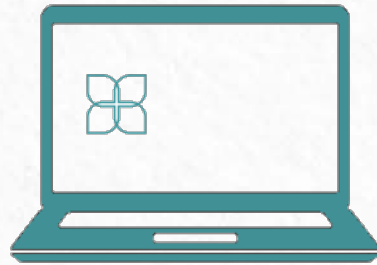


## Moving from Clinics to Continuity:



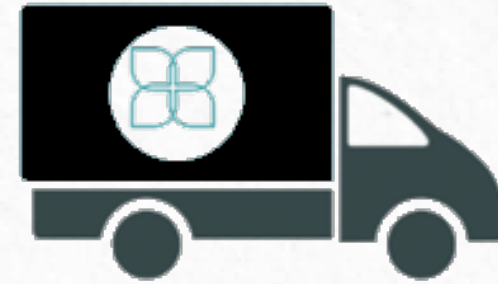
### Activate

- Agreement
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- Standing Orders
- Life Enrichment List



### Order

- Maintain patient's roster
- e-Consent
- EUA
- 2 Days in Advance
- Rx Verification



### Receive

- Consistent Location
- Secure
- Do Not Open



### Vaccinate



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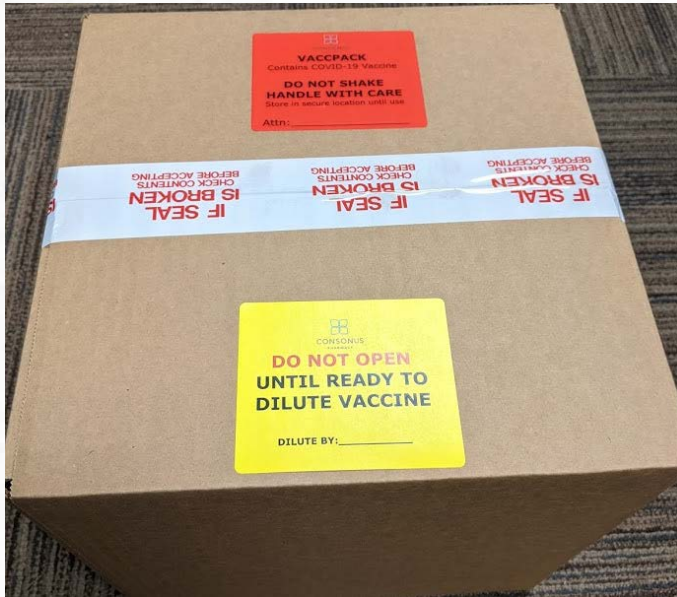
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# Introducing the VaccPack:



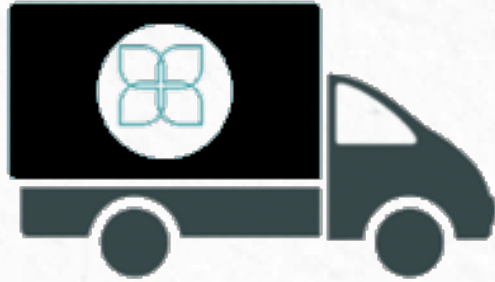
Quantity 6

Product is to stay in pharmacy packaging until vaccine is given.

- Vaccine
- Diluent – 0.9% NS Vial
- Vaccine Dilution Instructions

- Syringe (s) (7)
- Alcohol Swabs (16)
- CDC Cards (7)
- Band-Aides (7)

## Clinic Schedule:



## Routine Delivery:

- Used within 24 hours of receipt

## Moving from Clinics to Continuity:



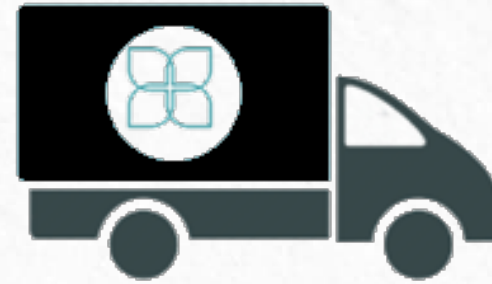
### Activate

- Agreement
- Identify FC
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- FC Website Training
- Vaccinator Training
- Physician Order
- Standing Orders
- Life Enrichment List



### Order

- Maintain patient's roster
- e-Consent
- EUA
- 2 Days in Advance
- Rx Verification



### Receive

- Central Location
- Secure
- Do Not Open



### Vaccinate

- Allergic Reaction Kit
- Vaccination Kit
- Screen
- Monitor
- Inventory Report



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# Screening Protocols Remain in Place

## Pre-Vaccination Form for Pfizer-BioNTech COVID-19 Vaccine



Information for Healthcare Professionals about the Pre-Vaccination Form for Pfizer-BioNTech COVID-19 Vaccine.

For additional information on COVID-19 vaccine recommendations, see:  
<https://www.cdc.gov/vaccines/covid-19/info-by-product/pfizer/clinical-considerations.html>  
For additional information on ACIP general recommendations, see:  
<https://www.cdc.gov/vaccines/hcp/acip-recs/general-recs/index.html>

### Are you feeling sick today?

There is no evidence that acute illness reduces vaccine efficacy or increases vaccine adverse events. However, as a precaution with moderate or severe acute illness, all vaccines should be delayed until the illness has improved. Mild illnesses (e.g., upper respiratory infections, diarrhea) are NOT contraindications to vaccination. Do not withhold vaccination if a person is taking antibiotics. Vaccination of persons with current SARS-CoV-2 infection should be deferred until the person has recovered from acute illness and they can discontinue isolation. While there is no minimum interval between infection and vaccination, current evidence suggests reinfection is uncommon in the 90 days after initial infection. Persons with documented acute SARS-CoV-2 infection in the preceding 90 days may delay vaccination until near the end of this period, if desired.

### Have you ever received a dose of COVID-19 vaccine?

Two doses of the same COVID-19 vaccine product are recommended. Check medical records, immunization information systems, and vaccination record cards to help determine the initial product received. Those who received a trial vaccine should consult with the trial sponsors to determine if it is feasible to receive additional doses.

Have you ever had a severe allergic reaction (e.g., anaphylaxis) to something? For example, a reaction for which you were treated with epinephrine or EpiPen®, or for which you had to go to the hospital? Allergic reactions, including severe allergic reactions, NOT related to vaccines or injectable therapies (e.g., food, pet, venom, environmental, or latex allergies; oral medications) are NOT a contraindication or precaution to vaccination with currently authorized COVID-19 vaccine. HOWEVER, individuals who have had severe allergic reactions to something, regardless of cause, should be observed for 30 minutes after vaccination. All other persons should be observed for 15 minutes.

### Was the severe allergic reaction after receiving a COVID-19 vaccine?

History of severe allergic reaction (e.g., anaphylaxis) to a previous dose or component of the COVID-19 vaccine product being offered is a contraindication to that COVID-19 vaccine.

### Was the severe allergic reaction after receiving another vaccine or another injectable medication?

A history of mild allergic reaction to a vaccine or injectable therapy is not a precaution to vaccination. History of severe allergic reaction (e.g., anaphylaxis) to another vaccine or a component of another vaccine OR anaphylactic reaction to any other injectable medication is a precaution to currently authorized COVID-19 vaccine. Vaccine may be given, but counsel patients about unknown risks of developing a severe allergic reaction and balance these risks against the benefits of vaccination. These individuals should be observed for 30 minutes after vaccination.

### Do you have a bleeding disorder or are you taking a blood thinner?

COVID-19 vaccine may be given to these patients, if a physician familiar with the patient's bleeding risk determines that the vaccine can be administered intramuscularly with reasonable safety. ACIP recommends the following technique for intramuscular vaccination in patients with bleeding disorders or taking blood thinners: a fine-gauge needle (23-gauge or smaller caliber) should be used for the vaccination, followed by firm pressure on the site, without rubbing, for at least 2 minutes.

### Have you received passive antibody therapy as treatment for COVID-19?

Based on the estimated half-life of monoclonal antibodies or convalescent plasma as part of COVID-19 treatment, as well as evidence suggesting that reinfection is uncommon in the 90 days after initial infection, vaccination should be deferred for at least 90 days, as a precautionary measure until additional information becomes available, to avoid interference of the antibody treatment with vaccine-induced immune responses.

### » Considerations

Immunocompromise is not a contraindication to current COVID-19 vaccine, including those with cancer, leukemia, HIV/AIDS and other immune system problems or taking medication that affects their immune systems. However, patients should be informed that the vaccine might be less effective than in someone who is immunocompetent.

Pregnancy is not a contraindication to current COVID-19 vaccine. While there are currently no available data on the safety of COVID-19 vaccines in pregnant people, studies and results are expected soon. Pregnant people may choose to get vaccinated. Observational data demonstrate that while the absolute risk is low, pregnant people with COVID-19 have an increased risk of severe illness.

Lactation is not a contraindication to current COVID-19 vaccine. Lactating people may choose to be vaccinated. There is no data available for lactating people on the effects of mRNA vaccines.

12/16/20

CS316204

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## Pre-Vaccination Form for Pfizer-BioNTech COVID-19 Vaccine



### For vaccine recipients:

The following questions will help us determine if there is any reason you should not get the COVID-19 vaccine today. If you answer "yes" to any question, it does not necessarily mean you should not be vaccinated. It just means additional questions may be asked. If a question is not clear, please ask your healthcare provider to explain it.

Patient Name \_\_\_\_\_

Age \_\_\_\_\_

	Yes	No	Don't know
1. Are you feeling sick today?			
2. Have you ever received a dose of COVID-19 vaccine? If yes, which vaccine product? <input type="checkbox"/> Pfizer <input type="checkbox"/> Another product _____			
3. Have you ever had a severe allergic reaction (e.g., anaphylaxis) to something? For example, a reaction for which you were treated with epinephrine or EpiPen®, or for which you had to go to the hospital?			
• Was the severe allergic reaction after receiving a COVID-19 vaccine?			
• Was the severe allergic reaction after receiving another vaccine or another injectable medication?			
4. Do you have a bleeding disorder or are you taking a blood thinner?			
5. Have you received passive antibody therapy as treatment for COVID-19?			

Form completed by \_\_\_\_\_ Date \_\_\_\_\_

Form reviewed by \_\_\_\_\_ Date \_\_\_\_\_

Adapted with appreciation from the Immunization Action Coalition (IAC) screening checklist

12/16/20

CS316204

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Consonus Healthcare  
[consonushealth.com](https://consonushealth.com)

Plan provided by  
Consonus Pharmacy



ROAD TO  
RECOVERY  
Ready, Set, Go

# Report Inventory:



My Patients

Upcoming Clinics

Vaccine Inventory

Manage Accounts

My Clinics

Help

Order Pfizer

Inventory Report Out

Patient Corrections

My Patients at

Total Number of Residents

22

Obtained (82%)

Refused (0%)

Needed (18%)

Stop - Do Not Give (0%)

Total Number of Staff

3

Obtained (67%)

Refused (0%)

Needed (33%)

Stop - Do Not Give (0%)

Vaccination Progress

27

1st Dose (11%)

None (44%)

Booster (44%)

Hope Village Patients

Search...

FILTER

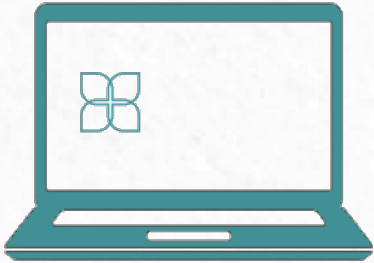
EXPORT PATIENTS

ADD PATIENT

!	First Name	Last Name	Patient Type	First Dose	Booster Dose	Scheduled	Consent Status
	Another	Patient3	Resident	12/20/2020	12/23/2020	Yes	Obtained
	Cool	Guy	Resident	02/02/2021	02/26/2021	Yes	Obtained



# Report Inventory:



## Post-Clinic Inventory Report Out

Facility Coordinator to complete the below Inventory Report

\* Required

1. Date of Vaccine Adminsitration \*

Please input date in format of M/d/yyyy



2. Name of Facility (Include ALF, SNF, MC in Name) \*

Enter your answer

3. Vaccine Manufacturer: \*

☐ Pfizer-BioNTech

☐ Moderna

4. Number of Vials Diluted: \*








# COVID Vaccines Authorized for Use:









- Pfizer
- Moderna
- Janssen

- 
- |                     |                           |
|---------------------|---------------------------|
| 1 Efficacy          | 4 Clinical Considerations |
| 2 Safety            | 5 Population              |
| 3 Contraindications | 6 VAERS Reporting         |

## Storage and Handling:

				
	Age	18 yo	16 yo	18 yo
	Between doses	28 days	21 Days	Single Dose
	Room Temp (unopened vials)	12 hours	2 hours	12 hours
	In use vials	6 hours after puncture	6 hours after dilution	2 hours- rm temp 6 hours - refrigerator

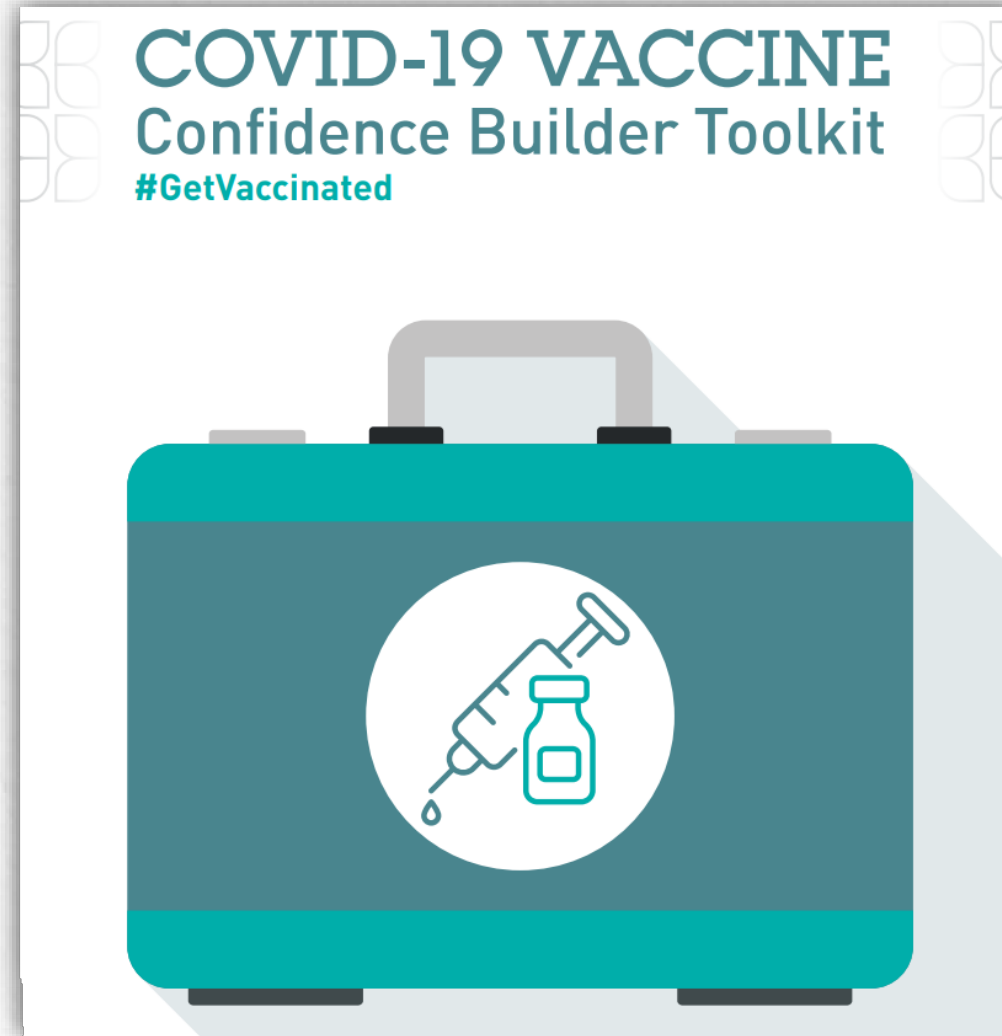
# Product Comparison:

				
	Dilution	None, ready to use	Dilute with saline	None, ready to use
	Doses/Vial	10	6	5
	Dose	0.5 ml IM	0.3ml IM	0.5 ml IM





#GetVaccinated:



## Best Case Schedule:



### Get Ready!

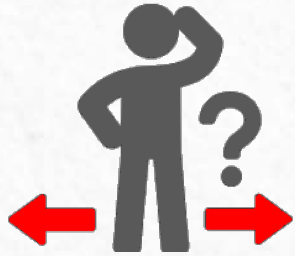
Week 1 – 2 Pilot facilities

Week 2 - 21 facilities

Week 3 – Continue SNF + 4 ALF's

Week 4 – All SNF Customers + Additional ALF's

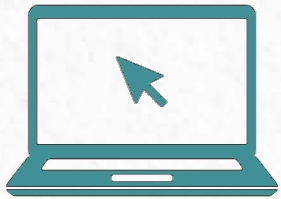
## What to Work on Now:



- Continue to address vaccine hesitancy
- Build patient rosters



Evaluate if this program is right for your community

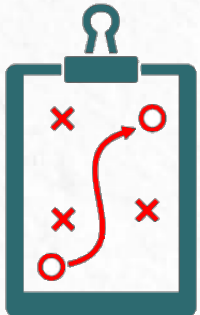


Activation Phase

- Sign Vaccinator Agreement



Standing orders



Get internal processes in place



Review new admit / staff data

## LTC Continuity Program:

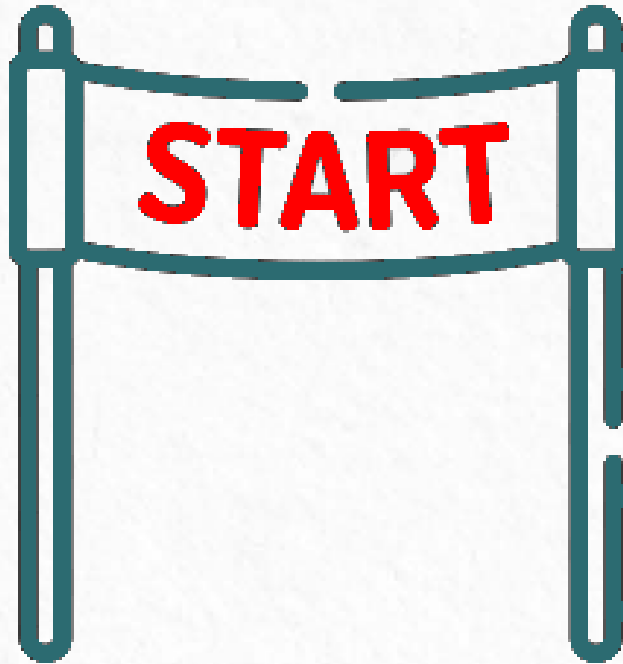


This program may not be for everyone. Other options may be:

- County public health
- Rural access partnerships
- Health systems in your area

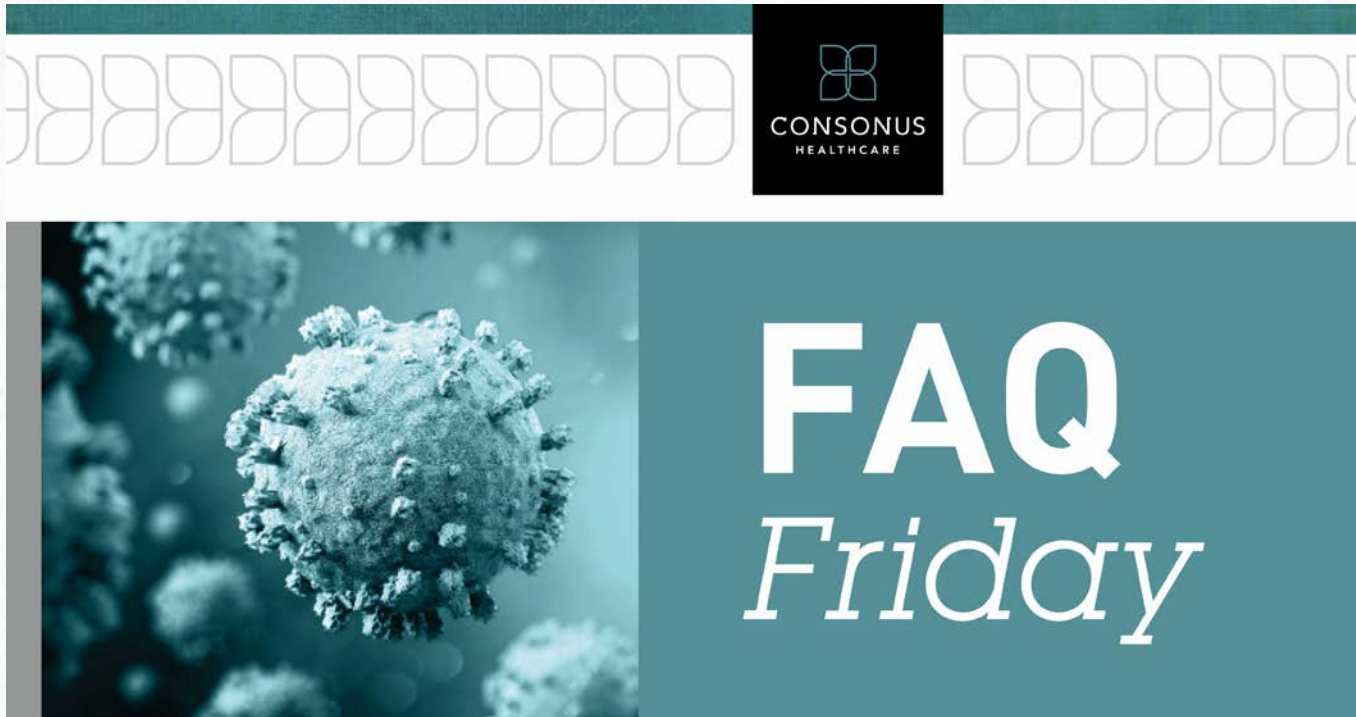


# When Do We Get Started?



- 1 Return your signed addendum
- 2 Validate access to Consonus Vaccine Application
- 3 Once the federal program is activated you will receive notification of acceptance.

# Questions?



- Check our website for the latest information
- FAQ Friday answering your commonly asked questions
- When we have information, you will hear from us