PROVIDER COVID-19 IMMUNIZATION CONSENT FORM

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For COVID-19 Provider Use Only Clinic Name/Code:		
Location type:(clinic, health department, pharmacy, etc.,)		
Address:City:County:		
State:Zip Code: Date of Service:		
Person Receiving Vaccine:		
(Legal) First Name: MI: Last Name:		_
Date of Birth:		
1. MEDICAL HISTORY: Complete the following questions for the individual receiving the vaccine.		
If you answer "YES" you may not be able to receive the COVID-19 vaccine.		
If YES refer to Pfizer website at www.PfizerMedInfo.com. Moderna COVID-19 vaccine www.modernatx.com. Janssen		
COVID-19 vaccine www.janssencovid19vaccine.com.	*YES	NO
Refer to Pre-vaccination Checklist for COVID-19 Vaccines Information for Healthcare Professionals (cdc.gov) to		
clarify further questions: www.cdc.gov/vaccines/covid-19/downloads/pre-vaccination-screening-form.pdf. Have you had a previous COVID-19 vaccine? If yes, what type and date?		
Do you have a fever today? Are you sick today? Do you have COVID-19 infection and are currently in isolation? Are		\vdash
you currently in quarantine for known exposure to COVID-19?		
Have you ever had an allergic reaction to a COVID-19 vaccine or a COVID-19 vaccine component (including		
polyethylene glycol [PEG], which is found in some medications, or laxatives, and preparations for colonoscopy; or		
polysorbate, which is found in some vaccines, coated tablets, or IV steroids)?		
Have you ever had an immediate allergic reaction of any severity to any vaccine or injectable therapy? This would		
include an anaphylactic reaction that required treatment with epinephrine (or EpiPen) or treatment at a hospital, as well		
as an allergic reaction that occurred within 4 hours, such as difficulty breathing, hives, swelling of your face and throat,		
fast heartbeat, bad rash all over your body, dizziness, and weakness.		
Did you develop myocarditis or pericarditis after the first dose of COVID-19 vaccine? Do you have history of		
myocarditis or pericarditis prior to COVID-19 vaccination? Are you a male between age 12 through 29 years?		
Do you have a bleeding disorder or are you taking a blood thinner?		
Have you ever had a severe allergic reaction (anaphylaxis) to something other than a component of COVID-19 vaccine or any vaccine or injectable medication such as food, pet, venom, environmental, or oral medication allergies?		
Are you pregnant, breastfeeding, planning to become pregnant? Women in this group may receive any FDA-authorized		
COVID-19 vaccine. Women 18 through 49 years of age can receive any FDA-authorized COVID-19 vaccine and should		
be informed of the increased risk of thrombosis with thrombocytopenia syndrome (TTS) after receiving Janssen		
COVID-19 Vaccine and the availability of other (mRNA)COVID-19 vaccines.		
Are you immunocompromised? Do you have a condition that weakens your immune system? Are you receiving any		
immunosuppressive therapy? You are still eligible to receive Pfizer-BioNTech, Moderna, or Janssen COVID-19 vaccine		
unless you have a contraindication for some other reason. However, you will need special counseling about the vaccine.		
Have you had history of heparin-induced thrombocytopenia? If it has been 90 days or less since illness resolved, you		
may receive Pfizer-BioNTech or Moderna COVID-19 vaccine. After 90 days since illness resolved, you may be		
vaccinated with any FDA-authorized COVID-19 vaccine. Have you received monoclonal antibodies or convalescent plasma as part of COVID-19 treatment? Have you had		
Multisystem Inflammatory Syndrome (MIS)? Pfizer-BioNTech, Moderna, or Janssen COVID-19 vaccination should be		
deferred for at least 90 days to avoid interference with vaccine-induced immune responses.		
NOTE: Recipients of Janssen COVID-19 vaccine should be instructed to seek immediate medical attention if they develo	p shortn	ess
of breath, chest pain, leg pain or swelling, persistent abdominal pain, neurological symptoms (including severe or persistent		
or blurred vision), nausea, vomiting, petechiae, or easy bleeding beyond the site of vaccination within 4 to 30 days of rece	ipt of Ja	nssen
vaccine. Most people who have developed blood clots and low platelets were females ages 18 through 49 years.		
NOTE: Depending on vaccine type, a second dose of COVID-19 vaccine may be due in 21 days or 28 days after initial vaccine. Refer		
to your COVID-19 vaccination record card for second dose due date. Contact your vaccination provider, PCP, or your ADH Local		
Health Unit in 21 days or 28 days for more information. Keep your COVID-19 vaccination record card for your records for proof of		
initial vaccine date. Janssen COVID-19 vaccine is a ONE dose series.		
2. RELEASE AND ASSIGNMENT: My signature below indicates I have read, understand, and	nd agre	e to '
Please read the section on the reverse side of this form.; section 2. Release and Assignment of the COVID-19 Ir		ation
The Providers Privacy Notice is available at the clinic I Consent Form and Vaccine Recipient Emergency Use of	f	1
site or accompanies this form. Authorization Fact Sheet (EUA).		
Then sign in the box at right. Please sign here Signature of Patient/Parent/Guardian:		!
Date		

RELEASE AND ASSIGNMENT: • I have read or had explained to me the Vaccine Recipient Emergency Use Authorization (EUA) Fact Sheet for COVID-19 vaccine risks and benefits. To read the Vaccine Recipient EUA Fact Sheet for Pfizer COVID-19 vaccine, Moderna COVID-19 vaccine, or Janssen COVID-19 vaccine visit https://www.cdc.gov/vaccines/covid-19/eua/index.html or you may also visit your Local Health Unit or PCP to receive a printed copy of the EUA Fact Sheet. • I give consent to this COVID-19 provider/staff for the individual named below to be vaccinated with COVID-19 vaccine. • I hereby acknowledge that I have reviewed a copy of the Provider's Privacy Notice. • I understand that information about this COVID-19 vaccination will be included in (WebIZ) Arkansas Immunization Information System. To My Insurance Carrier(s): • I authorize the release of any medical information necessary to process my insurance claim(s). • I authorize and request payment of medical benefits directly to this COVID-19 Provider. • I agree that the authorization will cover all medical services rendered until I revoke the authorization. • I agree that the photocopy of this form may be used instead of the original. PATIENT INFORMATION: _____MI: ____ Last Name: ______ (Legal) First Name: ___ **Gender:** □ Male □ Female **Phone** #: Date of Birth: Street Address: _____ P.O. Box ____ Apt. No. ____ State: _____ Zip Code: Race: Asian Black/African American Native American /Alaska Native Native Hawaiian/Other Pacific Islander White Other **Ethnicity:** Hispanic/Latino Non-Hispanic **INSURANCE STATUS (Check appropriate box):** Patient's Relationship to Insurance Policy Holder: Self Spouse Child Other ☐ Medicaid/ARKids Number: ☐ Medicare Number: ☐ Insurance Company Name: Member ID/Policy #: REQUIRED POLICY HOLDER INFORMATION: (Legal) First Name: ____ _MI:___Last Name: ______ **Policy Holder Date of Birth:** Email Address: _____ Policy Holder's Employer Name: COVID-19 VACCINE ADMINISTRATION (Completed by staff only) Co-administration of COVID-19 vaccines and other vaccines. COVID-19 vaccines and other vaccines may be administered without regard to timing. This includes simultaneous administration of COVID-19 vaccines and other vaccines during the same visit. Other vaccines can also be administered any time before or after COVID-19 vaccination. Refrigerated COVID-19 Vaccine Ultra-cold COVID-19 Vaccine Frozen COVID-19 Vaccine ☐ Astra Zeneca ☐ Janssen (Johnson & Johnson) ☐ Pfizer-BioNTech ☐ Moderna □ Novavax-Matrix-M1 □ Other COVID-19 Vaccine _ MFG Code **Route** Site Code Dosage mL Lot Number \square IM MFG Codes: PFR=Pfizer, MOD=Moderna, ASZ=AstraZeneca, JSN=Janssen, NVX=Novavax, MSD=Merck Site Codes: Right Deltoid = RD, Left Deltoid = LD, Right Leg = RL, Left Leg = LL, Right Arm = RA, Left Arm = LA

Signature and Title of Vaccine Administrator: _____

Date Vaccine Administered: ____/__/

Revised 8/5/21/21