TEMPLATE ON LY

THIS TEMPLATE IS NOT LEGAL ADVICE. TALK TO YOUR ATTORNEY FOR GUIDANCE.

တာ်သမံသမ်း COVID-19 ကသံဉ်ဒီသဏ နီးတာ်အာဉ်လီးတုာ်လိဉ်

တဂ်ဂုံတကြိုလာတာ်ထာဖြိုင်အီးလာလံာ်ကွီဉ်ဒိအပူးအံး တာ်ကစူးကါအီးလာတာ်မေနီဉ်မေးယါလာနင်းနှုံဘဉ်လံ ကသံဉ်ဒိသေး(တဖဉ်) နှဉ်လီး တဂ်ဂုံတကြိုသာဉ်ပေးကသံဉ်ဒိသဒး(တဖဉ်) ဘဉ်သှဉ်သှဉ်တာ်ကနီးဟုဉ်လီးအီးဆုမံဉ်နံဉ်စိထဉ် (Minnesota) တာ်ဆုံးကသံဉ်ဒိသဒာ တဂ်ဂုံတကြိုတာ်ဘူးစဲ (MIIC) ယုာ်င်းပှာဟုဉ်လီးတာ်အိဉ်ဆူဉ်အိဉ်ချုတာ်မေးအဂၤ, ကွိတဖဉ်, တာ်အိဉ်ဆူဉ်အိဉ်ချုတ်ကျို, ငီးပှာလာအိဉ်င်းတာ်စိ တက်မီးအဂၤလာအိဉ်လာသဲစေးတာ်ဘူးအဖိလာ်ဒ်သီးကင်းနှုံဘဉ်အီးအင်္ဂါနှဉ်လီး. နမ္မာ်အိဉ်င်းတာ်သံကွာ်တမ်းမီးနှဉ်, ဝံသးစူးသံကွာ်န ကသံဉ်သရဉ် မှတမှာ ပူးဟုဉ်လီးတာ်အိဉ်ဆူဉ်အိဉ်ချုတာ်မေးအဂၤတက္နာ့ နမ္မာ်အိဉ်င်းတာ်သံကွာ်ဘဉ်ပေး MIIC နှဉ်, ကွာ်သယ်းဆူ MIIC င်း ပူးကမျာ်တဖဉ် (www.health.state.mn.us/people/immunize/miic/public.html) မှတမှာ ကိုးလီတဲစိဆု ၁-၈၀၀-၆၅၇-၃၉၇၀ တက္ခါး

တာ်ဟုဉ်လီးတာ်နှုံးဘျူး**ီးမူေါတဖဉ်လာတာ်ဟုဉ်စံ့အင်္ဂီ** တာ်အားပျဲပဆျာကဖွဲ့စရီဆူနတာ်အိဉ်ဆူဉ်အိဉ်ချုတာ်တိာ်ကျဲး မဲ့တမာ့် ခ်ီပနံဉ် ဇီးဒ်သီးကဒီးနှာ် တာ်ဟုဉ်စုတဘိုယ်အင်္ဂါနှဉ်လီး

ယဟ္နာ်ပြဲအခွဲးဆုပှာဟုဉ်လီးဆုဉ်ချတာ်မာစားအံး လာကဆျကနွဲစရီဆူ ယတာ်အိဉ်ဆုဉ်အိဉ်ချတာ်တိာ်ကျဲး မဲ့တမာ့် ပှာအဂလာအဟုဉ်စ့လာယချာ်စး, ဒီးလာကဒီးနှာ်ဘဉ်တာ်ဟုဉ်စ့တာ်ဟုဉ်ခွဲးတာ်နှာ်ဘူးအဂ်ီးနှဉ်လီး

တာ်ဆဲးကိုးတာ်ဂ်ာတာကြိုး ပူးလာအဆဲးတာ့ကသံဉ်ဒီသဇာ

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ပှာဆါအမံး (မံးကတာဂ်, ခီဉ်ထံး, ချဉ်သး)–

အီဉ်ဖျဲဉ်မုဂ်နံး–

သးနံဉိ–

လိတဲစိအခီဉ်ထံးတဖျာဉ်–

လိဂ်အိဉ်ဆီးထံး (ကွဲ မှတမှာ် လံာ်တားနိုဉ်က်)–

တုံ–

ဟိဉ်ကဝီးကိဂ်စဉ်–

မံး(ပ)နိုဉ်က်၊–

မိဂ်အမံး (မံးကတာဂ်, ခီဉ်ထံး, ချဉ်သး– သးနံဉ်မှာ်စုံနှာ် ၁၈ နံဉ်)–

မိဂ်အမံးကတာဂ်လာအပူးကွဲဉ် (သးနံဉ်မှာ်စနာ့် ၁၈ နံဉ်)–
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တၢ်ဟုဉ်စုတၢ်ဂ့ၢ်တၢ်ကြို

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ဟဲစိ၁်နတာ်အှဉ်ကီးခးကဲ့ငီးနာ.
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ပျယ**့်**ဦဂ်ါခ်ီဦထီးတါအူဦကီး–

ဖီလစံဉ်/တာ်အုဉ်သးနိုဉ်ဂံ၊ ID/ကရာဖိနိုဉ်ဂံ၊ – ကၡုံအနိုဉ်ဂံ၊ –

ပူးဟူဉ်တါအူဉ်ကီးခံဖုတဖု–

ဖိလစ်ဉ်တာ်အူဉ်သးနိုဉ်ဂံ၊ ID/ကရာဖိနိုဉ်ဂံ၊ – က၍အနိုဉ်ဂံ၊ –

ပူးဖီဉ်ဖီလစံဉ်, မှု်လီးဆီလိဉ်အသးဖီးပူလာအဆုံးကသံဉ်ဖီသစာ–

ψ̂\-
အိန်ဖျဲန်ဖုံနံး–
ခ်ီပနံ်ာ်တာ်ဟူဉ်စ္–
ခ်ီပနံ်ာ်အမံ၊–
ပူးလာအဆုံးကသံဉ်ဒီသရာမှါတအိုဉ်ဒီးတါအှဉ်ကီးနှဉ် တိုးနိုဉ်ဖွဲ့အုံးတက္ခါ

တာ်အာဉ်လီးတူာ်လိာ်

ခံဖိုလာတာ်ဆုံးလီးမုံးလာလာအားအယိ, ယနာ်ဟာ, ဟာသူဉ်ဟာသး, အာဉ်လီးတူဂ်လိဉ်, နီးသူးလီးပလိဉ်လာ–

- ယဒီးနှုံဘဉ်ဒီးဖူး မှတမှာ် တာ်တဲနာ်ဟာ်ယာလာ ဂဲ့ာဂ်ီးဆူတာ်စူးကါတာ်ဆာဉ်လီးဟုဉ်ခွဲးတာ်မူးဆသီးနှီးနှီးဆကဘျုံး (Emergency Use Authorization Fact Sheet) လာကသံဉ်ဒီသဇာလာဆပ်ာတွဲထိဉ်အခံဆုံးအင်္ဂါလံ— [ဖြနင်္ခလီးကသံဉ်ဒီသဇာမုံးပနိုဉ်အမုံး]တက္ပါ.
- ယအိဉ်တုံးေတာ်ခွဲးတာယာလာကသံကွာ်တာသံကွာ်တဖဉ်လာအိဉ်ေးတာ်စုံးဆာလာအမာမံသးယာ, ေးယနာ်ဟာ်တာန္ခ်ာဘူးတဖဉ်ေးတာ်ဘဉ်ယိဉ်ဘဉ်ဘီ လာအဘဉ်ဃးမီး COVID-19 ကသံဉ်ဖိသဏလာတာပတ်၍ထိဉ်အီးနှဉ်လီး.
- ယအာဉ်လီးတူဂ်လိဉ်လ၊ကဆဲး COVID-19 ကသံဉ်နီသရလ၊ယနီးကစာ မ္ ϕ မှာ ပူးအမံးလ၊ထးအံးအဂ်ိုန္ဉ်ာလီး

ပူးဆါ မ့တမ့်၊ မိါ်ပါ/ပူးကွက်ထွဲတာ် ဆဲးလီးမံး–	
မှါနံး/	

တာ်အိုဉ်ဆှဉ်အိုဉ်ချ အတာ်စံဉ်စီး

နမ့်၊စံးဆ၊ မ့၊ ဆူတာ်သံကွာ်တဖဉ်အံးတမံးမံးနှဉ်, ပူးလာကဆဲးနာကသံဉ်ဒီသဇာတဂၢ ကလိဉ်ဘဉ်အါထီဉ်တာ်က်တာ်ကြိုးလာနအိဉ်တချုး လာကဆဲးနှာ်နာကသံဉ်ဒီသဇာလီ၊ –

မ့်	တမ့်	တသ္5ညါဘဉ်	တာ်သံကွာ်
ခ်ု	တမ့်		မှန်မှာ်ပြာလာအသးနံဉ်ဘဉ်ဘိျီးဘဉ်ဒါလာကဆဲး COVID-19 ကသံဉ်ဒီသရာအဂ်ိုခါ. • Pfizer-BioNTech—အသးကဘဉ်မှာ ၆ လါ မှတမှာ ပြုနှာ်အနှဉ်. • Moderna ကသံဉ်ဒီသရ—အသးကဘဉ်မှာ ၆ လါ မှတမှာ ပြုနှာ်အနှဉ်. • Novavax ကသံဉ်ဒီသရ— နသးနံဉ်ကဘဉ်မှာ ၁၂ နံဉ်ဒီးဆူအဖီခိဉ်နှဉ်လီး.
မွ်	တမ်ု	တသ့ဉ်ညါဘဉ်	မ့်၊နအို််င်းမှုတ်၊ဘဉ်က်, နိုုံခိဂ်ကြီဆာတလာတပြဲတာအိုဉ်သးအဂၤတဖဉ်, မ့တမ့်၊ အီ နှီါခိဂ်ကြီဆာ တလာတပြဲးကသံဉ်ကသိ မ့တမ့်၊ တာ်ကူစါယါဘျါတဖဉ်ပေါ.
မွ်	တဓံ့၊	တသ့ဉ်ညါဘဉ်	ကသံဉ်အတၢ်ဒိဘဉ်လ၊အဒိဉ်အမှာ် (အဒိ, anaphylaxis) စဲဆဲးကသံဉ်လ၊အပူးကွံာ်တဖျဉ်မှတမှ1 ဘဉ်ဃးဒီး COVID-19 ကသံဉ်ဒီသရတမံးမံးခါ.
မွ်	တမ့်	တသ္5်ညါဘဉ်	မှန်အို်င်းတါတဘဉ်လိဂ်င်းသွံဉ်တစ်ဆာသတုဂ်ကလာ်, တမှါနးနားကလာ် (၄ နဉ်ရံဉ်အတိါ်ပူး) င်းတါဆဲး COVID- 19 ကသံဉ်င်္ဒိသင်္ဂလာအပူးကွဲဂ်တဘို မဲ့တမှါ သဲ့ဉ်ညါ (အို််င်္ဂီး) တါတဘဉ်လိဂ်င်းတါပဉ်ယုဂ် လာကသံဉ်ဆဲးဒိသင်္ခအပူး မဲ့တမှါ တါပဉ်ယုဂ်အဂၤတမ်းလာ်လာ်အပူးခါ.
မှ်	တမ့်	တသူဉ်ညါဘဉ်	မ့်၊ နအိဉ်ဒီးတာ်တဘဉ်လိာ်ဒီးသွှဲဉ်တာ်ခီဆာသတု်ကလာ်ဒီး ကသံဉ်ဒာသဒာဆဂၤတမ်းလာ်လာ (တမှာ –COVID-19) မှာ်တာ်ကူစါယါဘျါလာအိဉ်ဒီးတာ်ဆဲးကသံဉ် (အဒီဆဲးနှာ်ကသံဉ်လာညဉ်လာ်အပူး (ညဉ်ထုဉ်ညဉ်မျာ်), လာ သွှဲဉ်ကျိုးအပူး (သွှဲဉ်ကျိုး), မှတမှာ် ဖီးဘဲ့ဉ်ထံးရှူအသိအကထာ (ဖီးဘဲ့ဉ်အဖီလာ်) ဧါ. တပဉ်ယှာ်ဒီးတာ်ဆဲးတာ်တ ဘဉ်လိာ်ဒီးသွဲ့ဉ်ဘဉ်.
မ့်	တမ့်	တသ့ဉ်ညါဘဉ်	တနံးအံးနဆိမိဉ်လ၊နဆိးက္ ဓါ ့

မှ်	တမ့်	တသ္နဉ်ညါဘဉ်	တာ်သံကွာ်
မ့်	တမ့်	တသ့ဉ်ညါဘဉ်	အိဉ်ငီးတၢ်စံဉ်စီးဘဉ်ယးငီး တာ်ညီးကဖိထိဉ်တာ်ဆါတဘဉ်လိာ်ဘဉ်စးအကျိုအကျဲအကလှာ်ကလှာ် စဲဘဉ် COVID-19 တာ်ဆါဝံးအလိၢိခံ၏.
ဗို	တမ်ု	တသ့ဉ်ညါဘဉ်	မ့်၊နအို််ငီးတါစံဉ်စီးလ၊ သးဖျဉ်အညဉ်ထူဉ်ညီး (myocarditis) မ့တမ့်၊ သးဖျဉ်သလီညီး (pericarditis) ဖဲနဆဲးဝဲငဉ် COVID-19 ကသံဉ်ငီသရလာအမ့်၊ Moderna, Pfizer-BioNTech မှတမ့်၊ Novavax ကသံဉ်ငီသရဝံးအလို်ခံ၏.
မွ်	သရုံ	တသ့ဉ်ညါဘဉ်	မှ1်နဆဲးတ္1်လံဝဲဧဉ်ကသံဉ်ဒီသဒၢအဂၤတမံးမံးဖဲအပူးကွံဉ် 4 နွံအတိ1်ပူးဓါ
မွ်	တမ်ု	တသ့ဉ်ညါဘဉ်	တာ်မာအိုဉ်လာတာ်ပဉ်အိုဉ်လီးဖြိုဉ်သးနီးပှာက မှာ်လာဘဉ်သင်္ကျီနီးပှာအကတကလာအိုဉ်နီး COVID-19 တာ်ဆါလာတာ်ပဉ်က်ကြာအီးခါ.
မှ်	တမ့်	တဘဉ်ထွဲဘဉ်	မ့်၊ နဆဲးဘဉ်တ့ဂ် COVID-19 ကသံဉ်ဇိသဇာတဘျိတဘျိမါ. မှု်ဆဲးတ့ဂ်န္နဉ်, ကွဲးလီးကသံဉ်ဇိသဇာမံးပနိုဉ်ဇီးမုဂ်နံးလာနဆဲးအီးတက္ဂ် –

တဘဉ်ကွဲးတါလ၊လံဉ်ကြိုးအံုးအဖီလဉ်ဘဉ်

Vaccine information

COVID-19 Vaccine Presentation ¹	EUA Fact Sheet Date	Route ²	Manufacturer ³	Lot Number	Admin Site ⁴	Person Admin ⁵
COVID-19 Comirnaty (Pfizer) 12 years and older (gray cap), 0.3 mL		IM	PFR			
COVID-19 Pfizer 5-11 years (blue cap), 0.3 mL		IM	PFR			
COVID-19 Pfizer 6 months-4 years (yellow cap), 0.3 mL		IM	PFR			
COVID-19 Spikevax (Moderna) 12 years and older (blue cap/blue label)		IM	MOD			
COVID-19 Moderna 6-11 years (blue cap/green label), 0.25 mL		IM	MOD			
COVID-19 (Novavax)		IM	NVX			

1.	COVID-19 Vaccine Presentation =	lists specific	product name (e.g., Pfizer,	. Moderna,	Novavax, e	tc.)
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- 2. Route: IM = Intramuscular
- 3. **Manufacturer:** MOD = Moderna, PFR = Pfizer, NVX= Novavax
- 4. **Site Vaccine Given:** LD = Left Deltoid, RD = Right Deltoid, LT = Left Thigh, RT = Right Thigh
- 5. **Signature or initials of person administering vaccine:** Can be used if more than one person is administering vaccines.

Signature and title of person administering vaccine: _	
Date administered://	

Information for health care professionals about the pre-vaccination form for COVID-19 vaccine

[For health care providers, not for the patient]

This information is derived from the <u>CDC</u>: <u>Use of COVID-19 Vaccines in the United States</u> <u>(www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html)</u>. We will reference this document as CDC's Interim Clinical Considerations below and note specific sections where information can be found.

Age

Follow recommendations for vaccine administration to authorized age groups found under each vaccine product's emergency use authorization (EUA) or package insert. For Moderna, Novavax and Pfizer-BioNTech COVID-19 vaccine primary series doses, an 8-week interval is suggested between dose one and two for immunocompetent people 6 months to 64 years of age, and especially males 12-39 years.

Immediate allergic reaction

An immediate allergic reaction to a vaccine or medication is defined as any hypersensitivity-related signs or symptoms such as urticaria (hives), angioedema (painless swelling under the skin, often happens with hives), respiratory distress (e.g., wheezing, stridor), or anaphylaxis that occur within four hours following administration.

Have you had a severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a component of the COVID-19 vaccine?

CDC considers this to be a contraindication to vaccination with COVID-19 vaccines. People with an allergy-related contraindication to one type of COVID-19 vaccine have a contraindication or precaution to the other types of COVID-19 vaccines. For additional details, refer to the following sections of CDC's Interim Clinical Considerations: Contraindications and precautions and Appendix E: Triage of people with a history of allergies or allergic reactions. For a full list of ingredients included in COVID-19 vaccines, refer to FDA: COVID-19 Vaccines (www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/covid-19-vaccines) and CDC: U.S. COVID-19 Vaccine Product Information (www.cdc.gov/vaccines/covid-19/info-by-product/index.html).

Have you had an immediate, non-severe allergic reaction to a previous dose or known (diagnosed) allergy to a component of the COVID-19 vaccine or any of its ingredients?

CDC considers this to be a precaution to vaccination with COVID-19 vaccines. Non-severe allergic reactions may include urticaria (hives) beyond the injection site and angioedema (visible swelling) involving lips, facial skin, or skin in other locations. Angioedema affecting the airway (e.g., tongue, uvula, or larynx) would be a **severe** allergic reaction. For additional details, refer to the following sections of CDC's Interim Clinical Considerations: Contraindications and precautions and Appendix E: Triage of people with a history of allergies or allergic reactions. For a full list of ingredients included in COVID-19 vaccines, refer to COVID-19 vaccine-specific <u>FDA:</u> COVID-19 Vaccines and <u>U.S. COVID-19 Vaccine Product Information</u>.

Immediate allergic reaction to any other vaccines (non-COVID-19) or injectable therapy (intramuscular, intravenous, or subcutaneous)? Does not include allergy shots.

People with a history of an immediate allergic reaction to a non-COVID-19 vaccine or injectable therapy that contains multiple components, one or more of which is a component of a COVID-19 vaccine, but it is unknown which component elicited the allergic reaction, have a precaution to vaccination with that COVID-19 vaccine. These people may benefit from consultation with an allergist-immunologist who can perform a more detailed risk assessment for COVID-19 vaccine receipt and possibly allergy testing. For additional details, refer to the following

sections of CDC's Interim Clinical Considerations: Contraindications and precautions and Appendix E: Triage of people with a history of allergies or allergic reactions.

Are you feeling sick today?

There is no evidence that someone who is sick when vaccinated will decrease the vaccine's effectiveness or increase vaccine adverse events. If a person has COVID-19 symptoms, they should isolate following current guidelines, get tested, and if necessary, seek medical care. As a precaution, when someone is moderately to severely ill, all vaccines should be delayed until the illness has improved. A person who is mildly ill (e.g., diarrhea, upper respiratory infection, etc.), can still receive a vaccine, including people who are taking an antibiotic.

People should be offered vaccination regardless of their history of symptomatic or asymptomatic COVID-19 infection, including people with prolonged post-COVID-19 symptoms. Vaccination of people with known current COVID-19 infection should be deferred until at least the person has recovered from the acute illness (if the person had symptoms) and they have met criteria to discontinue isolation. This recommendation applies to people who experience COVID-19 infection before receiving any COVID-19 vaccine dose. For details, including additional information on delaying vaccine doses, refer to the following section of CDC's Interim Clinical Considerations: COVID-19 vaccination and SARS-COV-2 infection.

Have a history of Multisystem Inflammatory Syndrome after SARS-CoV-2 infection?

Given the lack of data on the safety of COVID-19 vaccines in people with a history of MIS-C and MIS-A, a conversation between the patient, their guardian(s), and their clinical team or a specialist (e.g., specialist in infectious diseases, rheumatology, or cardiology) is strongly encouraged to assist with decisions about the use of COVID-19 vaccines. Additional details can be found in the following section of CDC's Interim Clinical Considerations: COVID-19 vaccination and MIS-C and MIS-A section.

Have a history of myocarditis or pericarditis after a previous dose of Moderna, Pfizer-BioNTech, or Novavax COVID-19 vaccine?

CDC considers this to be a precaution to vaccination with mRNA COVID-19 vaccines (i.e., Moderna or Pfizer-BioNTech) or Novavax COVID-19 vaccine. Refer to their primary care provider to receive an assessment of their current health condition and assessment of individual benefits and risks. For more details, refer to the following section of CDC's Interim Clinical Considerations: COVID-19 vaccination and myocarditis and pericarditis.

Have you had any other vaccinations in the last 4 weeks?

Because of the observed risk for myocarditis after receipt of ACAM2000 orthopoxvirus vaccine and mRNA (i.e., Moderna and Pfizer-BioNTech) and Novavax COVID-19 vaccines and the unknown risk for myocarditis after JYNNEOS, people, particularly adolescent or young adult males, might consider waiting 4 weeks after orthopoxvirus vaccination (either JYNNEOS or ACAM2000) before receiving a Moderna, Novavax, or Pfizer-BioNTech COVID-19 vaccine. However, if an orthopoxvirus vaccine is recommended for prophylaxis in the setting of an outbreak, orthopoxvirus vaccination should not be delayed because of recent receipt of a Moderna, Novavax, or Pfizer-BioNTech COVID-19 vaccine; no minimum interval between COVID-19 vaccination with these vaccines and orthopoxvirus vaccination is necessary.

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

Refer to the following sections of CDC's Interim Clinical Considerations: COVID-19 vaccination overview and timing, spacing and interchangeability. Verify a person's age, what vaccine they have received, and the date(s) of prior dose(s) to assure the correct vaccine product and dose interval is used.

Other considerations

Chronic health condition – is not a contraindication or precaution for vaccination.

- Immunocompromised conditions (e.g., HIV infection, immunosuppressive medications, or therapies, etc.) immunocompromised people age 6 months and older should receive a COVID-19 vaccine series as soon as possible. They should be counseled regarding the potential for reduced immune responses and that the vaccine may not fully protect them. People need to continue to follow current guidance to protect themselves.
 - Moderately or severely immunocompromised Because the immune response following COVID-19 vaccination may differ in moderately or severely immunocompromised people, CDC has specific guidance for this population. For more details refer to the following sections of the CDC's Interim Clinical Considerations for Use of COVID-19 Vaccines: COVID-19 Vaccines, Recommendations, and Schedule for People who are moderately or severely immunocompromised.
- **Bleeding disorder or are taking a blood thinner** recommended to use a fine-gauge needle (23 gauge or smaller) for the vaccination, followed by firm pressure on the site, without rubbing, for at least 2 minutes.
- **Dermal filler(s)** advise to contact their health care provider for evaluation if they develop swelling at or near the site of dermal filler following vaccination. For additional details, refer to the following section of CDC's Interim Clinical Considerations: Special Populations.
- Pregnant Both CDC and ACOG urge that pregnant people be vaccinated. Pregnant and recently pregnant
 people with COVID-19 are at increased risk for severe illness when compared with non-pregnant people. Early
 data supports that vaccination is well-tolerated and elicits a protective immune response. For details, refer to
 the following section of CDC's Interim Clinical Considerations: Consideration involving pregnancy, lactation,
 and fertility.
- Passive antibody therapy for prevention or treatment for COVID-19 COVID-19 vaccination can be given at any time following receipt of antibody products as part of COVID-19 treatment, post-exposure prophylaxis, or pre-exposure prophylaxis once the isolation or quarantine period has been completed. For details, refer to the following section of CDC's Interim Clinical Considerations: COVID-19 vaccination and SARS-CoV-2 infection.
- COVID-19 vaccines and myocarditis and pericarditis Ongoing safety monitoring of the mRNA and Novavax COVID-19 vaccines has found increased risks of myocarditis and pericarditis, predominantly in males 12-39 years of age within the first week of receiving the second dose.

An 8-week interval between the first and second doses of an Moderna, Pfizer-BioNTech, and Novavax COVID-19 vaccine primary series may be optimal for some people ages 6 months—64 years, especially for males ages 12—39 years, as it may reduce the small risk of myocarditis/pericarditis associated with mRNA and Novavax COVID-19 vaccines.

Clinicians should consult the following section of CDC's Interim Clinical Considerations for Use of COVID-19 Vaccines: Safety considerations for COVID-19 vaccines or the CDC: Clinical Considerations:
CDC: Clinical Considerations:
Mttps://www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html#myocarditis-pericarditis when deciding whether to administer aCOVID-19 vaccine to someone with a history of myocarditis or pericarditis or when a patient presents with symptoms of myocarditis or pericarditis.

Find EUA fact sheets for health care providers and recipients and caregivers at <u>FDA: COVID-19 Vaccines</u> (www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/covid-19-vaccines).