

SMCHS FLU & PFIZER VACCINATION CLINIC

<https://www.signupgenius.com/go/30E0D4FACAA2CA13-smchs1>

Reserve your flu shot and/or Pfizer vaccination by Monday, September 20 at 10:00pm

Date: **Friday, September 24, 2021**

Time: **12:00 pm - 1:30 pm**

Location: **SMCHS Nurse's Office / Administration Building**

Pavilions Pharmacy will be on campus to provide flu shots and Pfizer vaccinations for SMCHS staff and students on Friday, September 24th.

Please come to the clinic with the following forms:

1. **COVID-19 screening form**
2. **COPY of the driver's license or ID card of the insur card holder and person receiving vaccination**
3. **COPY of the front and back of your PHARMACY insurance card**
4. **Copy of the front and back of your Medicare card if you have reserved a High Dose Influenza Vaccine.**

The CDC recommends that everyone 6 months of age and older get a flu vaccine every year. Remember that flu vaccine not only protects you, but it also can help protect those around you. Please help us keep our school healthy! For more information visit www.cdc.gov/flu

FLU SHOT: Quadrivalent influenza (flu) vaccine will be offered. It is designed to protect against four different flu viruses, including two influenza A viruses and two influenza B viruses. Most insurance plans cover 100% of the cost of the flu shot; the cash cost is \$44.00 by check or credit card.

HIGH DOSE FLU SHOT FOR 65+: A High Dose Influenza Vaccine (Flublock HD/Fluad) is indicated for adults 65 years and older. If you would like to reserve a dose of this vaccine, please write "High Dose" in the note section of the SignUp Genius or call the Nurse's office at (949) 279-9236. Most insurance plans cover 100% of the cost of the flu shot; the cash cost is \$73.00.

PFIZER COVID VACCINATION: Pfizer vaccine first and second doses will be offered.

A third dose is available for immunocompromised people*. Not all people who are immunocompromised qualify for the booster. Anyone that states they qualify for a third booster dose will have to attest to a document and will need to bring in their COVID vaccine card. *CDC recommendations are subject to change

The Pfizer booster for those who have had 2 doses of Pfizer or Moderna, has not been approved at this time. *CDC recommendations are subject to change

At this time, Orange County Catholic Schools are not requiring that all students and staff be vaccinated against COVID-19. Diocese of Orange Bishops' Statement <https://www.rcbo.org/diocese-of-orange-bishops-statement-vaccinate-to-protect-health-of-yourself-and-your-community/> . Updated CDPH guidelines state that fully vaccinated students may remain in school and avoid interruptions to in-person education, even if they are exposed to someone with COVID-19, so long as they remain without symptoms.

PFIZER ADMINISTRATION REQUIREMENTS FOR STUDENTS - PLEASE READ

The two-dose Pfizer vaccine is now fully approved for students age 16-17. They do NOT need a parent/guardian present but the consent form MUST be signed by the parent/guardian with full name printed next to the signature.

Students age 12-15 will need to have a parent/guardian present.

If you have any questions, please contact the Nurse's Office at (949) 766-6029

Attachments:

Inactivated Vaccine Universal Consent Form 2021

Who Needs an Additional COVID-19 Vaccine

Pfizer EUA Fact sheet for Recipients - Pfizer-BioNTech COVID-19 Vaccine_FINAL

Informed Consent for Immunization with Inactivated Vaccine

M F Other

Last Name	First Name	Middle	Date of Birth () -)
Home Address	City	State	Zip <input type="checkbox"/> Home <input type="checkbox"/> Cell

Medicare Part B ID#: _____ Last 4 digits of SSN: _____ Driver's License #: _____

Race: Asian Black or African American Hispanic American Indian Caucasian Pacific Islander Two or More Other: _____

Ethnicity: Hispanic or Latino Non-Hispanic or Latino Decline to State (Unknown)

Vaccine(s) requested: Flu COVID-19 Pneumonia Shingles Tetanus Other: (Please Specify) _____

Which arm do you prefer for vaccine? Enter weight IF LESS than 66 pounds: _____ lbs. Primary Care Provider Name: _____
(Please circle) Left Right Primary Care Provider Address: _____

Screening Questions – NOTE: IF COMPLETED ONLINE, REVIEW ANSWERS WITH PATIENT TO ENSURE NO CHANGES		Yes	No
1.	Are you sick today?	<input type="checkbox"/>	<input type="checkbox"/>
2.	Do you have a serious allergy to ANY medications, food, pet, environmental allergens, oral medication or latex? (e.g. eggs, gelatin, thimerosal, neomycin, gentamicin, polyethylene glycol (PEG), polysorbate etc.)? If yes, please list: _____	<input type="checkbox"/>	<input type="checkbox"/>
3.	Have you ever had a serious reaction or fainted after receiving any vaccination or injectable medication?	<input type="checkbox"/>	<input type="checkbox"/>
4.	Have you ever received a dose of COVID-19 vaccine? (COVID-19 only) If yes, which product did you receive? <input type="checkbox"/> Pfizer <input type="checkbox"/> Moderna <input type="checkbox"/> J&J Date: _____	<input type="checkbox"/>	<input type="checkbox"/>
5.	Have you received passive antibody therapy (monoclonal antibodies or convalescent serum) as a treatment for COVID-19 within the last 90 days? (COVID-19 only)	<input type="checkbox"/>	<input type="checkbox"/>
6.	Do you have a seizure disorder or a brain disorder? (Tdap only)	<input type="checkbox"/>	<input type="checkbox"/>
7.	Do you have a medical condition or take medication(s) that may weaken your immune system? If yes, please list: _____	<input type="checkbox"/>	<input type="checkbox"/>
8.	For women: Are you pregnant or are you considering becoming pregnant in the next month?	<input type="checkbox"/>	<input type="checkbox"/>
Immunization Needs		Yes	No
9.	Please check all that apply to you: <input type="checkbox"/> Asthma <input type="checkbox"/> Diabetes <input type="checkbox"/> Heart Disease <input type="checkbox"/> Tobacco Smoker <input type="checkbox"/> 65 Years or older. - If you checked any of the above, have you ever received a PNEUMONIA vaccine? If yes, when? _____	<input type="checkbox"/>	<input type="checkbox"/>
10.	Patients 50 and older: Have you ever received the SHINGLES vaccine?	<input type="checkbox"/>	<input type="checkbox"/>
11.	How many years has it been since your last TETANUS vaccine?	_____ yrs	<input type="checkbox"/>
12.	Patients 45 and under: Have you received the HPV (Human Papillomavirus) vaccine?	<input type="checkbox"/>	<input type="checkbox"/>
13.	Patients aged 11 to 23: Have you received a meningitis vaccine?	<input type="checkbox"/>	<input type="checkbox"/>
14.	Please indicate which vaccine(s) you would like more information about? <input type="checkbox"/> Hepatitis A <input type="checkbox"/> Hepatitis B <input type="checkbox"/> MMR (Measles, Mumps, Rubella) <input type="checkbox"/> Travel Vaccines <input type="checkbox"/> Other: _____	<input type="checkbox"/>	<input type="checkbox"/>

Informed Consent: Please read and sign.

By my signature below, I consent to the administration of the vaccine(s) by a pharmacist or a supervised student pharmacist or technician, or other authorized person, where permitted by law or state/federal guidance, employed or contracted by Albertsons Companies or one of its affiliated pharmacies and to be contacted at the number provided above regarding other immunizations for which I am due or eligible to receive. The above information is true and correct. I attest I meet eligibility criteria for the vaccination (if any); if I am the parent/guardian of the minor patient, I attest the minor patient meets eligibility criteria for the vaccination. I also release Albertsons Companies and its subsidiaries, affiliates, officers, directors, employees, and agents from all liability, including acts of omission or commission, resulting, or arising from my receipt or the minor's receipt of this vaccination. I understand that: 1) I have voluntarily chosen to receive the vaccination and understand that I am obligated to pay for all products and services received, if applicable. 2) I may be responsible for payment after the date of service if the product or service is billed to my medical benefit. 3) I am of legal age and authorized to execute this consent form or I am the parent/guardian of the minor patient. 4) I will immediately alert the pharmacist of any medical conditions which may adversely affect my personal health or effectiveness of the vaccine. 5) I have been counseled about potential side effects after vaccination, when they may occur, and when and where I should seek treatment. I am responsible for following up with my physician at my expense if I experience any side effects. 6) I should remain in the area for observation for 15 minutes unless I have a history of an immediate allergic reaction of any severity to a vaccine or injectable therapy or if I have a history of anaphylaxis due to any cause I should remain in the area for observation for 30 minutes after the vaccination. If I leave the area without waiting, I acknowledge that I am doing so at my own risk and against the advice of the professional who administered the vaccine. 7) I have read, or have had read to me, the Vaccine Information Statement(s) ("VIS") or Emergency Use Authorization ("EUA") provided for the vaccine(s) to be administered. I have had the opportunity to ask questions, and all my questions have been answered to my satisfaction. I understand the benefits and risks of the vaccine(s). 8) I have been offered and/or provided a copy of the company's Notice of Privacy Practices in compliance with the Health Insurance Portability and Accountability Act (HIPAA). 9) This vaccination, including any vaccination granted additional privacy protections under state or federal law, is subject to reporting by my pharmacy or its business associate to an immunization registry, which may share my immunization data with others, and to my primary care physician, the authorizing physician, or the local Department of Health, if applicable, and I authorize these disclosures. (New Jersey Only: I authorize _____ do not authorize _____ reporting of my receipt of this vaccination to my primary care provider I understand that failure to check authorize/do not authorize will serve as authorization.) (South Dakota and Massachusetts only: I understand I have the right to object to the sharing of my data to the above-mentioned parties through such registries.)

X

Signature of Patient or Parent/Guardian of Minor Patient

Date

For Pharmacy Use Only

Vaccine Name	Lot #	Expiration Date	Manufacturer	Dose (ml)	Dose #	Route	Site (circle)	VIS/EUA Publication Date
							<input type="checkbox"/> R / <input type="checkbox"/> L Deltoid	
							<input type="checkbox"/> R / <input type="checkbox"/> L Deltoid	
							<input type="checkbox"/> R / <input type="checkbox"/> L Deltoid	

Name of Administrator: _____ Administration Date: _____ NPP Offered RPh Counseling (Please circle): Accepted / Declined

RPh Signature [Indicates (1) VIS/EUA Provided (2) Counseling Offered and (3) Patient Eligibility Verified]: _____

WA ONLY: Substitution Permitted: _____ Dispense as Written: _____

RxBIN: _____ PCN: _____ Group #: _____ ID#: _____

Medical (Name, ID#, Group#, Payer ID - if UHC): _____

Billing Info (off-site only) Clinic Name: _____ Clinic Address: _____

Currently, CDC is recommending that moderately to severely immunocompromised people receive an additional dose. This includes people who have:

- Been receiving active cancer treatment for tumors or cancers of the blood
- Received an organ transplant and are taking medicine to suppress the immune system
- Received a stem cell transplant within the last 2 years or are taking medicine to suppress the immune system
- Moderate or severe primary immunodeficiency (such as DiGeorge syndrome, Wiskott-Aldrich syndrome)
- Advanced or untreated HIV infection
- Active treatment with high-dose corticosteroids or other drugs that may suppress your immune response

People should talk to their healthcare provider about their medical condition, and whether getting an additional dose is appropriate for them.

FACT SHEET FOR RECIPIENTS AND CAREGIVERS

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

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

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

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

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

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

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

WHAT YOU NEED TO KNOW BEFORE YOU GET THIS VACCINE

WHAT IS COVID-19?

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

WHAT IS THE PFIZER-BIONTECH COVID-19 VACCINE?

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

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

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

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

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

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

WHO SHOULD GET THE PFIZER-BIONTECH COVID-19 VACCINE?

FDA has authorized the emergency use of the Pfizer-BioNTech COVID-19 Vaccine in individuals 12 years of age and older.

WHO SHOULD NOT GET THE PFIZER-BIONTECH COVID-19 VACCINE?

You should not get the Pfizer-BioNTech COVID-19 Vaccine if you:

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

WHAT ARE THE INGREDIENTS IN THE PFIZER-BIONTECH COVID-19 VACCINE?

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

HOW IS THE PFIZER-BIONTECH COVID-19 VACCINE GIVEN?

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

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

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

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

HAS THE PFIZER-BIONTECH COVID-19 VACCINE BEEN USED BEFORE?

The Pfizer-BioNTech COVID-19 Vaccine is an unapproved vaccine. In clinical trials, approximately 23,000 individuals 12 years of age and older have received at least 1 dose of the Pfizer-BioNTech COVID-19 Vaccine.

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

In an ongoing clinical trial, the Pfizer-BioNTech COVID-19 Vaccine has been shown to prevent COVID-19 following 2 doses given 3 weeks apart. The duration of protection against COVID-19 is currently unknown.

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

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

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

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

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

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

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

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

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

WHAT SHOULD I DO ABOUT SIDE EFFECTS?

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

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

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

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

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

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

WHAT IF I DECIDE NOT TO GET THE PFIZER-BIONTECH COVID-19 VACCINE?

It is your choice to receive or not receive the Pfizer-BioNTech COVID-19 Vaccine. Should you decide not to receive it, it will not change your standard medical care.

ARE OTHER CHOICES AVAILABLE FOR PREVENTING COVID-19 BESIDES PFIZER-BIONTECH COVID-19 VACCINE?

Currently, there is no approved alternative vaccine available for prevention of COVID-19. Other vaccines to prevent COVID-19 may be available under Emergency Use Authorization.

CAN I RECEIVE THE PFIZER-BIONTECH COVID-19 VACCINE WITH OTHER VACCINES?

There is no information on the use of the Pfizer-BioNTech COVID-19 Vaccine with other vaccines.

WHAT IF I AM IMMUNOCOMPROMISED?

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

WHAT IF I AM PREGNANT OR BREASTFEEDING?

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

WILL THE PFIZER-BIONTECH COVID-19 VACCINE GIVE ME COVID-19?

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

KEEP YOUR VACCINATION CARD

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

ADDITIONAL INFORMATION

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

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

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

HOW CAN I LEARN MORE?

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

WHERE WILL MY VACCINATION INFORMATION BE RECORDED?

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

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

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

WHERE CAN I REPORT CASES OF SUSPECTED FRAUD?

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

WHAT IS THE COUNTERMEASURES INJURY COMPENSATION PROGRAM?

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

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

WHAT IS AN EMERGENCY USE AUTHORIZATION (EUA)?

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

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

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



Manufactured by
Pfizer Inc., New York, NY 10017

BIONTECH

Manufactured for
BioNTech Manufacturing GmbH
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55131 Mainz, Germany

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

Barcode Date: 05/2021