

# ST. JESUS PHARMACY 4180 BROADWAY NEW YORK, NY 10033

# **COVID-19 Vaccine Screening and Consent Form for Children and Adults**

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N	lame (please print)							
A	ddress	City		State	Zip			
Parent/Guardian/Surrogate (if applicable, please print)  Phone								
DC	DB Email Ad	ldress				·		
	•	tate Status Below: W -	Single D – Divorce - Widowed V – Civ - Unknown SEPAR RTNER – Life Partner	il Union RATED –		ed y Separated		
Prii	mary Care Physician Address/Phone Number		wine on -o	aici oi i	viaitii a	Ciui		
Sci	reening Questionnaire: The following questions will help us determine if th question, it does not necessarily mean the vaccine cannot be given. It jus healthcare		_	-				
1. 2.	Are you feeling sick today?  In the last 10 days, have you had a COVID-19 test because you had results or been told by a health care provider or health department COVID-19 infection or exposure?	□ Yes	□ No	□ Unknown □ Unknown				
3.	Has the person to be vaccinated ever received a dose of COVID-19 vaccine?  If yes, which product was administered?  Pfizer-BioTech   Janssen (Johnson & Johnson)   Another product  Moderna   Novavax  How many doses of COVID-19 vaccine were previously administered?  Did you bring the vaccination record card or other documentation?							
4.	Have you ever had an immediate allergic reaction (e.g., hives, facial swelling, difficulty breathing, anaphylaxis) to any vaccine, injection, or shot or to any component of the COVID-19 vaccine, or a severe allergic reaction (anaphylaxis) to anything?							
5.								
6.	Are you pregnant or considering becoming pregnant?			□ Yes	□ No	□ Unknown		
7.	Do you have a bleeding disorder, a history of blood clots or are you			□ Yes	□ No	□ Unknown		
8.	around the heart)?							
9.	Do you have a history of MIS-C or MIS-A (multisystem inflammator inflammatory syndrome in adults)?	y synarome in chilaren or	muitisystem	□ Yes	□ No	□ Unknown		
10	Have you recently received an orthopoxvirus vaccine within the las	st 4 weeks (e.g., JYNNEOS	or ACAM2000)?	□ 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 undergone the same type of review as an FDA-approved or cleared product. However, the FDA's decision to make the vaccine available is based on the totality of scientific evidence available, showing that known and potential benefits of the vaccine outweigh the known and potential risks. The Novavax COVID-19 vaccine is EUA authorized for those individuals 12 years and older. Please note: FDA approved the Pfizer-BioNTech COVID-19 vaccine in individuals 12 years of age and older; and approved the Moderna COVID-19 vaccine in individuals 18 years of age and older. These vaccines continue to be available under an EUA for certain populations, including Pfizer-BioNTech COVID-19 vaccine for those individuals 6 months through 11 years old, and Moderna COVID-19 vaccine for individuals 6 months through 17 years old and for the administration of an additional dose in the populations set forth in the consent section below.

### **Emergency Use Instruction**

Recipient/Surrogate/Guardian Signature

Emergency Use Instructions (EUIs) are issued by the CDC to provide information about emergency use of FDA-approved medical products that may not be included in or differ in some way from the information provided in the FDA-approved labeling (package insert). The COVID-19 vaccine by Pfizer-BioNTech is an FDA-approved COVID-19 vaccine (brand name Comirnaty, mRNA) to prevent COVID-19 in persons 12 years of age and older. CDC is issuing EUI to provide information about use of this vaccine as an additional primary dose in certain immunocompromised persons (12 years of age and older) and a booster dose in certain adults (18 years of age and older) who received certain non-FDA authorized or approved COVID-19 vaccine (e.g., certain vaccines available outside of the United States or from clinical trial participation).

#### Consent

I have read, or had explained to me, the information sheet about the COVID-19 vaccination. I understand that if my vaccine requires two doses, I will need to be administered (given) two doses to be considered fully vaccinated. Further, I understand that a booster dose of COVID-19 vaccine is recommended at least 2 months following the completion of a COVID-19 vaccine primary series or a monovalent booster dose to increase my protection.

I have had a chance 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 the vaccination as described.

Date / Time

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). I understand there will be no cost to me for this vaccine. I understand that any monies or benefits for administering the vaccine will be assigned and transferred to the vaccinating provider, including benefits/monies from my health plan, Medicare 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.

**Print Name** 

Relationship to Patient (if other than recipient)

Vaccine Name	Administration			Manufacturer EUA Fact S	
	1st Dose 2nd Dose		Booster	& Lot #	Date
Pfizer/BioNTech					
Moderna					
Novavax	□ First Dose	□ Second Dose			
Accounting for an	y previous vaccine doses	administered, which number	dose is this?	<del>,, '</del>	
Injection Site	□ Left Deltoid	□ Right Deltoid	□ Left Thigh	□ Right Thigh	
Dosage	□ 0.2 ml	□ 0.25 ml	□ 0.3 ml	□ 0.5 ml	
☐ I have provion vaccination was		r parent, guardian, or surro	ogate, as applicable) with i	information about the vac	cine and consent t
Vaccinator Signa	ature:				