

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.

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

	ANNUAL INCOME	For You	For Spouse	For Children	For Others	Sub Total
SOURCE OF INCOME	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					\$0.00

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 6 months to 17-years-old receiving the COVID-19 vaccine.

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 and Moderna COVID-19 vaccine onsite or online at www.fda.gov 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 Variant-Specific Booster

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 Booster 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: _____

Booster 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 Variant-Specific Booster for ages 12 years and above and Emergency Use Authorization (EUA) for the Pfizer-BioNTech COVID-19 Vaccine for ages 12 years and above, 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).

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

_____ The significant known and potential risks and benefits of Pfizer-BioNTech COVID-19 Booster, 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 Bivalent Booster is administered intramuscularly as a single dose for ages 5 years of age and older. Recipients must receive all doses of the Pfizer-BioNTech COVID-19 Vaccine prior to receiving this booster dose.

_____ Recipient is 5 years of age or older and completed a primary series or last booster 2 months before today.

_____ Booster may not protect all vaccine recipients.

_____ Pfizer-BioNTech COVID-19 Vaccine, Bivalent include the following ingredients: mRNA and lipids,

tromethamine, tromethamine hydrochloride, sucrose, and sodium chloride.

_____ 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 Booster. 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 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

FACT SHEET FOR RECIPIENTS AND CAREGIVERS ABOUT PFIZER-BIONTECH COVID-19 VACCINE, BIVALENT WHICH HAS EMERGENCY USE AUTHORIZATION (EUA) TO PREVENT CORONAVIRUS DISEASE 2019 (COVID-19)

You or your child is being offered the Pfizer-BioNTech COVID-19 Vaccine, Bivalent to prevent coronavirus disease 2019 (COVID-19) which is caused by the virus SARS-CoV-2. This fact sheet contains information to help you understand the risks and benefits of the Pfizer-BioNTech COVID-19 Vaccine, Bivalent which you or your child 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.

The U.S. Food and Drug Administration (FDA) has issued an Emergency Use Authorization (EUA) to make the Pfizer-BioNTech COVID-19 Vaccine, Bivalent available during the COVID-19 pandemic (for more details about an EUA please see “**WHAT IS AN EMERGENCY USE AUTHORIZATION?**” at the end of this document). The Pfizer-BioNTech COVID-19 Vaccine, Bivalent is not an FDA-approved vaccine in the United States. Read this Fact Sheet for information about the Pfizer-BioNTech COVID-19 Vaccine, Bivalent.

WHAT IS COVID-19?

COVID-19 is caused by a coronavirus called SARS-CoV-2. You can get COVID-19 through close 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 THE PFIZER-BIONTECH COVID-19 VACCINE, BIVALENT?

The Pfizer-BioNTech COVID-19 Vaccine, Bivalent is a vaccine for use in individuals 6 months of age and older to prevent COVID-19¹. The FDA has authorized the emergency use of the Pfizer-BioNTech COVID-19 Vaccine, Bivalent under an EUA.

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

¹ The Pfizer-BioNTech COVID-19 Vaccine, Bivalent encodes the spike protein of the Original SARS-CoV-2 and the Omicron BA.4/BA.5 SARS-CoV-2.

WHAT SHOULD YOU MENTION TO THE VACCINATION PROVIDER BEFORE YOU OR YOUR CHILD GET THE PFIZER-BIONTECH COVID-19 VACCINE, BIVALENT?

Tell the vaccination provider about all your or your child's medical conditions, including if you or your child:

- 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 or your child's 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, Bivalent is given as an injection into the muscle.

Individuals 6 months through 4 years of age:

- **Unvaccinated individuals²:** Three doses of Pfizer-BioNTech COVID-19 Vaccine, Bivalent are administered over at least 11 weeks. The first two doses are administered three weeks apart. The third dose is administered at least 8 weeks after the second dose.
- **Individuals who have received one dose of the monovalent³ Pfizer-BioNTech COVID-19 Vaccine:** Two doses of Pfizer-BioNTech COVID-19 Vaccine, Bivalent are administered. The first dose of Pfizer-BioNTech COVID-19 Vaccine, Bivalent is given three weeks after the monovalent Pfizer-BioNTech COVID-19 Vaccine and the second dose at least 8 weeks later.
- **Individuals who have received two doses of the monovalent Pfizer-BioNTech COVID-19 Vaccine:** A single dose of Pfizer-BioNTech COVID-19 Vaccine, Bivalent is administered at least 8 weeks after the monovalent Pfizer-BioNTech COVID-19 Vaccine.
- **Individuals who have received three doses of the monovalent Pfizer-BioNTech COVID-19 Vaccine:** A single dose of Pfizer-BioNTech COVID-19 Vaccine, Bivalent is administered at least 2 months after the monovalent Pfizer-BioNTech COVID-19 Vaccine.

² If your child will turn 5 years old in the next 11 weeks and has not started their vaccination series, please discuss the options with your provider.

³ The Pfizer-BioNTech COVID-19 Vaccine, a monovalent vaccine, encodes the spike protein of only the Original SARS-CoV-2.

Individuals 5 years of age and older:

- **Unvaccinated individuals:** A single dose of Pfizer-BioNTech COVID-19 Vaccine, Bivalent.
- **Individuals who have received one or more doses of a monovalent COVID-19 vaccine⁴:** A single dose of Pfizer-BioNTech COVID-19 Vaccine, Bivalent is administered at least 2 months after any monovalent COVID-19 vaccine.
- **Individuals 65 years of age and older who have received one dose of a bivalent COVID-19 vaccine:** A dose of Pfizer-BioNTech COVID-19 Vaccine, Bivalent may be administered at least 4 months after the dose of the bivalent COVID-19 vaccine.
- **Immunocompromised individuals 5 years of age and older who have received one dose of a bivalent COVID-19 vaccine:** An additional dose with Pfizer-BioNTech COVID-19 Vaccine, Bivalent may be administered at least 2 months following the dose of the bivalent COVID-19 vaccine; additional doses may be administered at the discretion of the healthcare provider.

WHO SHOULD NOT GET PFIZER-BIONTECH COVID-19 VACCINE, BIVALENT?

A person should not get Pfizer-BioNTech COVID-19 Vaccine, Bivalent if they had:

- a severe allergic reaction after a previous dose of Pfizer-BioNTech COVID-19 Vaccine, Pfizer-BioNTech COVID-19 Vaccine, Bivalent, or COMIRNATY (COVID-19 Vaccine, mRNA)⁵
- a severe allergic reaction to any ingredient in these vaccines.

WHAT ARE THE INGREDIENTS IN THIS VACCINE?

Pfizer-BioNTech COVID-19 Vaccine, Bivalent contains the following ingredients: messenger ribonucleic acid (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), tromethamine, tromethamine hydrochloride, and sucrose. Pfizer-BioNTech COVID-19 Vaccine, Bivalent for individuals 6 months through 11 years of age also contains sodium chloride.

HAS THIS VACCINE BEEN USED BEFORE?

Millions of individuals 6 months of age and older have received the Pfizer-BioNTech COVID-19 Vaccine, Bivalent under EUA. In clinical trials, 60 individuals 6 months through 4 years of age, 113 individuals 5 through 11 years of age, 107 individuals 12 through 17 years of age, 103 individuals 18 through 55 years of age, and 106 individuals greater than 55 years of age received a dose of Pfizer-BioNTech COVID-19 Vaccine, Bivalent.

In addition, millions of individuals 6 months of age and older have received the monovalent Pfizer-BioNTech COVID-19 Vaccine under EUA. In a clinical trial,

⁴ Monovalent refers to a COVID-19 vaccine that contains or encodes the spike protein of only the Original SARS-CoV-2.

⁵ COMIRNATY (COVID-19 Vaccine, mRNA) is an FDA-approved COVID-19 vaccine. COMIRNATY encodes the spike protein of only the Original SARS-CoV-2.

approximately 1,200 individuals 6 months through 23 months of age, approximately 1,800 individuals 2 through 4 years of age, and approximately 3,100 individuals 5 through 11 years of age have received at least 1 dose of Pfizer-BioNTech COVID-19 Vaccine. In another clinical trial, approximately 23,000 individuals 12 years of age and older have received at least 1 dose of the Pfizer-BioNTech COVID-19 Vaccine.

The Pfizer-BioNTech COVID-19 Vaccine, Bivalent is made in the same way as the Pfizer-BioNTech COVID-19 Vaccine, but it also contains an Omicron component to help prevent COVID-19 caused by the Omicron variant of SARS-CoV-2.

WHAT ARE THE BENEFITS OF PFIZER-BIONTECH COVID-19 VACCINE, BIVALENT?

FDA has authorized the Pfizer-BioNTech COVID-19 Vaccine, Bivalent to provide protection against COVID-19.

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

WHAT ARE THE RISKS OF PFIZER-BIONTECH COVID-19 VACCINE, BIVALENT?

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 one hour after getting a dose. For this reason, the vaccination provider may ask you or your child to stay at the place where you or your child received the vaccine for monitoring after vaccination. Signs of a severe allergic reaction can include:

- Difficulty breathing
- Swelling of the face and throat
- A fast heartbeat
- A bad rash all over the 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, Bivalent, the Pfizer-BioNTech COVID-19 Vaccine, or COMIRNATY (COVID-19 Vaccine, mRNA), more commonly in adolescent males and adult males under 40 years of age than among females and older males. In most of these people, symptoms began within a few days following vaccination. The chance of having this occur is very low. You should seek medical attention right away if you or your child have any of the following symptoms after receiving the vaccine, particularly during the 2 weeks after you or your child receives a dose of either vaccine:

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

Additional symptoms, particularly in children, may include:

- Fainting
- Unusual and persistent irritability
- Unusual and persistent poor feeding

- Unusual and persistent fatigue or lack of energy
- Persistent vomiting
- Persistent pain in the abdomen
- Unusual and persistent cool, pale skin

Side effects that have been reported with Pfizer-BioNTech COVID-19 Vaccine, Bivalent, Pfizer-BioNTech COVID-19 Vaccine, or COMIRNATY (COVID-19 Vaccine, mRNA) 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/tenderness
- 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
- Dizziness
- Irritability

These may not be all the possible side effects. Serious and unexpected side effects may occur. The possible side effects are still being studied.

WHAT SHOULD I DO ABOUT SIDE EFFECTS?

If you or your child experience a severe allergic reaction, call 9-1-1, or go to the nearest hospital.

Call the vaccination provider or your or your child's healthcare provider if you or your child have any side effects that bother you or your child 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, Bivalent 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 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 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 OR NOT TO HAVE MY CHILD GET THE PFIZER-BIONTECH COVID-19 VACCINE, BIVALENT?

Under the EUA, there is an option to accept or refuse receiving this vaccine. Should you decide for yourself or your child not to receive this vaccine, it will not change the standard medical care.

ARE THERE OTHER VACCINES FOR PREVENTING COVID-19 BESIDES THE PFIZER-BIONTECH COVID-19 VACCINE, BIVALENT?

Other vaccines to prevent COVID-19 may be available under EUA, including bivalent vaccines that contain an Omicron component of SARS-CoV-2. COMIRNATY and SPIKEVAX (COVID-19 Vaccine, mRNA) are FDA-approved monovalent COVID-19 vaccines.

CAN I OR MY CHILD RECEIVE PFIZER-BIONTECH COVID-19 VACCINE, BIVALENT AT THE SAME TIME AS OTHER VACCINES?

Data have not been submitted to FDA on administration of Pfizer-BioNTech COVID-19 Vaccine, Bivalent at the same time as other vaccines. If you are considering receiving or having your child receive Pfizer-BioNTech COVID-19 Vaccine, Bivalent with other vaccines, discuss your options with your or your child’s healthcare provider.

WHAT IF I AM, OR MY CHILD IS, IMMUNOCOMPROMISED?

Immunocompromised individuals 5 years of age and older may receive one or more additional doses of Pfizer-BioNTech COVID-19 Vaccine, Bivalent (see **HOW IS THE VACCINE GIVEN?** above).

Vaccinations may not provide full immunity to COVID-19 in people who are immunocompromised; therefore, you or your child should continue to maintain physical precautions to help prevent COVID-19. Your close contacts should be vaccinated as appropriate.

WHAT ABOUT PREGNANCY OR BREASTFEEDING?

If you are, or your child is, pregnant or breastfeeding, discuss the options with your healthcare provider.

WILL THIS VACCINE GIVE ME OR MY CHILD COVID-19?

No. This vaccine does not contain SARS-CoV-2 and cannot give you or your child COVID-19.


KEEP THE VACCINATION CARD

When you, or your child, receive the first COVID-19 vaccine, you will get a vaccination card. Remember to bring the card if you or your child receive additional doses.

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="347 1043 651 1073">www.cvdvaccine.com</p> 	<p data-bbox="980 1115 1252 1184">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 VACCINATION INFORMATION BE RECORDED?

The vaccination provider may include your or your child’s vaccination information in your state/local jurisdiction’s Immunization Information System (IIS) or other designated system. 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 FDA has made Pfizer-BioNTech COVID-19 Vaccine, Bivalent available under an emergency access mechanism called an EUA. 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. A product authorized for emergency use has not undergone the same type of review by FDA as an FDA-approved product.

FDA may issue an EUA when certain criteria are met, which includes that there are no adequate, approved, and available alternatives. In addition, the FDA decision is based on the totality of the 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 under EUA during the COVID-19 pandemic.

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

BIONTECH

Manufactured for

BioNTech Manufacturing GmbH

An der Goldgrube 12

55131 Mainz, Germany



Manufactured by

Pfizer Inc., New York, NY 10001

LAB-1572-0.3

Revised: 18 April 2023



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

GDTI: 0886983000585



**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



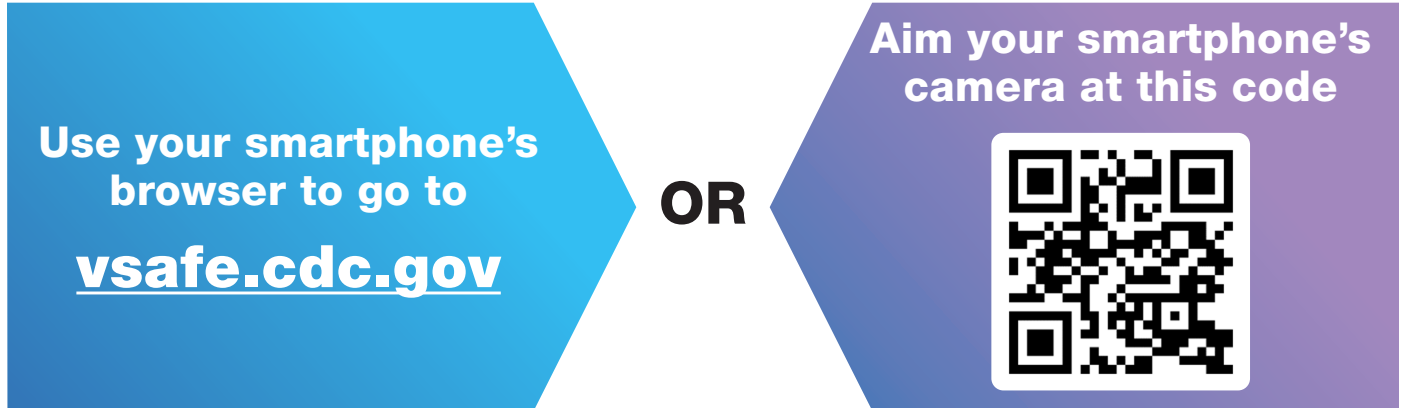
*To the extent **v-safe** uses existing information systems managed by CDC, FDA, and other federal agencies, the systems employ strict security measures appropriate for the data's level of sensitivity.

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

