

Registration Form for COVID Vaccines

Please fill out this form in its entirety.

Patient Name (Last, First)		Date of Birth (mm/dd/yyyy)	
Address:	City:	State:	Zip Code:
Phone Number:		Emergency Contact:	
Email:		Name:	
Marital Status:		Relation:	Phone:
Race: <input type="checkbox"/> Asian <input type="checkbox"/> Black or African American <input type="checkbox"/> American Indian or Alaska Native <input type="checkbox"/> Native Hawaiian <input type="checkbox"/> Other Pacific Islander <input type="checkbox"/> White/Latino <input type="checkbox"/> White/Non-Latino <input type="checkbox"/> Other Race		Gender Identity: <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Trans M/F <input type="checkbox"/> Non Binary <input type="checkbox"/> Other	
Ethnicity: <input type="checkbox"/> Latino <input type="checkbox"/> Non-Latino		Preferred Language:	
Latino Origin: <input type="checkbox"/> Cuban <input type="checkbox"/> General Latino <input type="checkbox"/> Mexican/Mexican-American/Chicano <input type="checkbox"/> Other Spanish/Latino <input type="checkbox"/> Puerto Rican <input type="checkbox"/> Unknown		Veteran <input type="checkbox"/> Yes <input type="checkbox"/> No	
		Homeless <input type="checkbox"/> Yes <input type="checkbox"/> No	
		Farmworker <input type="checkbox"/> Yes <input type="checkbox"/> No	

Income Information

Sea Mar requests this information from all patients for anonymous reporting purposes. Please circle the category that applies to you.

Family Size	Income Level					
1	0 - \$12,880	\$12,881-\$16,100	\$16,101-\$19,320	\$19,321-\$22,540	\$22,541-\$25,760	\$25,761+
2	0 - \$17,420	\$17,421-\$21,775	\$21,776-\$26,130	\$26,131-\$30,485	\$30,486-\$34,840	\$34,841+
3	0 - \$21,960	\$21,961-\$27,450	\$27,451-\$32,940	\$32,941-\$38,430	\$38,431-\$43,920	\$43,921+
4	0 - \$26,500	\$26,501-\$33,125	\$33,126-\$39,750	\$39,751-\$46,375	\$46,376-\$53,000	\$53,001+
5	0 - \$31,040	\$31,041-\$38,800	\$38,801-\$46,560	\$46,561-\$54,320	\$54,321-\$62,800	\$62,801+
6	0 - \$35,580	\$35,581-\$44,475	\$44,476-\$53,370	\$53,371-\$62,265	\$62,266-\$71,160	\$71,161+
7	0 - \$40,120	\$40,121-\$50,150	\$50,151-\$60,180	\$60,181-\$70,210	\$70,211-\$80,240	\$80,241+
8	0 - \$44,660	\$44,661-\$55,825	\$55,826-\$66,990	\$66,991-\$78,155	\$78,156-\$89,320	\$89,321+
Other (Provide Write-In Household Size and Income):						



Notice of Privacy Practices Acknowledgement

The Notice of Privacy Practices for Protected Health Information describes how medical information about you may be used and disclosed, how you can get access to this information and who to contact if you have questions, concerns or complaints.

Sea Mar has the responsibility to protect the privacy of your information, provide a Notice of Privacy Practices, and follow information practices that are described in this notice. If you have any questions, please contact Sea Mar's Vice President of Corporate and Legal Affairs at 206.763.5277.

By signing this form, you acknowledge receipt of Sea Mar Community Health Centers' Notice of Privacy Practices and Patient Rights and Responsibilities. Sea Mar encourages you to review these notices carefully.

I acknowledge receipt of Sea Mar Community Health Centers' Notice of Privacy Practices and Patient Rights and Responsibilities.

Patient or legally authorized individual signature

Date

Time

Printed name if signed on behalf of the patient

Relationship
(parent, legal guardian, personal representative)

Patient Name: <<PName>>

DOB: <<PDOB>>

Patient ID: <<PNumber>>

This form will be retained in your medical record.

Sliding Fee Scale Application

To comply with federal regulations and provide you a discount on Sea Mar services, it is necessary for you to fill out this form, answer some personal questions, and provide proof of income. Your answers will be kept on file and in strict confidence.

Patient Name:	DOB:	Patient ID:
Household Size:	Annual Income:	<input type="checkbox"/> I choose <u>NOT</u> to provide my income.

I choose **NOT** to apply for the sliding fee scale. Please sign and date below.

Signature

Date

I choose to apply for the sliding fee scale discount. The sliding fee scale is available for all patients, regardless of insurance status. If you have insurance, the sliding fee scale discount can be applied to charges not covered by insurance. Please complete the entire form to determine eligible discount.

Household Members	NAME	BIRTHDATE (MM/DD/YYYY)	HEALTH INSURANCE	RELATIONSHIP	SEA MAR PATIENT?	
	1					
	2					
	3					
	4					
	5					
	6					

SOURCE OF INCOME	ANNUAL INCOME	For You	For Spouse	For Children	For Others	Sub Total
	Gross Wages, Salaries, Tips					\$ 0.00
	Social Security & Pensions					\$ 0.00
	Annuity & Veteran Benefits					\$ 0.00
	Child Support & Alimony					\$ 0.00
	Self-Employment & Other					\$ 0.00
	For "Other," please explain:					TOTAL

By signing below, I agree to provide Sea Mar Community Health Centers with a proof of income for all persons listed above. Acceptable proof of income includes, but is not limited to, social security statements, paycheck stubs (two most recent), public assistance letter, tax return form, W-2 form, L&I check stub, unemployment check stub.

I understand that I will be asked to reapply for the sliding fee scale at least once a year so Sea Mar can maintain an updated application on file. I certify that the information provided is accurate and complete to the best of my knowledge. I understand that if I knowingly give false information that results in assistance for which I am not eligible, I will be subject to criminal prosecution. I give my consent to release any and all information from whatever source needed to verify the information I have given.

Signature

Date

OFFICE USE ONLY

Patient is eligible for Sliding Fee Scale: Yes No SFS Status (circle one): A B C D E F

POI Requested: _____ Initial: _____ POI Received: _____ Initial: _____

Authorized Adult Consent For COVID-19 Vaccinations

This form must be signed for patients ages 5 to 17-years-old receiving the COVID-19 vaccine. As of November 3, 2021, Pfizer is the only vaccine approved for ages 5 to 17.

Patient Name:	DOB:	Patient MRN:
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Acknowledgement

I have been provided an opportunity to review the COVID-19 Vaccine Fact Sheet for Recipients and Caregivers. I understand that I can review the Pfizer-BioNTech COVID-19 vaccine onsite or online at www.fda.gov/media/144414/download or by using the QR Code below.

Authorized Adult Consent

I am authorized to consent for the patient named above to receive this vaccine. I request that the vaccine be given to the patient named above. I understand that the patient should stay at the vaccine location for 15 to 30 minutes after receiving the vaccine to be monitored for potential immediate vaccine-related reactions and side effects and receive medical intervention if needed.

Printed Name of Authorized Adult

Relationship/Authority of Consenting Party

Signature of Authorized Adult

Date

Minor Consent

I am a legally emancipated minor, a minor married to an adult, or have been determined a mature minor. I request that I be given the vaccine. I understand that I should stay at the vaccine location for 15 to 30 minutes after receiving the vaccine to be monitored for potential immediate vaccine-related reactions and side effects and receive medical intervention if needed.

Signature of Emancipated/Married to An Adult Minor/Mature Minor

Date



QR CODE FOR VACCINE FACTSHEETS

VACCINATION CONSENT FORM

Pfizer-BioNTech COVID-19 Vaccine

The novel coronavirus SARS-CoV-2 (a/k/a COVID-19) is an infectious disease that appeared in late 2019.

I request that the Pfizer-BioNTech COVID-19 Vaccine be given to me or to the person named hereafter for whom I am authorized to make this request (select one): MYSELF PERSON NAMED BELOW

Recipient's Information:

_____ Last Name _____ First Name _____ Date of Birth _____ Gender _____

Address: _____

City: _____ State: _____ Zip: _____

Authorized Individual's Information (complete if different from vaccine recipient):

_____ Last Name _____ First Name _____ Date of Birth _____ Gender _____

Address: _____

City: _____ State: _____ Zip: _____

Relationship to recipient: _____

Vaccine is for (check one): Physician Contractor Employee Volunteer Other: _____

Company/Organization: _____

ACKNOWLEDGEMENTS (INITIAL EACH STATEMENT):

_____ Prior to vaccination, I was given a copy of the FDA's *Fact Sheet for Recipients and Caregivers* in connection with the administration of Pfizer-BioNTech COVID-19 Vaccine for ages 12 and above and Emergency Use Authorization (EUA) for the Pfizer-BioNTech COVID-19 Vaccine for ages 5-11, or was directed to the FDA's COVID-19 vaccination website at: [Pfizer-BioNTech COVID-19 Vaccine | cvdvaccine.com](https://www.fda.gov/covid19/vaccine/cvdvaccine.com).

_____ The recipient or their caregiver has the option to accept or refuse Pfizer-BioNTech COVID-19 Vaccine.

_____ The significant known and potential risks and benefits of Pfizer-BioNTech COVID-19 Vaccine, and the extent to which such risks and benefits are unknown, have been disclosed to me. Information about available alternative vaccines and the risks and benefits of those alternatives, to the extent reasonably known, have been disclosed to me.

_____ The Pfizer-BioNTech COVID-19 Vaccine is administered intramuscularly as a series of two doses (0.3 mL each) 3 weeks apart. Recipients must receive both doses of the Pfizer-BioNTech COVID-19 Vaccine to complete vaccination.

_____ Recipient is 5 years of age or older.

_____ Immunocompromised persons, including individuals receiving immunosuppressant therapy, may have a diminished immune response to the Pfizer-BioNTech COVID-19 Vaccine.

_____ Vaccine may not protect all vaccine recipients.

_____ 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.

_____ I have read or have had explained to me the information identified in the FDA's *Fact Sheet for Recipients and Caregivers* regarding the Pfizer-BioNTech COVID-19 Vaccine. I have had an opportunity to discuss the benefits and risks of this COVID-19 vaccine with a healthcare provider of my choice before vaccination. I have had a chance to ask questions which were answered to my satisfaction.

_____ I believe I understand the benefits and risks of this vaccine and ask that this vaccine be given to me or the person named for whom I am authorized to make this request.

MEDICAL SCREENING QUESTIONS: Check yes or no to each question below. Tell your vaccination provider about all your medical conditions, including if you answer “yes” to any question. Except for the last two (2) questions, a “yes” response to any other question means you may wish to consult with your individual healthcare provider before proceeding. Answering “yes” to either of the last two (2) questions means you should not be vaccinated today.

Question	Yes	No
Do you have any allergies?		
Do you have a fever?		
Do you have a bleeding disorder or are on a blood thinner?		
Are you immunocompromised or are you on a medicine that affects your immune system?		
Are you pregnant or plan to become pregnant?		
Are you breastfeeding?		
Have you received another COVID-19 vaccine?		
Have you had a severe allergic reaction after a previous dose of this vaccine?		
Have you had a severe allergic reaction to any ingredient of this vaccine?		
Have you ever fainted or felt lightheaded after receiving an injection or having blood drawn?		

Signature of Recipient OR Recipient's Authorized Individual

Date

DO NOT WRITE IN THIS SPACE—OFFICE USE ONLY VIS Edition Provided: _____

Vaccine: _____

Administration Date: _____

Manufacturer: _____

Lot #: _____

Exp. Date: _____

Route: _____

Site: _____

Volume (ml): _____

Nurse/ Provider's Signature

Date

Time

**VACCINE INFORMATION FACT SHEET FOR RECIPIENTS AND CAREGIVERS
ABOUT COMIRNATY (COVID-19 VACCINE, mRNA)
AND THE PFIZER-BIONTECH COVID-19 VACCINE TO PREVENT CORONAVIRUS
DISEASE 2019 (COVID-19) FOR USE IN INDIVIDUALS 12 YEARS OF AGE AND
OLDER**

FOR 12 YEARS OF AGE AND OLDER

You are being offered either COMIRNATY (COVID-19 Vaccine, mRNA) or the Pfizer-BioNTech COVID-19 Vaccine to prevent Coronavirus Disease 2019 (COVID-19) caused by SARS-CoV-2.

This Vaccine Information Fact Sheet for Recipients and Caregivers comprises the Fact Sheet for the authorized Pfizer-BioNTech COVID-19 Vaccine and also includes information about the FDA-licensed vaccine, COMIRNATY (COVID-19 Vaccine, mRNA) for use in individuals 12 years of age and older.

The FDA-approved COMIRNATY (COVID-19 Vaccine, mRNA) and the Pfizer-BioNTech COVID-19 Vaccine authorized for Emergency Use Authorization (EUA) for individuals 12 years of age and older, when prepared according to their respective instructions for use, can be used interchangeably.¹

COMIRNATY (COVID-19 Vaccine, mRNA) is an FDA-approved COVID-19 vaccine made by Pfizer for BioNTech. It is approved as a 2-dose series for prevention of COVID-19 in individuals 16 years of age and older. It is also authorized under EUA to provide:

- a 2-dose primary series to individuals 12 through 15 years of age;**
- a third primary series dose to individuals 12 years of age and older who have been determined to have certain kinds of immunocompromise;**
- a single booster dose to individuals 12 years of age and older who have completed a primary series with Pfizer-BioNTech COVID-19 Vaccine or COMIRNATY (COVID-19 Vaccine, mRNA); and**
- a single booster dose to individuals 18 years of age and older who have completed primary vaccination with a different authorized COVID-19 vaccine. The booster schedule is based on the labeling information of the vaccine used for the primary series.**

¹ When prepared according to their respective instructions for use, the FDA-approved COMIRNATY (COVID-19 Vaccine, mRNA) and the EUA-authorized Pfizer-BioNTech COVID-19 Vaccine for individuals 12 years of age and older can be used interchangeably without presenting any safety or effectiveness concerns.

The Pfizer-BioNTech COVID-19 Vaccine has received EUA from FDA to provide:

- **a 2-dose primary series to individuals 12 years of age and older;**
 - **a third primary series dose to individuals 12 years of age and older who have been determined to have certain kinds of immunocompromise;**
 - **a single booster dose to individuals 12 years of age and older who have completed a primary series with Pfizer-BioNTech COVID-19 Vaccine or COMIRNATY (COVID-19 Vaccine, mRNA); and**
 - **a single booster dose to individuals 18 years of age and older who have completed primary vaccination with a different authorized COVID-19 vaccine. The booster schedule is based on the labeling information of the vaccine used for the primary series.**
-

This Vaccine Information Fact Sheet contains information to help you understand the risks and benefits of COMIRNATY (COVID-19 Vaccine, mRNA) and the Pfizer-BioNTech COVID-19 Vaccine, which you may receive because there is currently a pandemic of COVID-19. Talk to your vaccination provider if you have questions.

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. 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 leading to death. 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 COMIRNATY (COVID-19 VACCINE, mRNA) AND HOW IS IT RELATED TO THE PFIZER-BIONTECH COVID-19 VACCINE?

COMIRNATY (COVID-19 Vaccine, mRNA) and the Pfizer-BioNTech COVID-19 Vaccine, when prepared according to their respective instructions for use, can be used interchangeably.

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 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

HOW IS THE VACCINE GIVEN?

The Pfizer-BioNTech COVID-19 Vaccine or COMIRNATY (COVID-19 Vaccine, mRNA) will be given to you as an injection into the muscle.

Primary Series: The vaccine is administered as a 2-dose series, 3 weeks apart. A third primary series dose may be administered at least 4 weeks after the second dose to individuals who are determined to have certain kinds of immunocompromise.

Booster Dose:

- A single booster dose of the vaccine may be administered at least 5 months after completion of a primary series of the Pfizer-BioNTech COVID-19 Vaccine or COMIRNATY (COVID-19 Vaccine, mRNA) to individuals 12 years of age and older.
- A single booster dose of the vaccine may be administered to individuals 18 years of age and older who have completed primary vaccination with a different authorized COVID-19 vaccine. Please check with your healthcare provider regarding timing of the booster dose.

The vaccine may not protect everyone.

WHO SHOULD NOT GET THE VACCINE?

You should not get the 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 VACCINES?

COMIRNATY (COVID-19 Vaccine, mRNA) and the authorized formulations of the vaccine include the following ingredients:

- mRNA and 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).

Pfizer-BioNTech COVID-19 vaccines for individuals 12 years of age and older contain 1 of the following sets of additional ingredients; ask the vaccination provider which version is being administered:

- potassium chloride, monobasic potassium phosphate, sodium chloride, dibasic sodium phosphate dihydrate, and sucrose

OR

- tromethamine, tromethamine hydrochloride, and sucrose

COMIRNATY (COVID-19 Vaccine, mRNA) contains 1 of the following sets of additional ingredients; ask the vaccination provider which version is being administered:

- potassium chloride, monobasic potassium phosphate, sodium chloride, dibasic sodium phosphate dihydrate, and sucrose

OR

- tromethamine, tromethamine hydrochloride, and sucrose

HAS THE VACCINE BEEN USED BEFORE?

Yes. In clinical trials, approximately 23,000 individuals 12 years of age and older have received at least 1 dose of the vaccine. Data from these clinical trials supported the Emergency Use Authorization of the Pfizer-BioNTech COVID-19 Vaccines and the approval of COMIRNATY (COVID-19 Vaccine, mRNA). Millions of individuals have received the vaccine under EUA since December 11, 2020. The vaccine that is authorized for use in individuals 12 years of age and older includes two formulations; one that was studied in clinical trials and used under EUA, and one with the same mRNA and lipids but different inactive ingredients. The use of the different inactive ingredients helps stabilize the vaccine under refrigerated temperatures and the formulation can be administered without dilution.

WHAT ARE THE BENEFITS OF THE VACCINE?

The vaccine has been shown to prevent COVID-19.

The duration of protection against COVID-19 is currently unknown.

WHAT ARE THE RISKS OF THE VACCINE?

There is a remote chance that the vaccine could cause a severe allergic reaction. A severe allergic reaction would usually occur within a few minutes to 1 hour after getting a dose of the 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 vaccine, more commonly in males under 40 years of age than among females and older males.

In most of these people, symptoms began within a few days following receipt of the second dose of 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 vaccine:

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

Side effects that have been reported with the 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)
- decreased appetite
- diarrhea
- vomiting
- arm pain
- fainting in association with injection of the vaccine

These may not be all the possible side effects of the vaccine. Serious and unexpected side effects may occur. The possible side effects of the vaccine are 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 either “COMIRNATY (COVID-19 Vaccine, mRNA)” or “Pfizer-BioNTech COVID-19 Vaccine EUA”, as appropriate, 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 COMIRNATY (COVID-19 VACCINE, mRNA) OR THE PFIZER-BIONTECH COVID-19 VACCINE?

Under the EUA, it is your choice to receive or not receive the 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 COMIRNATY (COVID-19 VACCINE, mRNA) OR THE PFIZER-BIONTECH COVID-19 VACCINE?

Other vaccines to prevent COVID-19 may be available under Emergency Use Authorization.

CAN I RECEIVE THE COMIRNATY (COVID-19 VACCINE, mRNA) OR PFIZER-BIONTECH COVID-19 VACCINE AT THE SAME TIME AS OTHER VACCINES?

Data have not yet been submitted to FDA on administration of COMIRNATY (COVID-19 Vaccine, mRNA) or the Pfizer-BioNTech COVID-19 Vaccine at the same time with other vaccines. If you are considering receiving COMIRNATY (COVID-19 Vaccine, mRNA) or the Pfizer-BioNTech COVID-19 Vaccine with other vaccines, discuss your options with your healthcare provider.

WHAT IF I AM IMMUNOCOMPROMISED?

If you are immunocompromised, you may receive a third dose of the 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 VACCINE GIVE ME COVID-19?

No. The 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 next dose(s) of the 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
<p data-bbox="315 705 620 737">www.cvdvaccine.com</p> 	<p data-bbox="950 779 1222 848">1-877-829-2619 (1-877-VAX-CO19)</p>

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)?

An Emergency Use Authorization (EUA) is a mechanism to facilitate the availability and use of medical products, including vaccines, during public health emergencies, such as the current COVID-19 pandemic. An 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 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.

This EUA for the Pfizer-BioNTech COVID-19 Vaccine and COMIRNATY (COVID-19 Vaccine, mRNA) will end when the Secretary of HHS determines that the circumstances justifying the EUA no longer exist or when there is a change in the approval status of the product such that an EUA is no longer needed.



Manufactured by
Pfizer Inc., New York, NY 10017

BIONTECH

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

LAB-1451-16.0

Revised: 03 January 2022



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

GDTI: 0886983000332



**Get vaccinated.
Get your smartphone.
Get started with v-safe.**

What is v-safe?

V-safe is a smartphone-based tool that uses text messaging and web surveys to provide personalized health check-ins after you receive a COVID-19 vaccination. Through **v-safe**, you can quickly tell CDC if you have any side effects after getting the COVID-19 vaccine. Depending on your answers, someone from CDC may call to check on you. And **v-safe** will remind you to get your second COVID-19 vaccine dose if you need one.

Your participation in CDC's **v-safe** makes a difference—it helps keep COVID-19 vaccines safe.

How can I participate?

Once you get a COVID-19 vaccine, you can enroll in **v-safe** using your smartphone. Participation is voluntary and you can opt out at any time. You will receive text messages from **v-safe** around 2 p.m. local time. To opt out, simply text "STOP" when **v-safe** sends you a text message. You can also start **v-safe** again by texting "START."

How long do v-safe check-ins last?

During the first week after you get your vaccine, **v-safe** will send you a text message each day to ask how you are doing. Then you will get check-in messages once a week for up to 5 weeks. The questions **v-safe** asks should take less than 5 minutes to answer. If you need a second dose of vaccine, **v-safe** will provide a new 6-week check-in process so you can share your second-dose vaccine experience as well. You'll also receive check-ins 3, 6, and 12 months after your final dose of vaccine.

Is my health information safe?

Yes. Your personal information in **v-safe** is protected so that it stays confidential and private.*



Use your smartphone to tell CDC about any side effects after getting the COVID-19 vaccine. You'll also get reminders if you need a second vaccine dose.



Sign up with your smartphone's browser at vsafe.cdc.gov

OR

Aim your smartphone's camera at this code



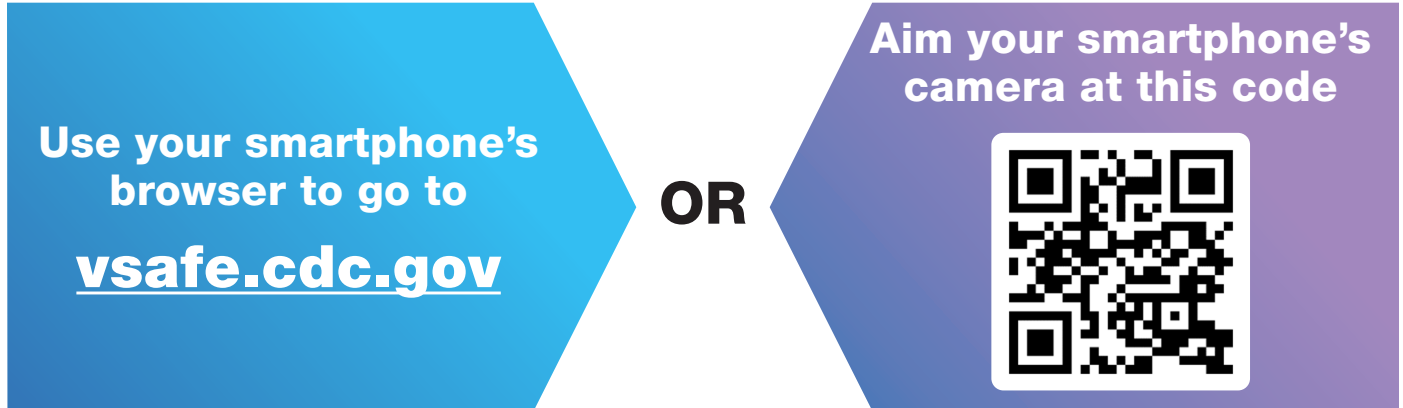
*To the extent **v-safe** uses existing information systems managed by CDC, FDA, and other federal agencies, the systems employ strict security measures appropriate for the data's level of sensitivity.

How to register and use v-safe

You will need your smartphone and information about the COVID-19 vaccine you received. This information can be found on your vaccination record card; if you cannot find your card, please contact your healthcare provider.

Register

1. Go to the **v-safe** website using one of the two options below:



2. Read the instructions. Click **Get Started**.
3. Enter your name, mobile number, and other requested information. Click **Register**.
4. You will receive a text message with a verification code on your smartphone. Enter the code in **v-safe** and click **Verify**.
5. At the top of the screen, click **Enter vaccine information**.
6. Select which COVID-19 vaccine you received (found on your vaccination record card; if you cannot find your card, please contact your healthcare provider). Then enter the date you were vaccinated. Click **Next**.
7. Review your vaccine information. If correct, click **Submit**. If not, click **Go Back**.
8. **Congrats! You're all set!** If you complete your registration before 2 p.m. local time, **v-safe** will start your initial health check-in around 2pm that day. If you register after 2 p.m., **v-safe** will start your initial health check-in immediately after you register — just follow the instructions.

You will receive a reminder text message from **v-safe** when it's time for the next check-in — around 2 p.m. local time. Just click the link in the text message to start the check-in.

Complete a v-safe health check-in

1. When you receive a **v-safe** check-in text message on your smartphone, click the link when ready.
2. Follow the instructions to complete the check-in.

Troubleshooting

How can I come back and finish a check-in later if I'm interrupted?

- Click the link in the text message reminder to restart and complete your check-in.

How do I update my vaccine information after my second COVID-19 vaccine dose?

- **V-safe** will automatically ask you to update your second dose information. Just follow the instructions.

Need help with v-safe?

Call 800-CDC-INFO (800-232-4636)

TTY 888-232-6348

Open 24 hours, 7 days a week

Visit www.cdc.gov/vsafe

