



COVID-19 Vaccine Screening and Consent Form: *Ages 12 Years and Older

Recipient Name (please print)		Preferred Name			
Address		City	State	Zip	Email Address
Parent/Guardian/ Surrogate (if applicable, please print)		Phone		Preferred Language	
DOB	Current Gender ID Key: W – Woman/Girl TW – Transgender Woman/Girl M – Man/Boy Indicate ID Below: TM – Transgender Man/Boy NB – Non-Binary Person GNC – Gender Non-Conforming <input type="text"/> Q – Not Sure/Questioning NR – Chose not to Respond GNL - Gender not Listed (write-in) * Gender Pronouns: write-in by client’s name				
Sex Assigned at Birth Key: M – Male F – Female Indicate Sex Below: <input type="text"/> I – Intersex NR – Chose not to Respond		Marital Status Key: S – Single D – Divorced M – Married Indicate Status Below: <input type="text"/> W – Widowed V – Civil Union U – Unknown SEPARATED – Legally Separated PARTNER – Life Partner			
Ethnicity Key: DECL – Declined Indicate Ethnicity Below: <input type="text"/> HIS – Hispanic Origin NHL – Non-Hispanic Origin UNK – Unknown		Race Key: AIA – Native American or Alaskan ASN – Asian Indicate Race Below: <input type="text"/> BAA – African American or Black DECL – Declined NHP – Native Hawaiian or Pacific Islander WHT – White OTH – Other or Multiracial			
Primary Insurance Name		Primary Insurance ID#		Subscriber Name/DOB	Subscriber Relation to Patient
Primary Insurance Address		Primary Insurance Group #		Primary Insurance Phone #	
Secondary Insurance Name		Secondary Insurance ID#		Subscriber Name/DOB	Subscriber Relation to Patient
Secondary Insurance Address		Secondary Insurance Group #		Secondary Insurance Phone #	
Clinic/Office Site Where Vaccine is Administered		Primary Care Physician Address/Phone Number			

Screening Questionnaire

1.	Are you feeling sick today?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
2.	In the last 10 days, have you had a COVID-19 test because you had symptoms and are still awaiting your test results or been told by a health care provider or health department to isolate or quarantine at home due to COVID-19 infection or exposure?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
3.	Have you been treated with antibody therapy or convalescent plasma for COVID-19 in the past 90 days (3 months)? <i>If yes, when did you receive the last dose? Date:</i>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
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?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
5.	Are you pregnant or considering becoming pregnant?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
6.	Do you have cancer, leukemia, HIV/AIDS or any other condition that weakens the immune system?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
7.	Do you take any medications that affect your immune system, such as cortisone, prednisone or other steroids, anticancer drugs, or have you had any radiation treatments?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
8.	Do you have a bleeding disorder, a history of blood clots or are you taking a blood thinner?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown

9.	Do you have a history of myocarditis (inflammation of the heart muscle) or pericarditis (inflammation of the lining around the heart)?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
10.	Have you had Guillain-Barre Syndrome after receipt of the Janssen vaccine?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
11.	Do you have a history of MIS-C or MIS-A (multisystem inflammatory syndrome in children or multisystem inflammatory syndrome in adults)?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
12.*	Are you 12 years of age or older and have you received a complete COVID-19 vaccine primary series (e.g., 2 doses of Moderna, Pfizer, or Novavax vaccine, or 1 dose of Janssen vaccine) or any booster dose at least 2 months ago?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
				Date of last dose: (if applicable)
13.**	If you had a previous dose of Janssen (Johnson & Johnson), did you develop thrombosis with thrombocytopenia syndrome (TTS)?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
14. ¹	Have you received a previous dose of a non-FDA authorized or approved COVID-19 vaccine authorized by the WHO ¹ but not by the FDA (AstraZeneca – VAXZEVRIA, Sinovac – CORONAVAC, Serum Institute of India – COVISHIELD, Sinopharm/BIBP, COVAXIN, Nuvaxovid, COVOVAX, or CanSino Biologics – Convidecia)?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown

*Question #12 pertain to monovalent/bivalent booster dose eligibility for Pfizer, Moderna, Novavax or Janssen.

**Question #13 pertains to booster dose eligibility for Janssen.

¹ As set forth in the [CDC's Emergency Use Instructions \(EUI\)](#), a non-FDA authorized or approved COVID-19 vaccine such as those vaccines "listed for emergency use by the World Health Organization, or is included in CDC's Technical Instructions for Implementing Presidential Proclamation Advancing Safe Resumption of Global Travel During the COVID-19 Pandemic and CDC's Order, or that is a non-placebo part of a clinical trial within or outside the United States that is a WHO-EUL COVID-19 vaccine or a vaccine that is not listed for emergency use by WHO but for which a U.S. data and safety monitoring board or equivalent has independently confirmed efficacy in the United States (hereinafter "non-FDA authorized or approved COVID-19 vaccines").

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 Janssen (Johnson & Johnson) COVID-19 vaccine is EUA authorized for those individuals 18 years old and older. 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 as a two-dose series in individuals 12 years of age and older; and approved the Moderna COVID-19 vaccine as a two-dose series 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 a third dose in the populations set forth in the consent section below.

Emergency Use Instruction

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.

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.

Recipient/Surrogate/Guardian Signature Date / Time Print Name Relationship to Patient (if other than recipient)

Telephonic Interpreter's ID # Date / Time

OR

Signature: Interpreter Date/ Time Print Interpreter's Name Relationship to Patient

Area Below to be Completed by Vaccinator

Which vaccine is the patient receiving today?

Vaccine Name	Administration				Manufacturer & Lot #	EUA Fact Sheet Date
Pfizer/BioNTech	<input type="checkbox"/> First Dose	<input type="checkbox"/> Second Dose	<input type="checkbox"/> Bivalent mRNA Booster (≥ 5 years old)			
Moderna	<input type="checkbox"/> First Dose	<input type="checkbox"/> Second Dose	<input type="checkbox"/> Bivalent mRNA Booster (≥ 6 years old)			
Novavax	<input type="checkbox"/> First Dose	<input type="checkbox"/> Second Dose	<input type="checkbox"/> Bivalent mRNA Booster (≥ 18 years old)			
Janssen	<input type="checkbox"/> Single Dose	<input type="checkbox"/> Additional Dose	<input type="checkbox"/> Bivalent mRNA Booster (≥ 18 years old)			
Administration Site	<input type="checkbox"/> Left Deltoid	<input type="checkbox"/> Right Deltoid	<input type="checkbox"/> Left Thigh	<input type="checkbox"/> Right Thigh		
Dosage	<input type="checkbox"/> 0.2 ml	<input type="checkbox"/> 0.25 ml	<input type="checkbox"/> 0.3 ml	<input type="checkbox"/> 0.5 ml		

I have provided the patient (and/or parent, guardian, or surrogate, as applicable) with information about the vaccine and consent to vaccination was obtained.

Vaccinator Signature: _____

*** Use of this form is optional.**

Updated October 17, 2022