



Consonus Healthcare consonushealth.com

Plan provided by Consonus Pharmacy

Road to Recovery: Update









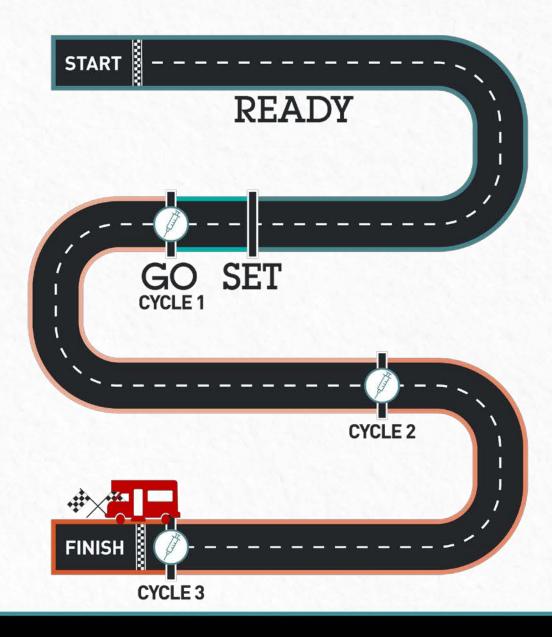


Information current as of 2.19.2021





Road to Recovery:







Current Consonus Stats:



Clinics Held

695

Total Vaccines to Date

48,018

Booster Doses

20,806



Resident Consent

87%

Staff Consent

72%

Done with 3rd Clinic

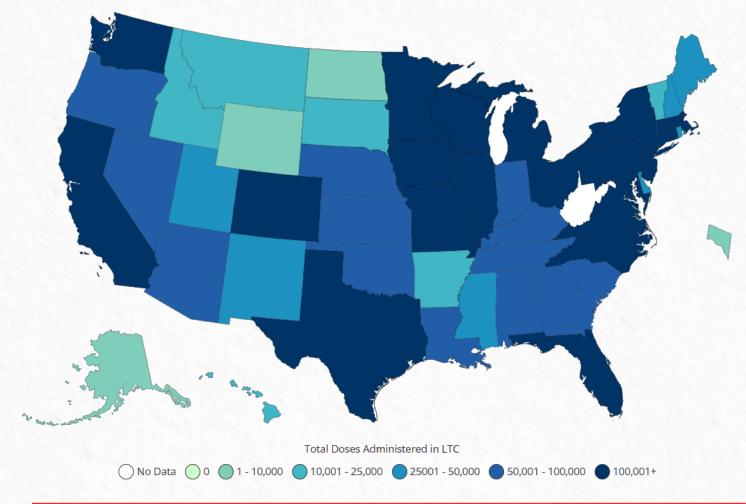
106







National Statistics



48% Decline in New Cases - After Clinic 1

LTC Pharmacy Partnership Program Subset

Total Number of Doses Administered in Long-Term Care Facilities

6,070,341

Number of People with 1 or More Doses in Long-Term Care Facilities

4,266,536

Number of People with 2 Doses in Long-Term Care Facilities

1,774,523

CDC | Data as of: Feb 17 2021 6:00am ET | Posted: Feb 17 2021 1:35PM ET





Vaccines & Immunizations

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











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 <u>Pharmacy Partnership for Long-Term Care (LTC) Program</u> 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

for LTC

Phase 2 Federal "Retail" Program:

Vaccine Continuity

Vaccine Continuity Program for LTC

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



Stakeholders



Everyone must work together





The Goal:



Maintain ≥ 70% 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





Our Shared Responsibility:



Manage Product



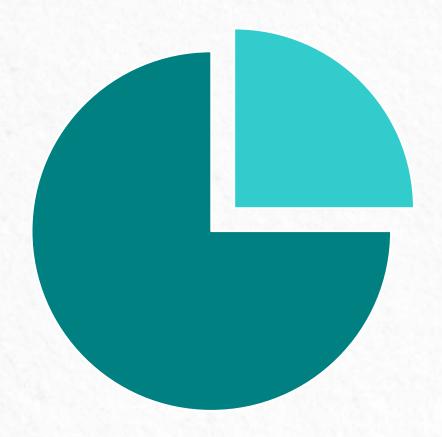
Manage Data



Manage Recipients



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





Product Similarities:



- mRNA technology
- Highly effective after second dose
- Not interchangeable



- SE/ ADE profile
- Precautions
- Contraindications
- VAERS reporting



- Clinical Screening
- 14 days- other vaccines
- 90- day deferral if received passive antibody therapy





Product Differences:

		moderna	Pizer
↑ 1 1 1 1 1 1 1 1 1 1	Age	18 yo	16 yo
	Between doses	28 days	21 Days
0	Dilution	None, ready to use	Dilute with saline
	Doses/Vial	10	6
	Dose	0.5 ml IM	0.3ml IM
00	Storage	Frozen	Ultra-cold





Storage and Handling:

	moderna	Pfizer
Thaw- room temp	1 hr	30 min- 2 hrs (dilute within 2 hrs)
Refrigerator	30 days	5 days
Room Temp (unopened vials)	12 hours	2 hours
In use vials	6 hours after puncture	6 hours after dilution





Clinical Screening and Standing Orders



Moderna COVID-19 Vaccine

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



Note: For more information/quidance, 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

· 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 mixedbrand series have been administered, no additional doses
- If the recipient has received 1 previous dose of Moderna COVID-19 Vaccine, a second dose of the same brand should be administered.
- o 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
- o 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.
- 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.

- » Severe allergic reaction (e.g., anaphylaxis) to a previous dose or component of either mRNA COVID-19 vaccine.
- » Immediate allergic reaction[±] 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).

- » 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.
- Prenare to administer the vaccine

Sex and Weight of Patient	Needle Gauge	Needle Length	Injection Site†
Female or male fewer than 130 lbs	22-25	% ⁵ −1"	Deltoid muscle of arm
Female or male 130–152 lbs	22-25	1"	Deltoid muscle of arm
Female 152–200 lbs	22-25	1-11/2"	Deltoid muscle of arm
Male 153–260 lbs	22-25	1-11/2"	Deltoid muscle of arm
Female 200+ lbs	22-25	11/2"	Deltoid muscle of arm
Male 260+ lbs	22-25	11/2"	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 situation:

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 aunid harriers or delays to mRNA COVID-19 vaccination

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*For the purpose of this guidance, an immediate alleroic reaction is urticaria, angioedema, respiratory distress (e.g., wheezing, stridor), or anaphylaxis that occur within 4 hours following exp vaccine or medication.

[†]Alternatively, the anterolateral thigh also can be used. ⁵Some experts recommend a 5/8-inch needle for men and women who weigh less 130 pounds. If used, skin must be stretched tightly (do not bunch subcutaneous tissue).



Pfizer-BioNTech COVID-19 Vaccine

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



Note: For more information/quidance, 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).

 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).

 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,
 Screen for contraindications and precautions. regardless of brand. If 2 doses of a same-brand or mixed-brand series have been administered, no additional doses are recommended.
- o If the recipient has received 1 previous dose of Pfizer-BioNTech COVID-19 Vaccine, the second dose of the same brand should be administered.
- o 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.
- o This vaccine is administered in a 2-dose series. Separate doses by at least 28 days, preferably within
- 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.5

- 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.
- - Contraindications:
 - Severe allergic reaction (e.g., anaphylaxis) to a previous dose or component of either mRNA COVID-19 vaccine.
 - Immediate allergic reaction* 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).

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†
Female or male fewer than 130 lbs	22-25	% 9 −1"	Deltoid muscle of arm
Female or male 130–152 lbs	22-25	1"	Deltoid muscle of arm
Female 152–200 lbs	22-25	1-11/2"	Deltoid muscle of arm
Male 153–260 lbs	22-25	1-11/2"	Deltoid muscle of arm
Female 200+ lbs	22-25	11/2"	Deltoid muscle of arm
Male 260+ lbs	22-25	11/2"	Deltoid muscle of arm

"Administer the second dose as close as possible to the recommended interval (21 days). If the second dose is not administered within 6 weeks of the first dose, the series does not need to be restarted. Second 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 old-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

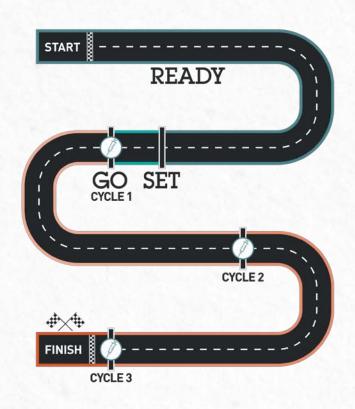
[†]Alternatively, the anterplateral thinh also can be used. 55ome experts recommend a 5/8-inch needle for men and women who weigh less 130 pounds. If used, skin must be stretched tightly (do not bunch subcutaneous tissue).



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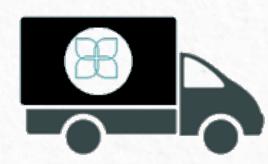


Moving from Clinics to Continuity:





Plan provided by Consonus Pharmacy





Ready, Set, Go

Introducing the VaccPack:

- Vaccine
- Syringe (s)
- Alcohol Swabs
- CDC Cards
- Band-Aides
- Diluent (if Pfizer)











Ordering the VaccPack:

- Build you roster of vaccine recipients in the Consonus vaccine app.
- 2 Select date vaccine is to be given
- Send order form to pharmacy indicating the date vaccine is to be given
- Prepare for receipt of vaccine on regular delivery the day prior to your clinic (shippers will receive same day)

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



Moderna COVID-19 VACCINE ORDER FORM

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none:	anc to Att	entron.	Fax:		
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	OVID-19 Ve				and corner and
	etains 50 doce				
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oderna O	OVID VACO	NE KIT (Moderna X	uccPack)		
	ntains:				
	Alcohol Swab Rand-Aid				
	COVID-CDC C	ands uringe for a diministration			
	and through		er of Individuals Receiving Vaccina	ition*	
		ion in the Consonus Va	ccination online application, please indicate th	e patients who ha	
vaccine, Entered	Signed	ntered into the applica Staff vs	tion, and have been scheduled to receive the Name	vaccination within	5 hours of reconstitution. 1st Dose vs 2nd Dose
into	Consen	Resident	Rame	500	1" Dose vs 2" Dose
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""This di	ite will be un If yo		ation**: ne delivery. Vaccine will be sent via regular del s or concerns, please contact Consonus Pl		



Requirement to Give Vaccine:



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



First Aid Kit



Stethoscope



Blood pressure cuff



Timing devise for pulse



Oxygen



Pulse Oximeter



Partnership Responsibilities:



- Consent
- EUA to patient
- Clinical screening
- Administration (reconstitution if required)
- Documentation
- Maintaining roster
- Ordering from pharmacy
- Being Accountable for product
- Having emergency supplies on hand



- Program Oversight
- Maintaining supply chain
- Appropriate packaging of VaccPack
- Delivering ordered supply to facility
- Validating roster
- Validate doses administered
- Reporting wasted doses
- Provide Consonus Vaccine APP
- Vaccine administration APP support
- Report mandatory data requirements
- Accountability to CDC
- Accountability state



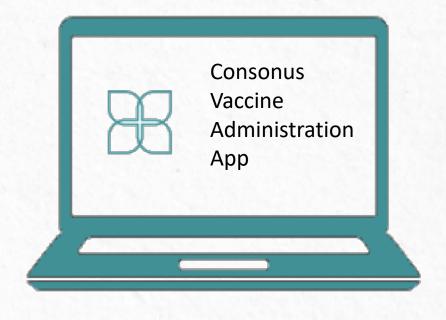


Documentation:

1 Consent

EUA for appropriate vaccine

Roster in Consonus Vaccine Administration App



Facility Vaccine Administrators

You will need a vaccinator for the continuity program.

Partnership Ideas

- Collaborate with established relationships
 - Home Health
 - Nurse Practitioner Groups
 - Skilled communities
 - Hospice organizations
 - Agency / per diem nursing
 - Retired Professionals (check state)







Considerations:



New Admission / Staff

- When were they vaccinated?
- What product?
- When are they due for second dose?

Vaccine Verification

- Immunization Information System
- CDC Cards



Process

- Documentation
- Internal policies
- Standing Orders per state
- •How do you move from clinic to continuity
- Maintain on call list to reduce waste (within state current phase)





What to Work on Now:



Continue to address vaccine hesitancy



Vaccine Verification



Get internal processes in place



Standing orders



Evaluate if this program is right for your community

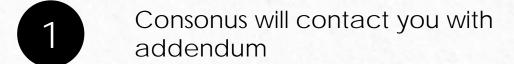


Review new admit / staff data



When Do We Get Started?





- Once you are activated you will receive order forms with acceptance verification.
- Access to Consonus Vaccine Application





Questions?



- Check our website for the latest information
- FAQ Friday answering your commonly asked questions
- When we have information, you will hear from us
- Next webinar: March 5th

