COVID-19 Vaccine Consent Form for Child Under 18 or Adult Conservatee

Please print information about the patient to receive vaccine

PATIENT'S NAME (Last)	(First)	(M.I.)	**SUFFIX** (eg. Jr, III)

DATE OF BIRTH (MM/DD/YYYY)	**AGE†**	**PHONE** ( )	☐ Cell ☐ Home

ADDRESS	**CITY**	**STATE**	**ZIP**

SEX AT BIRTH ☐ Female ☐ Male	**GENDER IDENTITY** (*optional*) ☐ Female ☐ Male ☐ Non-binary ☐ Unknown ☐ Prefer not to say	**Ethnicity:** ☐ Hispanic/Latino ☐ Unknown ☐ Not Hispanic/Latino

RACE ☐ American Indian/Alaska Native ☐ Asian ☐ Black/African American ☐ Native Hawaiian/Other Pacific Islander ☐ White ☐ Other

Guardian relationship to client: ☐ Father ☐ Mother ☐ Legal Guardian ☐ Other

I understand that the COVID-19 vaccine is a voluntary vaccine currently being given under the Emergency Use Authorization status and only a parent or legal guardian has the authority to consent to a minor or adult conservatee receiving this vaccine. By signing this form, I certify that I have the legal authority to do so on behalf of the patient identified above and will indemnify Oklahoma City-County Health Department against challenges to this consent or my status as legally able to provide consent for this vaccine.

Guardian's State or Federally issued ID # _____ (incl. State License, Passport, Consulate Card, etc.)

Screening for Vaccine Eligibility	**YES**	**NO**
Has the patient ever received a dose of the COVID-19 Pfizer-BioNTech vaccine? If yes, which one? ☐ Pfizer-BioNTech ☐ Moderna ☐ Janssen (Johnson & Johnson)		
Has the patient ever had an allergic reaction to: ☐ a component of a COVID-19 vaccine, including either of the following: -polyethylene glycol (PEG), which is found in some medications, such as laxatives and preps for colonoscopy procedures -polysorbate, which is found in some vaccines, film coated tablets, and intravenous steroids ☐ a previous dose of COVID-19 vaccine ☐ a vaccine or injectable therapy that contains multiple components, one of which is a COVID-19 component, but it is not known which component elicited the immediate reaction ☐ another vaccine (other than COVID-19 vaccine) or an injectable medication?		
Has the patient ever had a severe allergic reaction (e.g., anaphylaxis) to something other than a vaccine or injectable medication? This would include food, pet, venom, environmental, or oral medication allergies.		
Does the patient have a history of myocarditis or pericarditis?		
Has the patient ever had COVID-19 and been treated with monoclonal antibodies or convalescent plasma?		
Has the patient ever been diagnosed with Multisystem Inflammatory Syndrome (MIS-C or MIS-A) after a COVID-19 infection?		
Does the patient have a bleeding disorder or take a blood thinner?		
Does the patient have a history of heparin-induced thrombocytopenia (HIT)?		
Does the patient have a history of Guillian-Barré Syndrome (GBS)?		
If receiving a third dose of a COVID-19 vaccine, I attest that the patient is eligible for vaccination for this event because they: ☐ have undergone solid organ transplantation. ☐ are diagnosed with conditions that are considered to have an equivalent level of immunocompromise to solid organ transplantation.		

I understand that should I have any questions about the COVID-19 vaccine, need assistance filling out this form, or need any other information regarding COVID-19 I can contact the Oklahoma City County Health Department at (405) 425-4489 prior to signing this form or at the vaccine distribution location.

CONSENT FOR DEPENDENT'S VACCINATION AND RELEASE OF VACCINATION INFORMATION:

I have read or had explained to me the information contained in the *Emergency Use Authorization Fact Sheet for Recipients and Caregivers* for the COVID-19 vaccine and understand the risks and benefits of the vaccine. I have had a chance to ask questions which have been answered to my satisfaction I understand the benefits and risks of the vaccine. I understand that if my dependent exhibits disruptive behavior while staff is trying to administer the vaccine, they will not receive the vaccine at this clinic and will have to be taken to the health department or to their provider for this vaccine.

I authorize disclosure of this vaccination information to public health officials and other health care professionals. I understand that this vaccination will be recorded in the Oklahoma State Immunization Information System (OSIIS) for the purposes of sharing vaccination information with other health care providers and tracking vaccine inventory only.

I acknowledge that I can access a copy of Oklahoma City County Health Department's HIPPA Privacy Notice as required by the Health Information Portability and Accountability Act (HIPPA) at https://www.occhd.org/about/contact-us/hippa. I acknowledge a copy of the manufacturer's COVID-19 Fact Sheet for Recipients and Caregivers is provided prior to receiving the vaccine. This information can also be accessed at https://www.vaxokc.com/eua. Vaccine information statements for Pfizer can be at www.cvdvaccine.com.

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For health and safety reasons masks must be worn at all times during a vaccination event. If my child or adult conservatee does not have a mask one will be provided to him or her to wear during the vaccination event. By signing this form, I hereby give my consent to have my child or adult conservatee wear a mask during the vaccination process with OCCHD.

"In the event of an emergency situation, emergency medication (Epinephrine/Benadryl) and/or oxygen may be administered to my child or adult conservatee. In the event of an emergency situation where I am not present, I authorize Oklahoma City County Health Department staff or designee to obtain any necessary medical care they deem necessary including, but not limited to, obtaining paramedic assistance and transport to a local hospital for additional treatment or observation."

Signature of Parent/Guardian_____Date:_____

Please print Parent/Guardian name_____

****FOR OFFICIAL USE ONLY****

Client Name (Last, First, MI) _____ Client DOB (MM/DD/YYYY)_____

OFFICE USE ONLY – DO NOT WRITE BELOW		
Ask before administration: Is the client suffering from a moderate or severe acute illness with or without fever? ☐ Y ☐ N Is the client pregnant or breastfeeding? ☐ Y ☐ N Client completed the manufacturer's screening questions: ☐ Y ☐ N		
Vaccine Manufacturer: Lot #: Exp. Date:	Site: ☐ LT DELTOID IM ☐ RT DELTOID IM ☐ LT VAST LAT IM ☐ RT VAST LAT IM	Dose Number: ☐ 1st ☐ 2nd EUA*/VIS given? ☐ Y ☐ N Reaction? ☐ Y ☐ N
Vaccination Complete? ☐Complete ☐Refused ☐Not administered ☐Partially administered ☐No recorded completion status		
Provider Signature:		

FACT SHEET FOR RECIPIENTS AND CAREGIVERS

EMERGENCY USE AUTHORIZATION (EUA) OF THE PFIZER-BIONTECH COVID-19 VACCINE TO PREVENT CORONAVIRUS DISEASE 2019 (COVID-19) IN INDIVIDUALS 12 YEARS OF AGE AND OLDER

You are being offered the Pfizer-BioNTech COVID-19 Vaccine to prevent Coronavirus Disease 2019 (COVID-19) caused by SARS-CoV-2. This Fact Sheet contains information to help you understand the risks and benefits of the Pfizer-BioNTech COVID-19 Vaccine, which you may receive because there is currently a pandemic of COVID-19.

The Pfizer-BioNTech COVID-19 Vaccine is a vaccine and may prevent you from getting COVID-19. There is no U.S. Food and Drug Administration (FDA) approved vaccine to prevent COVID-19.

Read this Fact Sheet for information about the Pfizer-BioNTech COVID-19 Vaccine. Talk to the vaccination provider if you have questions. It is your choice to receive the Pfizer-BioNTech COVID-19 Vaccine.

The Pfizer-BioNTech COVID-19 Vaccine is administered as a 2-dose series, 3 weeks apart, into the muscle.

The Pfizer-BioNTech COVID-19 Vaccine may not protect everyone.

This Fact Sheet may have been updated. For the most recent Fact Sheet, please see www.cvdvaccine.com.

WHAT YOU NEED TO KNOW BEFORE YOU GET THIS VACCINE

WHAT IS COVID-19?

COVID-19 disease is caused by a coronavirus called SARS-CoV-2. This type of coronavirus has not been seen before. You can get COVID-19 through contact with another person who has the virus. It is predominantly a respiratory illness that can affect other organs. People with COVID-19 have had a wide range of symptoms reported, ranging from mild symptoms to severe illness. Symptoms may appear 2 to 14 days after exposure to the virus. Symptoms may include: fever or chills; cough; shortness of breath; fatigue; muscle or body aches; headache; new loss of taste or smell; sore throat; congestion or runny nose; nausea or vomiting; diarrhea.

WHAT IS THE PFIZER-BIONTECH COVID-19 VACCINE?

The Pfizer-BioNTech COVID-19 Vaccine is an unapproved vaccine that may prevent COVID-19. There is no FDA-approved vaccine to prevent COVID-19.

The FDA has authorized the emergency use of the Pfizer-BioNTech COVID-19 Vaccine to prevent COVID-19 in individuals 12 years of age and older under an Emergency Use Authorization (EUA).

For more information on EUA, see the "**What is an Emergency Use Authorization (EUA)?**" section at the end of this Fact Sheet.

WHAT SHOULD YOU MENTION TO YOUR VACCINATION PROVIDER BEFORE YOU GET THE PFIZER-BIONTECH COVID-19 VACCINE?
Tell the vaccination provider about all of your medical conditions, including if you:

- have any allergies
- have had myocarditis (inflammation of the heart muscle) or pericarditis (inflammation of the lining outside the heart)
- have a fever
- have a bleeding disorder or are on a blood thinner
- are immunocompromised or are on a medicine that affects your immune system
- are pregnant or plan to become pregnant
- are breastfeeding
- have received another COVID-19 vaccine
- have ever fainted in association with an injection

WHO SHOULD GET THE PFIZER-BIONTECH COVID-19 VACCINE?
FDA has authorized the emergency use of the Pfizer-BioNTech COVID-19 Vaccine in individuals 12 years of age and older.

WHO SHOULD <u>NOT</u> GET THE PFIZER-BIONTECH COVID-19 VACCINE?
You should not get the Pfizer-BioNTech COVID-19 Vaccine if you:

- had a severe allergic reaction after a previous dose of this vaccine
- had a severe allergic reaction to any ingredient of this vaccine.

WHAT ARE THE INGREDIENTS IN THE PFIZER-BIONTECH COVID-19 VACCINE?
The Pfizer-BioNTech COVID-19 Vaccine includes the following ingredients: mRNA, lipids ((4-hydroxybutyl)azanediyl)bis(hexane-6,1-diyl)bis(2-hexyldecanoate), 2 [(polyethylene glycol)-2000]-N,N-ditetradecylacetamide, 1,2-Distearoyl-sn-glycero-3-phosphocholine, and cholesterol), potassium chloride, monobasic potassium phosphate, sodium chloride, dibasic sodium phosphate dihydrate, and sucrose.

HOW IS THE PFIZER-BIONTECH COVID-19 VACCINE GIVEN?
The Pfizer-BioNTech COVID-19 Vaccine will be given to you as an injection into the muscle.

The Pfizer-BioNTech COVID-19 Vaccine vaccination series is 2 doses given 3 weeks apart.

If you receive one dose of the Pfizer-BioNTech COVID-19 Vaccine, you should receive a second dose of this same vaccine 3 weeks later to complete the vaccination series.

If you are immunocompromised, you may receive a third dose of the Pfizer-BioNTech COVID-19 Vaccine at least 1 month after the second dose.

HAS THE PFIZER-BIONTECH COVID-19 VACCINE BEEN USED BEFORE?
The Pfizer-BioNTech COVID-19 Vaccine is an unapproved vaccine. In clinical trials, approximately 23,000 individuals 12 years of age and older have received at least 1 dose of the Pfizer-BioNTech COVID-19 Vaccine.

WHAT ARE THE BENEFITS OF THE PFIZER-BIONTECH COVID-19 VACCINE?
In an ongoing clinical trial, the Pfizer-BioNTech COVID-19 Vaccine has been shown to prevent COVID-19 following 2 doses given 3 weeks apart. The duration of protection against COVID-19 is currently unknown.

WHAT ARE THE RISKS OF THE PFIZER-BIONTECH COVID-19 VACCINE?
There is a remote chance that the Pfizer-BioNTech COVID-19 Vaccine could cause a severe allergic reaction. A severe allergic reaction would usually occur within a few minutes to one hour after getting a dose of the Pfizer-BioNTech COVID-19 Vaccine. For this reason, your vaccination provider may ask you to stay at the place where you received your vaccine for monitoring after vaccination. Signs of a severe allergic reaction can include:

- Difficulty breathing
- Swelling of your face and throat
- A fast heartbeat
- A bad rash all over your body
- Dizziness and weakness

Myocarditis (inflammation of the heart muscle) and pericarditis (inflammation of the lining outside the heart) have occurred in some people who have received the Pfizer-BioNTech COVID-19 Vaccine. In most of these people, symptoms began within a few days following receipt of the second dose of the Pfizer-BioNTech COVID-19 Vaccine. The chance of having this occur is very low. You should seek medical attention right away if you have any of the following symptoms after receiving the Pfizer-BioNTech COVID-19 Vaccine:

- Chest pain
- Shortness of breath
- Feelings of having a fast-beating, fluttering, or pounding heart

Side effects that have been reported with the Pfizer-BioNTech COVID-19 Vaccine include:

- severe allergic reactions
- non-severe allergic reactions such as rash, itching, hives, or swelling of the face
- myocarditis (inflammation of the heart muscle)
- pericarditis (inflammation of the lining outside the heart)
- injection site pain
- tiredness
- headache

- muscle pain
- chills
- joint pain
- fever
- injection site swelling
- injection site redness
- nausea
- feeling unwell
- swollen lymph nodes (lymphadenopathy)
- diarrhea
- vomiting
- arm pain

These may not be all the possible side effects of the Pfizer-BioNTech COVID-19 Vaccine. Serious and unexpected side effects may occur. Pfizer-BioNTech COVID-19 Vaccine is still being studied in clinical trials.

WHAT SHOULD I DO ABOUT SIDE EFFECTS?
If you experience a severe allergic reaction, call 9-1-1, or go to the nearest hospital.

Call the vaccination provider or your healthcare provider if you have any side effects that bother you or do not go away.

Report vaccine side effects to FDA/CDC Vaccine Adverse Event Reporting System (VAERS). The VAERS toll-free number is 1-800-822-7967 or report online to https://vaers.hhs.gov/reportevent.html. Please include “Pfizer-BioNTech COVID-19 Vaccine EUA” in the first line of box #18 of the report form.

In addition, you can report side effects to Pfizer Inc. at the contact information provided below.

Website	Fax number	Telephone number
www.pfizersafetyreporting.com	1-866-635-8337	1-800-438-1985

You may also be given an option to enroll in v-safe. V-safe is a new voluntary smartphone-based tool that uses text messaging and web surveys to check in with people who have been vaccinated to identify potential side effects after COVID-19 vaccination. V-safe asks questions that help CDC monitor the safety of COVID-19 vaccines. V-safe also provides second-dose reminders if needed and live telephone follow-up by CDC if participants report a significant health impact following COVID-19 vaccination. For more information on how to sign up, visit: www.cdc.gov/vsafe.

WHAT IF I DECIDE NOT TO GET THE PFIZER-BIONTECH COVID-19 VACCINE?
It is your choice to receive or not receive the Pfizer-BioNTech COVID-19 Vaccine. Should you decide not to receive it, it will not change your standard medical care.

ARE OTHER CHOICES AVAILABLE FOR PREVENTING COVID-19 BESIDES PFIZER-BIONTECH COVID-19 VACCINE?
Currently, there is no approved alternative vaccine available for prevention of COVID-19. Other vaccines to prevent COVID-19 may be available under Emergency Use Authorization.

CAN I RECEIVE THE PFIZER-BIONTECH COVID-19 VACCINE WITH OTHER VACCINES?
There is no information on the use of the Pfizer-BioNTech COVID-19 Vaccine with other vaccines.

WHAT IF I AM IMMUNOCOMPROMISED?
If you are immunocompromised, you may receive a third dose of the Pfizer-BioNTech COVID-19 Vaccine. The third dose may still not provide full immunity to COVID-19 in people who are immunocompromised, and you should continue to maintain physical precautions to help prevent COVID-19. In addition, your close contacts should be vaccinated as appropriate.

WHAT IF I AM PREGNANT OR BREASTFEEDING?
If you are pregnant or breastfeeding, discuss your options with your healthcare provider.

WILL THE PFIZER-BIONTECH COVID-19 VACCINE GIVE ME COVID-19?
No. The Pfizer-BioNTech COVID-19 Vaccine does not contain SARS-CoV-2 and cannot give you COVID-19.

KEEP YOUR VACCINATION CARD
When you get your first dose, you will get a vaccination card to show you when to return for your second dose of Pfizer-BioNTech COVID-19 Vaccine. Remember to bring your card when you return.

ADDITIONAL INFORMATION

If you have questions, visit the website or call the telephone number provided below.

To access the most recent Fact Sheets, please scan the QR code provided below.

Global website	Telephone number
www.cvdvaccine.com	1-877-829-2619 (1-877-VAX-CO19)

HOW CAN I LEARN MORE?

- Ask the vaccination provider.
- Visit CDC at https://www.cdc.gov/coronavirus/2019-ncov/index.html.
- Visit FDA at https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization.
- Contact your local or state public health department.

WHERE WILL MY VACCINATION INFORMATION BE RECORDED?

The vaccination provider may include your vaccination information in your state/local jurisdiction's Immunization Information System (IIS) or other designated system. This will ensure that you receive the same vaccine when you return for the second dose. For more information about IISs visit: https://www.cdc.gov/vaccines/programs/iis/about.html.

CAN I BE CHARGED AN ADMINISTRATION FEE FOR RECEIPT OF THE COVID-19 VACCINE?

No. At this time, the provider cannot charge you for a vaccine dose and you cannot be charged an out-of-pocket vaccine administration fee or any other fee if only receiving a COVID-19 vaccination. However, vaccination providers may seek appropriate reimbursement from a program or plan that covers COVID-19 vaccine administration fees for the vaccine recipient (private insurance, Medicare, Medicaid, Health Resources & Services Administration [HRSA] COVID-19 Uninsured Program for non-insured recipients).

WHERE CAN I REPORT CASES OF SUSPECTED FRAUD?

Individuals becoming aware of any potential violations of the CDC COVID-19 Vaccination Program requirements are encouraged to report them to the Office of the Inspector General, U.S. Department of Health and Human Services, at 1-800-HHS-TIPS or https://TIPS.HHS.GOV.

WHAT IS THE COUNTERMEASURES INJURY COMPENSATION PROGRAM?

The Countermeasures Injury Compensation Program (CICP) is a federal program that may help pay for costs of medical care and other specific expenses of certain people who have been seriously injured by certain medicines or vaccines, including this vaccine. Generally, a claim must be submitted to the CICP within one (1) year from the

date of receiving the vaccine. To learn more about this program, visit www.hrsa.gov/cicp/ or call 1-855-266-2427.

WHAT IS AN EMERGENCY USE AUTHORIZATION (EUA)?

The United States FDA has made the Pfizer-BioNTech COVID-19 Vaccine available under an emergency access mechanism called an EUA. The EUA is supported by a Secretary of Health and Human Services (HHS) declaration that circumstances exist to justify the emergency use of drugs and biological products during the COVID-19 pandemic.

The Pfizer-BioNTech COVID-19 Vaccine has not undergone the same type of review as an FDA-approved or cleared product. FDA may issue an EUA when certain criteria are met, which includes that there are no adequate, approved, available alternatives. In addition, the FDA decision is based on the totality of scientific evidence available showing that the product may be effective to prevent COVID-19 during the COVID-19 pandemic and that the known and potential benefits of the product outweigh the known and potential risks of the product. All of these criteria must be met to allow for the product to be used in the treatment of patients during the COVID-19 pandemic.

The EUA for the Pfizer-BioNTech COVID-19 Vaccine is in effect for the duration of the COVID-19 EUA declaration justifying emergency use of these products, unless terminated or revoked (after which the products may no longer be used).

Manufactured by
Pfizer Inc., New York, NY 10017

Manufactured for
BioNTech Manufacturing GmbH
An der Goldgrube 12
55131 Mainz, Germany

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Scan to capture that this Fact Sheet was provided to vaccine recipient for the electronic medical records/immunization information systems.

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