

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: _____

Screening and Consent for Janssen (Johnson and Johnson) COVID-19 VACCINE 18 YEARS AND OLDER

The following questions will help us determine if there is any reason you should not get the Janssen (Johnson and Johnson) COVID-19 VACCINE today. If you answer “yes” to any question, it does not necessarily mean you should not be vaccinated. It just means additional questions must be asked. If a question is not clear, please ask your healthcare provider to explain it.

	yes	no	don't know
Have you tested positive for COVID-19 in the past 90 days:	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Do you have any of these COVID-related symptoms (fever, cough, shortness of breath, loss of taste and/or smell)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Have you been a close contact of a confirmed COVID case, in the last 14 days?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
1. Have you had a vaccine in the past 2 weeks?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
2. Are you sick today (aside from COVID symptoms)?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
3. Do you have an allergy to a component of the vaccine? <small>(Recombinant, Replication-incompetent adenovirus type 26 expressing the ARS-CoV-2 spike protein, citric acid, monohydrate, trisodium citrate dehydrate, ethanol, 2-hydroxypropyl-B-cyclodextrin (HBCD), polysorbate-80 sodium chloride)</small>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
4. Have you ever had a serious reaction to a vaccine in the past? <small>(hives, itching, difficulty breathing)</small>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
5. Have you EVER had anaphylaxis (severe, potentially life-threatening allergic reaction), NOT related to an injection?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
6. FOR FEMALES ONLY: Could you be pregnant or breastfeeding?	<input type="checkbox"/>	<input type="checkbox"/>	N/A
7. Have you ever fainted or felt lightheaded after receiving an injection or having blood drawn?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

I know the Food and Drug Administration (FDA) has authorized the emergency use of this vaccine. I know it is not a fully licensed FDA vaccine. I was asked to join the V-SAFE program. The program does health checks on the people who get the COVID-19 vaccine. I know I should report vaccine side effects to FDA/CDC Vaccine Adverse Event Reporting System (VAERS) at 1-800-822-7967 or <https://vaers.hhs.gov/reportevent.html>.

I know I should get only one dose of this COVID-19 vaccine.

I have been given a copy and have read or have had explained to me, the information in the Fact Sheet for Janssen (Johnson and Johnson) COVID-19 VACCINE. I have had a chance to ask questions which were answered to my satisfaction. I understand the benefits and risks of the vaccine.

I am aware of a remote chance of blood clots involving blood vessels in the brain, abdomen, and legs along with low levels of platelets (blood cells that help your body stop bleeding), that have occurred in some people who have received the Janssen COVID-19 Vaccine. In people who developed these blood clots and low levels of platelets, symptoms began approximately one to two-weeks following vaccination. Most people who developed these blood clots and low levels of platelets were females ages 18 through 49 years. You should seek medical attention right away if you have any of the following symptoms after receiving Janssen COVID-19 Vaccine: shortness of breath, chest pain, leg swelling, persistent abdominal pain, severe or persistent headaches or blurred vision, easy bruising or tiny blood spots under the skin beyond the site of the injection.

Vaccine: Janssen (Johnson and Johnson) COVID VACCINE Accept Immunization Decline Immunization

NAME: _____ DOB: _____ MRN: _____

PATIENT SIGNATURE: _____ Date: _____

FOR CLINIC STAFF ONLY:

WAIS reviewed – dose giving today: (date of Immunization: _____)

**** If yes to any questions, consult with medical provider or pharmacist**

MEDICAL PROVIDER/PHARMACIST SIGNATURE _____

VACCINE NAME _____ LOT # _____ EXP DATE _____ INITIALS _____

COVID screening question follow up:

1. If patient has a positive COVID test, must wait 90 days from date of positive test to receive vaccine when supplies are limited
2. if symptoms – test for COVID and wait for negative PCR to vaccinate. If positive, see #1. If negative, vaccinate.
3. If patient is a close contact of someone who tested positive for COVID within 14 days,
 - a. persons in the community or outpatient setting who have had a known COVID-19 exposure should not seek vaccination until their [quarantine period](#) has ended to avoid potentially exposing healthcare personnel and other persons to SARS-CoV-2 during the vaccination visit.
 - b. Healthcare workers and first responders – vaccinate

Have you had a vaccine in the past 2 weeks?

- Given the lack of data on the safety and efficacy of the Janssen (Johnson and Johnson) COVID-19 vaccine administered simultaneously with other vaccines, the Janssen (Johnson and Johnson) vaccine series should be administered alone, with a minimum interval of 14 days before or after administration with any other vaccines. If Janssen (Johnson and Johnson) COVID-19 vaccine is inadvertently administered within 14 days of another vaccine, doses do not need to be repeated for either vaccine.
- Watch for new hire vaccinations and prioritize COVID vaccine above others.

Is the person to be vaccinated sick today (aside from COVID symptoms)?

- Aside from COVID symptoms – evaluate as would normally for vaccines (afebrile, etc.)
- Provider must assess situation and sign off

Does the person to be vaccinated have an allergy to a component of the vaccine?

(Recombinant, Replication-incompetent adenovirus type 26 expressing the ARS-CoV-2 spike protein, citric acid, monohydrate, trisodium citrate dehydrate, ethanol, 2-hydroxypropyl-β-cyclodextrin (HBCD), polysorbate-80 sodium chloride)

Has the person to be vaccinated ever had a serious reaction to a vaccine in the past? (hives, itching, difficulty breathing)

- **Do not vaccinate at this time, per ACIP**

Has the person to be vaccinated EVER had anaphylaxis to a NON-injectable agent?

- If yes – must observe for 30 minutes post-dose vs 15 minutes
- If it was to an injectable (vaccine, drug) – DO NOT VACCINATE

Is the person to be vaccinated pregnant or breastfeeding? Patient must make informed decision; no thimerosal

- Available data on Janssen (Johnson and Johnson) COVID-19 Vaccine administered to pregnant women are insufficient to inform vaccine-associated risks in pregnancy.
- Data are not available to assess the effects of Janssen (Johnson and Johnson) COVID-19 Vaccine on the breastfed infant or on milk production/excretion.
- CDC recommends vaccination in both breastfeeding and lactation if appropriate ([link below](#)).

If the person has ever felt faint or lightheaded after receiving an injection or having blood drawn?

- The recommendation is to accommodate them in a room with appropriate positioning to decrease the chances of a vasovagal syncope reaction

Reference:

<https://www.cdc.gov/vaccines/covid-19/info-by-product/janssen/index.html>

<https://www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/janssen-covid-19-vaccine>

FACT SHEET FOR RECIPIENTS AND CAREGIVERS

EMERGENCY USE AUTHORIZATION (EUA) OF THE JANSSEN COVID-19 VACCINE TO PREVENT CORONAVIRUS DISEASE 2019 (COVID-19) IN INDIVIDUALS 18 YEARS OF AGE AND OLDER

You are being offered the Janssen 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 receiving the Janssen COVID-19 Vaccine, which you may receive because there is currently a pandemic of COVID-19.

The Janssen COVID-19 Vaccine may prevent you from getting COVID-19.

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

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

- A single dose primary vaccination to individuals 18 years of age and older.
- A single booster dose to individuals 18 years of age and older who have completed a primary vaccination with the Janssen COVID-19 Vaccine.
- A single booster dose to individuals 18 years of age and older who have completed primary vaccination with a different authorized or approved COVID-19 vaccine.

The Janssen COVID-19 Vaccine may not protect everyone.

This Fact Sheet may have been updated. For the most recent Fact Sheet, please visit www.janssencovid19vaccine.com.

WHAT YOU NEED TO KNOW BEFORE YOU GET THIS VACCINE

WHAT IS COVID-19?

COVID-19 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. Common 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 JANSSEN COVID-19 VACCINE?

The Janssen COVID-19 Vaccine is an unapproved vaccine that may prevent COVID-19.

The FDA has authorized the emergency use of the Janssen COVID-19 Vaccine to prevent COVID-19 in individuals 18 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 JANSSEN COVID-19 VACCINE?

Tell the vaccination provider about all of your medical conditions, including if you:

- have any allergies,
- 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 JANSSEN COVID-19 VACCINE?

FDA has authorized the emergency use of the Janssen COVID-19 Vaccine in individuals 18 years of age and older.

WHO SHOULD NOT GET THE JANSSEN COVID-19 VACCINE?

You should not get the Janssen 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.
- had a blood clot along with a low level of platelets (blood cells that help your body stop bleeding) following Janssen COVID-19 Vaccine or following AstraZeneca’s COVID-19 vaccine (not authorized or approved in the United States).

WHAT ARE THE INGREDIENTS IN THE JANSSEN COVID-19 VACCINE?

The Janssen COVID-19 Vaccine includes the following ingredients: recombinant, replication-incompetent adenovirus type 26 expressing the SARS-CoV-2 spike protein, citric acid

monohydrate, trisodium citrate dihydrate, ethanol, 2-hydroxypropyl- β -cyclodextrin (HBCD), polysorbate-80, sodium chloride.

HOW IS THE JANSSEN COVID -19 VACCINE GIVEN?

The Janssen COVID-19 Vaccine will be given to you as an injection into the muscle.

Primary Vaccination

The Janssen COVID-19 Vaccine is administered as a **single dose**.

Booster Dose

- A single booster dose of the Janssen COVID-19 Vaccine may be administered at least two months after primary vaccination with the Janssen COVID-19 Vaccine.
- A single booster dose of the Janssen COVID-19 Vaccine may be administered to individuals 18 years of age and older who have completed primary vaccination with a different authorized or approved COVID-19 vaccine. Please check with your health care provider regarding timing of the booster dose.

HAS THE JANSSEN COVID-19 VACCINE BEEN USED BEFORE?

The Janssen COVID-19 Vaccine is an unapproved vaccine. In clinical trials, more than 61,000 individuals 18 years of age and older have received the Janssen COVID-19 Vaccine. Millions of individuals have received the vaccine under EUA since February 27, 2021.

WHAT ARE THE BENEFITS OF THE JANSSEN COVID-19 VACCINE?

The Janssen COVID-19 Vaccine has been shown to prevent COVID-19. The duration of protection against COVID-19 is currently unknown.

WHAT ARE THE RISKS OF THE JANSSEN COVID-19 VACCINE?

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

- Injection site reactions: pain, redness of the skin and swelling.
- General side effects: headache, feeling very tired, muscle aches, nausea, and fever.
- Swollen lymph nodes.
- Blood clots.
- Unusual feeling in the skin (such as tingling or a crawling feeling) (paresthesia), decreased feeling or sensitivity, especially in the skin (hypoesthesia).
- Persistent ringing in the ears (tinnitus).

- Diarrhea, vomiting.

Severe Allergic Reactions

There is a remote chance that the Janssen 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 Janssen 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.

Blood Clots with Low Levels of Platelets

Blood clots involving blood vessels in the brain, lungs, abdomen, and legs along with low levels of platelets (blood cells that help your body stop bleeding), have occurred in some people who have received the Janssen COVID-19 Vaccine. In people who developed these blood clots and low levels of platelets, symptoms began approximately one to two weeks after vaccination. Blood clots with low levels of platelets following the Janssen COVID-19 Vaccine have been reported in males and females, across a wide age range of individuals 18 years and older; reporting has been highest in females ages 30 through 49 years (about 1 case for every 100,000 vaccine doses administered), and about 1 out of every 7 cases has been fatal. You should seek medical attention right away if you have any of the following symptoms after receiving the Janssen COVID-19 Vaccine:

- Shortness of breath,
- Chest pain,
- Leg swelling,
- Persistent abdominal pain,
- Severe or persistent headaches or blurred vision,
- Easy bruising or tiny blood spots under the skin beyond the site of the injection.

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

Guillain Barré Syndrome

Guillain Barré syndrome (a neurological disorder in which the body’s immune system damages nerve cells, causing muscle weakness and sometimes paralysis) has occurred in some people who have received the Janssen COVID-19 Vaccine. In most of these people, symptoms began within 42 days following receipt of the Janssen COVID-19 Vaccine. The chance of having this occur is very low. You should seek medical attention right away if you develop any of the following symptoms after receiving the Janssen COVID-19 Vaccine:

- Weakness or tingling sensations, especially in the legs or arms, that’s worsening and spreading to other parts of the body.
- Difficulty walking.
- Difficulty with facial movements, including speaking, chewing, or swallowing.
- Double vision or inability to move eyes.
- Difficulty with bladder control or bowel function.

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 “Janssen COVID-19 Vaccine EUA” in the first line of box #18 of the report form.

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

e-mail	Fax number	Telephone numbers
JNJvaccineAE@its.jnj.com	215-293-9955	US Toll Free: 1-800-565-4008 US Toll: (908) 455-9922

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 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 JANSSEN COVID-19 VACCINE?

It is your choice to receive or not receive the Janssen 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 JANSSEN COVID-19 VACCINE?

Another choice for preventing COVID-19 is Comirnaty, an FDA-approved COVID-19 vaccine. Other vaccines to prevent COVID-19 may be available under Emergency Use Authorization.

CAN I RECEIVE THE JANSSEN COVID-19 VACCINE AT THE SAME TIME AS OTHER VACCINES?

Data have not yet been submitted to FDA on administration of the Janssen COVID-19 Vaccine at the same time as other vaccines. If you are considering receiving the Janssen COVID-19 Vaccine with other vaccines, discuss your options with your healthcare provider.

WHAT IF I AM PREGNANT OR BREASTFEEDING?

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

WILL THE JANSSEN COVID-19 VACCINE GIVE ME COVID-19?


No. The Janssen COVID-19 Vaccine does not contain SARS-CoV-2 and cannot give you COVID-19.

KEEP YOUR VACCINATION CARD

When you receive the Janssen COVID-19 Vaccine, you will get a vaccination card to document the name of the vaccine and date of when you received the vaccine.

ADDITIONAL INFORMATION

If you have questions or to access the most recent Janssen COVID-19 Vaccine Fact Sheets, scan the QR code using your device, visit the website or call the telephone numbers provided below.

QR Code	Fact Sheets Website	Telephone numbers
	www.janssencovid19vaccine.com .	US Toll Free: 1-800-565-4008 US Toll: (908) 455-9922

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. 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, 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 TIPS.HHS.GOV.

WHAT IS THE COUNTERMEASURE 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 for 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 Janssen 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 Janssen 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, and 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 during the COVID-19 pandemic.

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

Manufactured by:
Janssen Biotech, Inc.
a Janssen Pharmaceutical Company of Johnson & Johnson
Horsham, PA 19044, USA



For more information, call US Toll Free: 1-800-565-4008, US Toll: (908) 455-9922 or go to www.janssencovid19vaccine.com

Revised: 12/14/2021



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

GDTI: 0886983000363



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



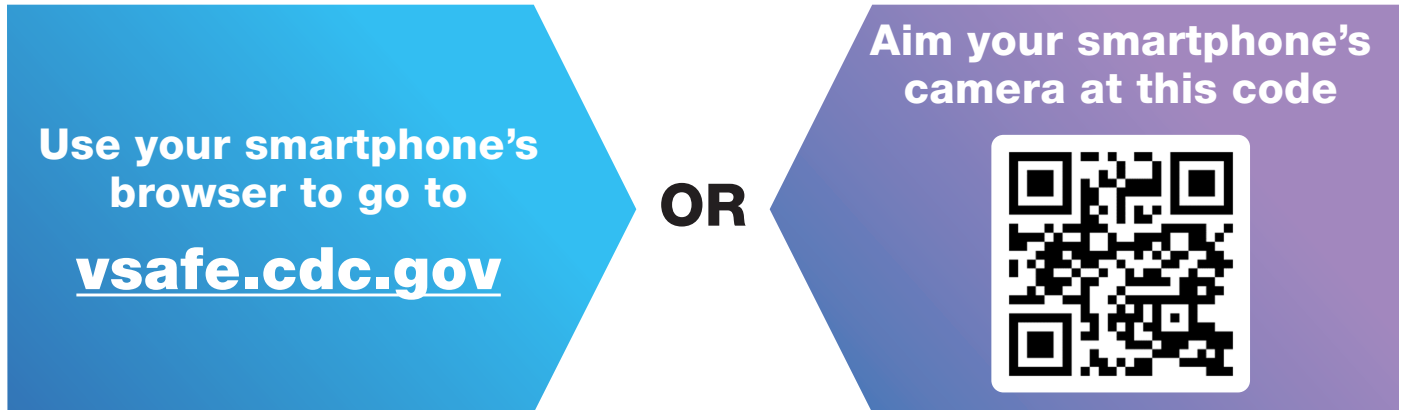
*To the extent **v-safe** uses existing information systems managed by CDC, FDA, and other federal agencies, the systems employ strict security measures appropriate for the data's level of sensitivity.

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

