

COVID – 19 Vaccine Acknowledgement and Consent Form

Moderna COVID-19 Vaccine FIRST DOSE

Recipient Information (Please Print Clearly)

Last Name:	First Name:	Date of Birth:			
Home Address:		Phone #:			
City:	State:	Zip:			
Ethnicity:					
☐ Black/African-American ☐ Caucasian ☐ Asian ☐ Native-American ☐ Other:					

The following questions will help us determine whether you can receive the COVID-19 vaccine. If you answer 'yes' to any question, it does not necessarily mean that you should not be vaccinated. It just means additional questions must be asked. If a question is not clear, please ask a staff member for further explanation:

	YES	NO	N/A
Are you sick today?			
Do you have a history of severe allergies?			
Have you ever had a serious reaction after receiving a vaccination,	П	П	П
including the first dose of the COVID-19 vaccine?			
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?			
Have you ever had a serious reaction after receiving a vaccine?			
Do you have a bleeding disorder or are you on a blood thinner?			
Are you immunocompromised or on a medication that affects your		П	П
immune system?			
For Women: Are you pregnant or planning to become pregnant?			
For Women: Are you breastfeeding?			
Have you received any vaccinations (other than the first dose of the		П	П
COVID-19 vaccination) in the past 2 weeks?			
Have you previously been diagnosed with COVID-19 and were treated			
with monoclonal antibodies in the last 90 days?			

I understand that the COVID-19 vaccine I will receive today requires two (2) doses from the same manufacturer to be fully effective. I understand I must return in 28 days to receive a second dose of the vaccine.



I consent to administration of the Moderna COVID-19 vaccination and acknowledge and agree with the following statements:

Initial before each statement (or put N/A if not applicable)

- I have received the Emergency Use Authorization (EUA) Fact Sheet for Recipients and Caregivers for the COVID-19 vaccine (the 'Fact Sheet').
- I have read the 'Fact Sheet' or had it read to me.
- The U.S. Food and Drug Administration (FDA) has authorized emergency use of the Moderna COVID-19 vaccine, which is not an FDA-approved vaccine. At this time, there is no FDA approved vaccine to prevent COVID-19.
- I understand the known and potential risks and benefits to this COVID-19 vaccine and the extent to which such benefits and risks are unknown.
- I acknowledge that I have the option to refuse vaccination and have been informed of any available alternatives to this COVID-19 vaccine and the risks and benefits of available alternatives.
- Recipients who are Pregnant or Breastfeeding: Pregnant and breast feeding persons were NOT included in the clinical trials for the COVID-19 vaccine. I have discussed the potential risks of COVID-19 infection versus the risk of vaccination with my healthcare provider and have made the informed decision to receive the COVID-19 vaccine.
- I understand that it is recommended that I remain at the vaccination clinic for fifteen (15) minutes following administration, but those with history of anaphylaxis should be monitored for thirty (30) minutes post vaccination to ensure that I have no immediate adverse reaction to the vaccine.
- I acknowledge that I have received information on V-Safe, a voluntary smartphone based tool operated by the Centers for Disease Control and Prevention (CDC). Through V-Safe, vaccine recipients can report any side effects of the vaccine to the CDC. This information helps CDC monitor the safety of the COVID-19 Vaccines in near real time.
- I have had the opportunity to ask questions regarding the administration of the Moderna COVID-19 vaccination which have been answered to my satisfaction.

If you experience an adverse reaction to the COVID-19 vaccine, please contact your primary care provider or present to the nearest emergency department. If you are experiencing a medical emergency, call 911.

Signature of Recipient/Authorized Representative:	Date:		
Print:			
If signed by Authorized Representative, please state relationship to Recipient:			



FOR CLINIC USE ONLY

Patient Name:		Date of Birth:			
Vaccine Administrator	(Print Name):				
Administration Date:	ministration Date:		Date Fact Sheet Provided:		
Scheduled date of 2 nd vaccination:					
Manufacturer	Lot Number	Expiration Date	Site of Administration		
			☐ Left Arm		
			□ Right Arm		
Monitoring period	od completed and no a	adverse reaction noted.			
☐ Recipient declined monitoring period. Waiver Completed.					
☐ Adverse reaction noted:					
Signature of Ohserver					

The COVID-19 Acknowledgement and Consent Form and COVID-19 Post-Vaccination Monitoring Period Waiver (if applicable) must be uploaded to the patient/employee chart.