Prevaccination Checklist for COVID-19 Vaccines

For vaccine recipients:

Patient Name _____

Age _____

The following questions will help us determine if there is any reason you should not get the 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 may be asked.
If a question is not clear, please ask your healthcare provider to explain it.

	Yes	No	Don't know
1. Are you feeling sick today?	☐	☐	☐
2. Have you ever received a dose of COVID-19 vaccine?	☐	☐	☐
• If yes, which vaccine product did you receive? ☐ Pfizer ☐ Moderna ☐ Another product _____			
3. Have you ever had an allergic reaction to: (This would include a severe allergic reaction [e.g., anaphylaxis] that required treatment with epinephrine or EpiPen® or that caused you to go to the hospital. It would also include an allergic reaction that occurred within 4 hours that caused hives, swelling, or respiratory distress, including wheezing.)			
• A component of the COVID-19 vaccine, including polyethylene glycol (PEG), which is found in some medications, such as laxatives and preparations for colonoscopy procedures	☐	☐	☐
• Polysorbate	☐	☐	☐
• A previous dose of COVID-19 vaccine	☐	☐	☐
4. Have you ever had an allergic reaction to another vaccine (other than COVID-19 vaccine) or an injectable medication? (This would include a severe allergic reaction [e.g., anaphylaxis] that required treatment with epinephrine or EpiPen® or that caused you to go to the hospital. It would also include an allergic reaction that occurred within 4 hours that caused hives, swelling, or respiratory distress, including wheezing.)	☐	☐	☐
5. Have you ever had a severe allergic reaction (e.g., anaphylaxis) to something other than a component of COVID-19 vaccine, polysorbate, or any vaccine or injectable medication? This would include food, pet, environmental, or oral medication allergies.	☐	☐	☐
6. Have you received any vaccine in the last 14 days?	☐	☐	☐
7. Have you ever had a positive test for COVID-19 or has a doctor ever told you that you had COVID-19?	☐	☐	☐
8. Have you received passive antibody therapy (monoclonal antibodies or convalescent serum) as treatment for COVID-19?	☐	☐	☐
9. Do you have a weakened immune system caused by something such as HIV infection or cancer or do you take immunosuppressive drugs or therapies?	☐	☐	☐
10. Do you have a bleeding disorder or are you taking a blood thinner?	☐	☐	☐
11. Are you pregnant or breastfeeding?	☐	☐	☐

Form reviewed by _____ **Date** _____

Adapted with appreciation from the Immunization Action Coalition (IAC) screening checklists

Prevaccination Checklist for COVID-19 Vaccines

Information for Healthcare Professionals

For additional information on COVID-19 vaccine clinical guidance, see: https://www.cdc.gov/vaccines/covid-19/info-by-product/clinical-considerations.html.

For additional information on ACIP general recommendations, see: https://www.cdc.gov/vaccines/hcp/acip-recs/general-recs/index.html.

Two COVID-19 vaccines are currently authorized for use in the United States. These vaccines are authorized for use among different age groups.

PRODUCT	AUTHORIZED AGE GROUPS
Pfizer-BioNTech COVID-19 Vaccine	16 years of age and older
Moderna COVID-19 Vaccine	18 years of age and older

Anyone outside of the authorized age groups for a product should not receive the vaccine.

Postvaccination Observation Times for Persons without Contraindications to COVID-19 Vaccination

- **30 minutes:** Persons with a history of an immediate allergic reaction of any severity to a vaccine or injectable therapy or a history of anaphylaxis due to any cause
- **15 minutes:** All other persons

Are you feeling sick today?

There is no evidence that acute illness reduces vaccine efficacy or increases vaccine adverse events. However, as a precaution with moderate or severe acute illness, all vaccines should be delayed until the illness has improved. **Mild illnesses (e.g., upper respiratory infections, diarrhea) are NOT contraindications to vaccination.** Do not withhold vaccination if a person is taking antibiotics.

Vaccination of persons with current SARS-CoV-2 infection should be deferred until the person has recovered from acute illness and they can discontinue isolation. This recommendation applies to persons who develop SARS-CoV-2 infection before receiving any vaccine doses as well as those who develop SARS-CoV-2 infection after the first dose but before receipt of the second dose.

Have you ever received a dose of COVID-19 vaccine?

COVID-19 vaccines are **NOT** interchangeable. Currently authorized COVID-19 vaccines require two doses. Both doses of the series should be completed with the same product. Product dosing schedules vary.

Check medical records, immunization information systems, and vaccination record cards to help determine the initial product received. Those who received a trial vaccine should consult with the trial sponsors to determine if it is feasible to receive additional doses.

PRODUCT	**DOSING SCHEDULE** between doses 1 and 2
Pfizer-BioNTech COVID-19 Vaccine	21 days
Moderna COVID-19 Vaccine	28 days

Prevaccination Checklist for COVID-19 Vaccines

Information for Healthcare Professionals

COVID-19 Vaccine Components

Description	Pfizer-BioNTech COVID-19 vaccine	Moderna COVID-19 vaccine
mRNA	Nucleoside-modified mRNA encoding the viral spike (S) glycoprotein of SARS-CoV-2	Nucleoside-modified mRNA encoding the viral spike (S) glycoprotein of SARS-CoV-2
Lipids	2[(polyethylene glycol)-2000]-N, N-ditetradecylacetamide	PEG2000-DMG: 1,2-dimyristoyl-rac-glycerol, methoxypolyethylene glycol
	1,2-distearoyl-sn-glycero-3-phosphocholine	1,2-distearoyl-sn-glycero-3-phosphocholine
	Cholesterol	Cholesterol
	(4-hydroxybutyl)azanediyl)bis(hexane-6,1-diyl) bis(2-hexyldecanoate)	SM-102: heptadecane-9-yl 8-((2-hydroxyethyl) (6-oxo-6-(undecyloxy) hexyl) amino) octanoate
Salts, sugars, buffers	Potassium chloride	Tromethamine
	Monobasic potassium phosphate	Tromethamine hydrochloride
	Sodium chloride	Acetic acid
	Dibasic sodium phosphate dihydrate	Sodium acetate
	Sucrose	Sucrose

Have you ever had an allergic reaction to:

- **Any component of a COVID-19 vaccine, including polyethylene glycol (PEG), which is found in some medications, such as laxatives and preparations for colonoscopy procedures?**
- **Polysorbate**
- **A previous COVID-19 vaccine**

History of anaphylaxis or an immediate allergic reaction (of any severity) to any COVID-19 vaccine or any component of an mRNA COVID-19 vaccine is a contraindication to any current COVID-19 vaccine. Polyethylene glycol (PEG) is an ingredient in mRNA COVID-19 vaccines. Because of potential cross-reactive hypersensitivity with the COVID-19 vaccine ingredient PEG, a history of allergic reaction to polysorbate is also a contraindication to an mRNA COVID-19 vaccine.

Healthcare professionals should be familiar with identifying immediate-type allergic reactions, including anaphylaxis, and be competent in treating these events at the time of vaccine administration. Appropriate medical treatment for severe allergic reactions must be immediately available in the event that an acute anaphylactic reaction occurs following administration of a COVID-19 vaccine.
See Management of Anaphylaxis at COVID-19 Vaccination Sites | CDC for additional guidance.

Have you ever had an allergic reaction to another vaccine (other than COVID-19 vaccine) or another injectable medication?

A history of any immediate allergic reaction to any other vaccine or injectable therapy (i.e., intramuscular, intravenous, or subcutaneous vaccines or therapies not related to a component of mRNA COVID-19 vaccines or polysorbate) is a precaution to currently authorized COVID-19 vaccines. Vaccine may be given, but counsel patients about unknown risks of developing a severe allergic reaction and balance these risks against the benefits of vaccination. Deferral of vaccination and/or consultation with an allergist-immunologist may be considered. Considerations for vaccination include risk of exposure to SARS-CoV-2, risk of severe disease or death due to COVID-19, previous infection with COVID-19, unknown risk of anaphylaxis following mRNA COVID-19 vaccination, and ability of recipient to receive care immediately for anaphylaxis if necessary. **These individuals should be observed for 30 minutes after vaccination.**

When vaccine recipients report an immediate allergic reaction, providers should attempt to determine whether reactions reported following vaccination are consistent with immediate allergic reactions versus other types of reactions commonly observed following vaccination, such as vasovagal reaction or postvaccination side effects (which are not contraindications to receiving the second vaccine dose). See page 6 for additional information.

Have you ever had a severe allergic reaction (e.g., anaphylaxis) to something other than a component of COVID-19 vaccine, polysorbate, or any vaccine or injectable medication? This would include food, pet, venom, environmental, or oral medication allergies.

Allergic reactions, including severe allergic reactions, NOT related to vaccines or injectable therapies, components of mRNA COVID-19 vaccines (including PEG), or polysorbates are NOT contraindications or precautions to vaccination with currently authorized COVID-19 vaccines. HOWEVER, individuals who have had severe allergic reactions to anything, regardless of cause, **should be observed for 30 minutes after vaccination.** All others, including those with immediate allergic reactions that were not severe, should be observed for 15 minutes.

Clinical Consideration Questions

Responses to these questions are not (on their own) contraindications or precautions to vaccination. However, healthcare professionals should be prepared to discuss information and options with patients based on their responses to the following questions.

Have you received another vaccine in the last 14 days?

The COVID-19 vaccine series should be administered alone, with a minimum interval of 14 days before or after administration of other vaccines. This recommendation is based on the lack of data on the safety and efficacy of mRNA COVID-19 vaccines administered simultaneously with other vaccines.

Have you had a positive test for COVID-19 or has a doctor ever told you that you had COVID-19?

Vaccination should be offered to persons regardless of history of prior symptomatic or asymptomatic SARS-CoV-2 infection. Vaccination of persons with known current SARS-CoV-2 infection should be deferred until the person has recovered from the acute illness (if the person had symptoms) and criteria have been met for them to discontinue isolation. Persons with documented acute SARS-CoV-2 infection in the preceding 90 days may delay vaccination until near the end of this period, if desired, because current evidence suggests reinfection is uncommon during this time. Viral testing to assess for acute SARS-CoV-2 infection or serologic testing to assess for prior infection solely for the purpose of vaccine decision-making is not recommended.

Have you received passive antibody therapy as treatment for COVID-19?

Based on the estimated half-life of monoclonal antibodies or convalescent plasma as part of COVID-19 treatment, as well as evidence suggesting that reinfection is uncommon in the 90 days after initial infection, vaccination should be deferred for at least 90 days, as a precautionary measure until additional information becomes available, to avoid interference of the antibody treatment with vaccine-induced immune responses.

Do you have a weakened immune system caused by something such as HIV infection or cancer or do you take immunosuppressive drugs or therapies?

Persons with HIV infection or other immunocompromising conditions, or who take immunosuppressive medications or therapies, might be at increased risk for severe COVID-19. mRNA COVID-19 vaccines may be administered to persons with underlying medical conditions who have no contraindications to vaccination. However, they should be counseled about the unknown vaccine safety profile and effectiveness in immunocompromised populations, as well as the potential for reduced immune responses and the need to continue to follow all current guidance to protect themselves against COVID-19, including wearing a mask, social distancing, and washing hands frequently. Revaccination is not recommended after immune competence is regained in persons who received mRNA COVID-19 vaccines during chemotherapy or treatment with other immunosuppressive drugs.

Do you have a bleeding disorder or are you taking a blood thinner?

As with all vaccines, COVID-19 vaccine may be given to these patients, if a physician familiar with the patient's bleeding risk determines that the vaccine can be administered intramuscularly with reasonable safety. ACIP recommends the following technique for intramuscular vaccination in patients with bleeding disorders or taking blood thinners: A fine-gauge needle (23-gauge or smaller caliber) should be used for the vaccination, followed by firm pressure on the site, without rubbing, for at least 2 minutes.

Prevaccination Checklist for COVID-19 Vaccines

Information for Healthcare Professionals

Are you pregnant or breastfeeding?

If pregnant people are part of a group that is recommended to receive a COVID-19 vaccine (e.g., healthcare personnel), they may choose to be vaccinated. For pregnant people seeking guidance in making a decision, pregnant people and their healthcare providers should consider the level of COVID-19 community transmission, the patient's personal risk of contracting COVID-19, the risks of COVID-19 to the patient and potential risks to the fetus, the efficacy of the vaccine, the side effects of the vaccine, and the lack of data about use of the vaccine during pregnancy.

A lactating person who is part of a group recommended to receive a COVID-19 vaccine (e.g., healthcare personnel) may choose to be vaccinated. There are no data on the safety of COVID-19 vaccines in lactating people or the effects of mRNA COVID-19 vaccines on the breastfed infant or milk production/excretion.

Potential characteristics of allergic reactions, vasovagal reactions, and vaccine side effects following mRNA COVID-19 vaccination

In patients who develop postvaccination symptoms, determining the etiology is important to decide whether a person can receive additional doses of mRNA COVID-19 vaccines. The following table of signs and symptoms is meant to serve as a resource but may not be exhaustive, and patients may not have all signs or symptoms. Providers should use their clinical judgement when assessing patients to determine the diagnosis and management.

Characteristic	Immediate allergic reactions (including anaphylaxis)	Vasovagal reaction	Vaccine side effects (local and systemic)
Timing after vaccination	Most occur within 15-30 minutes of vaccination	Most occur within 15 minutes	Median of 1 to 3 days after vaccination (with most occurring day after vaccination)
Signs and symptoms			
Constitutional	Feeling of impending doom	Feeling warm or cold	Fever, chills, fatigue
Cutaneous	Skin symptoms present in ~90% of people with anaphylaxis, including pruritus, urticaria, flushing, angioedema	Pallor, diaphoresis, clammy skin, sensation of facial warmth	Pain, erythema or swelling at injection site, lymphadenopathy in same arm as vaccination
Neurologic	Confusion, disorientation, dizziness, lightheadedness, weakness, loss of consciousness	Dizziness, lightheadedness, syncope (often after prodromal symptoms for a few seconds or minutes), weakness, changes in vision (such as spots of flickering lights, tunnel vision), changes in hearing	Headache
Respiratory	Shortness of breath, wheezing, bronchospasm, stridor, hypoxia	Variable; if accompanied by anxiety, may have an elevated respiratory rate	N/A
Cardiovascular	Hypotension, tachycardia	Variable; may have hypotension or bradycardia during syncopal event	N/A
Gastrointestinal	Nausea, vomiting, abdominal cramps, diarrhea	Nausea, vomiting	Vomiting or diarrhea may occur
Musculoskeletal	N/A	N/A	Myalgia, arthralgia
Vaccine recommendations			
Recommended to receive second dose of mRNA COVID-19 vaccine?	No	Yes	Yes

Healthcare professionals or health departments in the United States can request a consultation from the Clinical Immunization Safety Assessment COVIDvax project for a complex COVID-19 vaccine safety question not readily addressed by CDC guidance about an individual patient residing in the United States not readily addressed by CDC guidance.

TEXAS Health and Human Services | Texas Department of State Health Services

TEXAS IMMUNIZATION REGISTRY (ImmTrac2)
ADULT CONSENT FORM

(Please print clearly)

First Name _____ Middle Name _____ Last Name _____

Date of Birth (mm/dd/yyyy) ____/____/____ Gender: ☐ Female ☐ Male Telephone ___-___-____ Email address _____

Address _____ Apartment # / Building # _____

City _____ State _____ Zip Code _____ County _____

Mother's First Name _____ Mother's Maiden Name _____

Ethnicity (Select All That apply):

☐ American Indian or Alaskan Native ☐ Asian ☐ Black or African American
☐ Native Hawaiian or Other Pacific Islander ☐ White ☐ Other Race

Race (Select Only One):

☐ Hispanic or Latino
☐ Not Hispanic or Latino

The Texas Immunization Registry is a free service of the Texas Department of State Health Services (DSHS). The immunization registry is a secure and confidential service that consolidates immunization records for public health purposes (e.g., giving all doctors treating a patient a central place to see that patient's immunization records). With your consent, your immunization information will be included in ImmTrac2. *For a family member younger than 18 years of age, a parent, legal guardian, or managing conservator may grant consent for participation for that minor by completing the ImmTrac2 Minor Consent Form (# C-7) available for downloading at www.ImmTrac.com.*

Consent for Registration and Release of Immunization Records to Authorized Persons / Entities

I understand that, by granting the consent below, I am authorizing release of my immunization information to DSHS and I further understand that DSHS will include this information in the Texas Immunization Registry. Once in ImmTrac2, my immunization information may by law be accessed by: a Texas physician, or other health care provider legally authorized to administer vaccines, for treatment of the individual as a patient; a Texas school in which the individual is enrolled; a Texas public health district or local health department, for public health purposes within their areas of jurisdiction; a state agency having legal custody of the individual; a payor, currently authorized by the Texas Department of Insurance to operate in Texas for immunization records relating to the specific individual covered under the payor's policy. **I understand that I may withdraw this consent at any time.**

State law permits the inclusion of immunization records for First Responders and their immediate family members (older than 18 years of age) in the Registry. A "First Responder" is defined as a public safety employee or volunteer whose duties include responding rapidly to an emergency. An "immediate family member" is defined as a parent, spouse, child, or sibling who resides in the same household as the First Responder. For a family member younger than 18 years of age, a parent, legal guardian, or managing conservator may grant consent for participation as an "ImmTrac2 child" by completing the Immunization Registry (ImmTrac2) Consent Form (# C-7).

Please mark the appropriate box to indicate whether you are a First Responder or an Immediate Family Member.

☐ I am a FIRST RESPONDER. ☐ I am an IMMEDIATE FAMILY MEMBER (older than 18 years of age) of a First Responder.

By my signature below, I GRANT consent for registration. I wish to INCLUDE my information in the Texas immunization registry.

Individual (or individual's legally authorized representative): Printed Name _____

Date _____ Signature _____

Privacy Notification: With few exceptions, you have the right to request and be informed about information that the State of Texas collects about you. You are entitled to receive and review the information upon request. You also have the right to ask the state agency to correct any information that is determined to be incorrect. See http://www.dshs.texas.gov for more information on Privacy Notification. (Reference: Government Code, Section 552.021, 552.023, 559.003, and 559.004)

Questions? (800) 252-9152 • (512) 776-7284 • Fax: (866) 624-0180 • www.ImmTrac.com

Texas Department of State Health Services • ImmTrac Group • MC 1946 • P. O. Box 149347 • Austin, TX 78714-9347

PROVIDERS REGISTERED WITH ImmTrac2: Please enter client information in ImmTrac2 and **affirm** that consent has been granted. **DO NOT fax to ImmTrac2. Retain this form in your client's record.**

TEXAS Health and Human Services | Texas Department of State Health Services

ImmTrac2 Immunization Registry
DISASTER INFORMATION RETENTION CONSENT FORM

(Please print clearly)

Client's Last Name

Client's First Name

Client's Middle Name

Client's Date of Birth (__/__/____)

*A parent, legal guardian or managing conservator must sign this form if the client is younger than 18 years of age.

Client's Gender: ☐ Male ☐ Female

Client's Address

Apartment #

Client's Telephone (___-___-____)

City

State

Zip Code

County

Mother's First Name (if client is younger than 18 years of age)

Mother's Maiden Name (if client is younger than 18 years of age)

ImmTrac2, the Texas immunization registry, has been designated as the disaster-related reporting and tracking system for immunizations, antivirals, and other medications administered to individuals in preparation for, or in response to, a disaster or public health emergency. From the time the event is declared over, ImmTrac2 will retain disaster-related information received from health-care providers for a period of 5 years. At the end of the 5 year retention period, client-specific disaster-related information will be removed from the Registry unless consent is granted to retain the client information in ImmTrac2 beyond the 5 year retention period.

The Texas Department of State Health Services (DSHS) encourages your voluntary participation in the Texas immunization registry.

Consent for Retention of Disaster-Related Information and Release of Information to Authorized Entities

I understand that, by granting the consent below, I am authorizing retention of my (or my child's) disaster-related information by DSHS beyond the 5 year retention period. I further understand that DSHS will include this information in the state's central immunization registry ("ImmTrac2"). Once in ImmTrac2, my (or my child's) disaster-related information may by law be accessed by:

- a state agency, for the purpose of aiding and coordinating communicable disease prevention and control efforts, and / or
- a physician or other health-care provider legally authorized to administer immunizations, antivirals, and other medications, for treating the client as a patient;

I understand that I may withdraw this consent to retain information in the ImmTrac2 Registry beyond the 5 year retention period and my consent to release information from the Registry, at any time by written communication to the Texas Department of State Health Services, ImmTrac2 Group – MC 1946, P. O. Box 149347, Austin, Texas 78714-9347.

By my signature below, I GRANT consent to retain my disaster-related information (or my child's information if younger than age 18) in the Texas immunization registry beyond the 5 year retention period.

Client (or parent, legal guardian, or managing conservator): _____

Printed Name:

Date:_____ Signature: _____

Privacy Notification: With few exceptions, you have the right to request and be informed about information that the State of Texas collects about you. You are entitled to receive and review the information upon request. You also have the right to ask the state agency to correct any information that is determined to be incorrect. See http://www.dshs.texas.gov for more information on Privacy Notification. (Reference: Government Code, Section 552.021, 552.023, 559.003, and 559.004)

Upon completion, please fax or mail form to the DSHS ImmTrac2 Group or a registered Health-care provider.

Questions? (800) 252-9152 • (512) 776-7284 • Fax: (866) 624-0180 • www.ImmTrac.com • ImmTrac2 DC

Texas Department of State Health Services • ImmTrac2 Group – MC 1946 • P. O. Box 149347 • Austin, TX 78714-9347

PROVIDERS REGISTERED WITH ImmTrac2

Please enter client information in ImmTrac2 and **affirm** that consent has been granted.

DO NOT fax to ImmTrac2. **Retain this form in your client's record.**

TEXAS Health and Human Services | Texas Department of State Health Services

ImmTrac2 Immunization Registry
FORMULARIO DE CONSENTIMIENTO DE RETENCIÓN DE INFORMACIÓN SOBRE EL DESASTRE

(Favor de escribir claramente con letra de molde)

Apellido del Cliente

Nombre del Cliente

Segundo Nombre del Cliente

Fecha de Nacimiento del Cliente

*Uno de los padres, el tutor legal o el custodio administrador debe firmar este formulario si el cliente es menor de 18 años de edad.

Género: ☐ **Masculino** ☐ **Femenino**

Dirección del Cliente

Apartamento #

Teléfono del cliente

Ciudad

Estado

Código Postal

Condado

Nombre de la Madre (si el cliente es menor de 18 años de edad)

Apellido de Soltera de la Madre (si el cliente es menor de 18 años de edad)

Nombre de la Madre (si el cliente es menor de 18 años de edad) Apellido de Soltera de la Madre (si el cliente es menor de 18 años de edad) ImmTrac2, el registro de inmunización de Texas, ha sido designado como el sistema de información y seguimiento para inmunizaciones, antivirales y otros medicamentos administrados a individuos durante la preparación o respuesta a un desastre o emergencia de salud pública. A partir del momento en que se declare finalizado el evento, ImmTrac2 retendrá la información relacionada con el desastre recibida por profesionales de la salud durante un periodo de 5 años. Al final del periodo de retención de 5 años, la información del cliente relacionada con el desastre se removerá del registro a menos que se dé el consentimiento para retener la información en ImmTrac2 más allá del periodo de retención de 5 años.

El Departamento Estatal de Servicios de Salud de Texas (DSHS) le anima a participar voluntariamente en el registro de inmunización de Texas.

Consentimiento de retención de información relacionada con el desastre y dar a conocer la información a entidades autorizadas

Entiendo que, al dar mi consentimiento a continuación, autorizo al DSHS a que retenga mi información relacionada con el desastre (o la de mi hijo[a]) más allá del periodo de retención de 5 años. Además entiendo que el DSHS incluirá esta información en el registro central de inmunización del estado ("ImmTrac2"). Una vez que mi información relacionada con el desastre (o la de mi hijo[a]) esté en ImmTrac2, puede ésta por ley ser accedida por:

- las agencias estatales, con el propósito de ayudar con los esfuerzos de prevención y control de enfermedades transmisibles y su coordinación y
- los médicos o demás profesionales de la salud legalmente autorizados para administrar vacunas, antivirales y otros medicamentos, para el tratamiento del cliente como paciente.

Entiendo que puedo retirar el consentimiento para retener información en el registro ImmTrac2 más allá del periodo de retención de 5 años y mi consentimiento para dar a conocer la información del registro, en cualquier momento mediante comunicación escrita dirigida a Texas Department of State Health Services, ImmTrac2 Group – MC 1946, P. O. Box 149347, Austin, Texas 78714-9347.

Con mi firma a continuación, DOY mi consentimiento para que se retenga mi información relacionada con el desastre (o la de mi hijo[a] si es menor de 18 años de edad) en el registro de inmunizaciones de Texas más allá del periodo de retención de 5 años.

Cliente (o padre / madre, tutor legal o custodio administrador): _____

Nombre en letra de molde

Fecha: _____ Firma: _____

Notificación Sobre Privacidad: Tan solo por unas cuantas excepciones, usted tiene el derecho de solicitar y de ser informado sobre la información que el Estado de Texas reúne sobre usted. A usted se le debe conceder el derecho de recibir y revisar la información al requerirla. Usted también tiene el derecho de pedir que la agencia estatal corrija cualquier información que se ha determinado sea incorrecta. Diríjase a *http://www.dshs.texas.gov* para más información sobre la Notificación sobre privacidad. (Referencia: Government Code, sección 552.021, 552.023, 559.003 y 559.004)

Al rellenarlo, mándelo por fax o correo postal al Grupo ImmTrac2 del DSHS o a un proveedor de salud inscrito.

¿Tiene preguntas? (800) 252-9152 • (512) 776-7284 • Fax: (866) 624-0180 • www.ImmTrac.com • ImmTrac2 DC

Texas Department of State Health Services • ImmTrac2 Group – MC 1946 • P. O. Box 149347 • Austin, TX 78714-9347

PROVIDERS REGISTERED WITH ImmTrac2

Please enter client information in ImmTrac2 and **affirm** that consent has been granted.

DO NOT fax to ImmTrac2. **Retain this form in your client's record.**

SWEENY COMMUNITY HOSPITAL
You'll Know We Care

COVID-19 VACCINE CONSENT FORM Moderna COVID-19 Vaccine

- I certify that I am: (a) the patient and at least 18 years of age; (b) the legal guardian of the patient and confirm that the patient is at least 18 years of age; or (c) authorized to consent for vaccination for the patient named above. Further, I hereby give my consent to Sweeny Community Hospital to administer the COVID-19 vaccine.
- I understand that this product has not been approved or licensed by FDA, but has been authorized for emergency use by FDA, under an EUA to prevent Coronavirus Disease 2019 (COVID-19) for use in individuals 18 years of age and older; and the emergency use of this product is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of the medical product under Section 564(b)(1) of the FD&C Act unless the declaration is terminated or authorization revoked sooner.
- I understand that it is not possible to predict all possible side effects or complications associated with receiving vaccine(s). I understand the risks and benefits associated with the above vaccine and have received, read and/or had explained to me the Emergency Use Authorization Fact Sheet on the COVID19 vaccine I have elected to receive. I also acknowledge that I have had a chance to ask questions and that such questions were answered to my satisfaction.
- I acknowledge that I have been advised to remain at the vaccination location for approximately 15 minutes after administration for observation. If I experience a severe reaction, call 9-1-1 or go to the nearest hospital.
- On behalf of myself, my heirs and personal representatives, I hereby release and hold harmless Sweeny Community Hospital, their staff, agents, successors, divisions, affiliates, subsidiaries, officers, directors, contractors and employees from any and all liabilities or claims whether known or unknown arising out of, in connection with, or in any way related to the administration of the vaccine listed above.
- I acknowledge that: (a) I understand the purposes/benefits of Texas's immunization registry and my personal immunization information will be shared with the Centers for Disease Control (CDC) or other federal agencies

_____ _____

Signature of Patient or Authorized Representative Date

_____

Print Name of Patient or Authorized Representative

_____

Signature of Sweeny Community Hospital Staff

SWEENY COMMUNITY HOSPITAL
You'll Know We Care

Medication/Vaccination Administration Monitoring Consent Form

With any medication you take, especially the first time, there is a risk for adverse reactions, or even severe allergic reactions, such as anaphylaxis (characterized by throat swelling and difficulty breathing). These reactions usually occur in the first 15 minutes after administration.

FOR YOUR SAFETY, IT IS HIGHLY RECOMMENDED THAT YOU WAIT FOR 15 MINUTES AFTER BEING VACCINATED SO OUR STAFF CAN RESPOND TO ANY POTENTIAL REACTIONS

Please initial one:

_____ I consent to stay the recommended 15 minutes

(Initial)

_____ I DO NOT consent to wait the recommended 15 minutes.

(Initial) I understand the health risks involved and I do not hold Sweeny Community Hospital liable.

_____ _____

Signature of Patient or Authorized Representative Date

_____

Signature of Sweeny Community Hospital Staff