## http://www.regentmedicalcare.com/for-patients/

## Consent - COVID-19 Immunization Screening and Consent Form - 6 Months and Older

Name: Last:		First:	Middle Initial:	Middle Initial: Temperature:			
Date of Birth:		Phone Number: ( )	SS#	QB ID#			
Address: Apt #:							
City: State:		State:	Zip:				
Sex (Gender assigned at birth) Race		Race			Ethnicity		
☐ Female		☐ American Indian orAlaska Native ☐ Native Hawai	iian or other □ Other Asian □ Unknown	n Hispanic or Latino			
☐ Male		☐ Asian ☐ Pacific Islan	der		☐ Not Hispanic or Latino		
		☐ Black or African American ☐ White	$\square$ Other Pacific Islander		□ Unknown		
Parent/Guardian/ Surrogate (if applicable, pleas		(if applicable, please print)	Clinic Where Vaccine is Administered				
Pri	Primary Care Physician Address Phone Number						
	Screening Questionnaire						
1.	Are you feeling sick too	day?		□ Yes	□ No		
2.	Have you ever received a dose of Covid 19 Vaccine? If yes, which vaccine product?			□ Yes	□ No	□ Unknown	
	o Pfizer o Moderna o Janssen (Johnson & Johnson) o Other product:						
3.	3. Have you ever had an allergic reaction to: (This would include a severe allergic reaction [e.g., anaphylaxis] that required treatment with epinephrine or EpiPen® or that caused you to go to the hospital. It would also include an allergic reaction that occurred within 4 hours that caused hives, swelling, or respiratory distress, including wheezing.)						
	•	of the COVID-19 vaccine, including polyethylene glyco ves and preparations for colonoscopy procedures	l (PEG), which is found in some medications,	□ Yes	□ No	□ Unknown	
	<ul> <li>Polysorbate</li> </ul>			□ Yes	□ No	□ Unknown	
	A previous do	ose of COVID-19 vaccine		□ Yes	□ No	□ Unknown	
4.	Have you ever had and an allergic reaction to another vaccine other than COVID 19 or any injectable medication? (This					□ Unknown	
	would include a severe allergic reaction [e.g., anaphylaxis] that required treatment with epinephrine or EpiPen® or that caused						
	,	. It would also include an allergic reaction that occurre	d within 4 hours that caused hives, swelling, or				
_	respiratory distress, incl	<u> </u>			<u> </u>		
5.	•	evere allergic reaction (e.g., anaphylaxis) to something on the or injectable medication? This would include food, processed in the control of the control o	·	□ Yes	□ No	□ Unknown	
6.		ny vaccine in the last 14 days?	et, environmental, or oral medication allergies.	□ Yes	□ No	□ Unknown	
7.		positive test for COVID-19 or has a doctor ever told you	that you had COVID-19?	□ Yes	□ No	□ Unknown	
8.		assive antibody therapy (monoclonal antibodies or convi	· · · · · · · · · · · · · · · · · · ·	□ Yes	□ No	□ Unknown	
9.		ned immune system caused by something such as HIV in	•	□ Yes	□ No	□ Unknown	
٥.	immunosuppressive (			cs		_ omenown	
10.	Do you have a bleed	ling disorder or are you taking a blood thinner?		□ Yes	□ No	□ Unknown	
11.	Are you pregnant or	breastfeeding?		□ Yes	□ No	□ Unknown	

## **Emergency Use Authorization**

The FDA has made the COVID-19 vaccine available under an emergency use authorization (EUA). The EUA is used when circumstances exist to justify the emergency use of drugs and biological products during an emergency, such as the COVID-19 pandemic. This vaccine has not completed the same type of review as an FDA-approved or cleared product. However, the FDA's decision to make the vaccine available under an EUA is based on the existence of a public health emergency and the totality of scientific evidence available, showing that known and potential benefits of the vaccine outweigh the known and potential risks.

## Consent

I have been provided with the Vaccine Information sheet or patient fact sheet corresponding to the COVID-19 vaccination that I am receiving. I have read the information provided about the vaccine that I am about to receive. I have been given an opportunity to ask questions which were answered to my satisfaction (and ensured the person named above for whom I am authorized to provide surrogate consent was also given a chance to ask questions. I understand the benefits and risks of vaccination and I voluntarily assume full responsibility for any reactions that may result. I attest that I am eligible to receive the COVID-19 vaccination at this time based on any jurisdiction-based requirements that apply to me. I understand that I should remain in this area for 15 minutes after the vaccination to be monitored for any potential adverse reactions. I

1

07/07/2022

understand that two doses of this vaccine will need to be administered (given) for it to be effective. I request that the COVID-19 vaccination be given to me (or the person named above for whom I am authorized to make this request and provide surrogate consent). Lunderstand there will be no cost to me for this vaccine. Lunderstand that any monies or benefits for administering the vaccine will be assigned and transferred to the vaccinating provider, including benefits/monies from my health insurance plan, or other third parties who are financially responsible for my medical care. I authorize release of all information needed (including but not limited to medical records, copies of claims and itemized bills) to verify payment and as needed for other public health purposes, including reporting to applicable vaccine registries and employers. Recipient/Surrogate/Guardian(Signature) Date / Time Print Name Relationship to patient, if other than recipient To be completed by immunizer Vaccine Name Administration Administration Site -IM ☐ First Dose ☐ Second / Later Dose □ Left Deltoid ☐ Right Deltoid Moderna/ Pfizer/ Janssen (Johnson & Johnson) **EUA Date** □ 0.5 ml Dosage Vaccine Expiration Date **Lot Number** □ I have reviewed side effects with patient (and parent, guardian or surrogate, as applicable) □ Iconfirm that the patient (and their surrogate, if applicable) was given an opportunity to ask questions about the vaccination, and all the questions asked by them (and/ortheir surrogate) have been answered correctly and to the best of my ability. □ I have confirmed the patient eligibility based on evidence that the patient has provided Printed Name: Vaccinator Signature: \_\_ Date: \_\_\_\_\_ Time Injected: Time Left: or 🗆 Unknown Time patient waited ☐ Refused to wait in office after injection □ 15 min  $\square$  20 min □ 30 min **Adverse Reactions - Systemic Adverse Reactions - Systemic Adverse Reactions - Injection Site** ☐ No swelling, redness or pain noted at the injection site ☐ No reactions noted during visit □ Nausea/Vomiting □ Pain at injection site □ Fatigue □ Axillary Swelling/Tenderness □ Headache ☐ Swelling at injection site □ Fever ☐ Erythema at the injection site □ Myalgia □ Anaphylaxis □ Other: □ Arthralgia □ Rash □ Chills □ Other Provider notes (if applicable): To be completed by immunizer, as required by state immunization registry reporting, only for states listed State Instructions Check all fields for patients 18 years and younger MS Check Race and Ethnicity for all patients. Select Next of Kin for patients 18 years of age and younger. ОК Next of Kin- 18 Years or Younger Name: Phone: Relationship: Address: City/State: Zip: State of NJ Only Prescriber Address: **Prescriber Name:** For CA, MA, MT, NJ, NM, NY, TX (For CA, this indicator means the registry will not share with Universities, Schools, or other agencies.

□ Yes

**Registry Sharing Indicator**