PROVIDER COVID-19 IMMUNIZATION CONSENT FORM

For COVID-19 Provider use only Clinic Name/Code: Location type: (clin						
Address: <u>6802 Rogers Ave Ste 2</u> City: <u>Fort Smith</u>						
State: <u>Arkansas</u> Zip Code: <u>72903</u>	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 following websites at <u>www.PfizerMedInfo.com</u> . Moderna <u>www.modernatx.com</u> . Janssen						
<u>www.janssencovid19vaccine.com.</u> Refer to Pre-vaccination Checklist for COVID-19 vaccines to clarify questions: *YESNO <u>www.cdc.gov/vaccines/covid-19/downloads/pre-vaccination-screening-form.pdf.</u>						
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 you						
currently in quarantine for known exposure to COVID-19?						
Have you ever had an allergic reaction to a COVID-19 vaccine or a COV	TD-19 vaccine component (including polyethylene					
glycol [PEG], which is found in some medications, or laxatives, and prep	parations for colonoscopy; or polysorbate, which is					
found in some vaccines, coated tablets, or IV steroids)?						
Have you ever had an allergic reaction that caused hives, swelling, respirator	y distress (including wheezing) or anaphylaxis to a					
vaccine other than COVID-19 vaccine or an injectable medication that requi						
at a hospital? Severe reaction or anaphylaxis to food, pet, venom, environme						
contraindications or precautions to vaccination with any COVID-19 vaccine						
Do you have a bleeding disorder or are you taking a blood thinner?						
Have you received a hematopoietic cell transplant (HCT) or CAR-T-cell the						
be revaccinated with a primary vaccine series at least 12 weeks after transpla						
Did you develop myocarditis or pericarditis after the first dose of COVID-19						
of any COVID-19 vaccine. If you have developed myocarditis or pericarditis	s unrelated to an mRNA COVID vaccination, may					
receive COVID-19 vaccine after the episode has completely resolved.						
Are you immunocompromised? Do you have a condition that weakens your immune system? Are you receiving any						
immunosuppressive therapy? You are eligible to receive any FDA-authorized or FDA-approved COVID-19 vaccine unless you have a contraindication for some other reason						
have a contraindication for some other reason.						
Have you had history of Heparin-Induced Thrombocytopenia (HIT) or Thrombosis with Thrombocytopenia Syndrome (TTS)? You may receive Pfizer-BioNTech or Moderna COVID-19 vaccine.						
Have you had history of Thrombosis with Thrombocytopenia Syndrome (TT	S) following Janssen or any other adenovirus-vector					
(AstraZeneca) COVID-19 vaccine? Those who developed TTS after the initia						
other adenovirus-vector COVID-19 vaccine booster dose. You may receive a mRNA COVID-19 vaccine.						
Have you received monoclonal antibodies or convalescent plasma as part of COVID-19 treatment or for post-exposure prophylaxis						
(PEP)? Defer vaccination 90 days after treatment and defer 30 days after PEP.						
Have you had Multisystem Inflammatory Syndrome (MIS)? Defer vaccination for at least 90 days. The decision for						
COVID-19 vaccination should be between the patient, their guardian, clinical team, or a specialist.						
Have you had history of Guillain-Barre Syndrome (GBS)? People with a history of GBS can receive any FDA-authorized or						
approved COVID-19 vaccine. People who had GBS after receiving Janssen vaccine should receive a Pfizer-BioNTech or Moderna						
COVID-19 vaccine booster at least 8 weeks after the Janssen dose.						
NOTE: CDC has made a clinical preference for persons 18 years and older to receive an mRNA COVID-19 vaccine over Janssen COVID-19						
vaccine. Patients who cannot or unwilling to receive an mRNA vaccine will be able to access Janssen COVID-19 vaccine. The Janssen Fact						
Sheet must be provided and explained to the recipient or parent/legal representative about the risks and benefits and address any questions or						
concerns that the recipient or parent/legal representative may have prior to the vaccination. Recipients of Janssen COVID-19 vaccine should						
seek immediate medical attention if they develop shortness of breath, chest pain, leg pain or swelling, persistent abdominal pain, severe or persistent headaches or blurred vision, easy bleeding beyond the vaccination site within 30 days of a Janssen vaccination.						
NOTE: 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 proof of initial vaccine date and 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.						
2. RELEASE AND ASSIGNMENT: Please read the section						
on the reverse side of this form. The Providers Privacy Notice	My signature below indicates I have read, understand, and					
is available at the clinic site or accompanies this form.	agree to section 2. Release and Assignment of the COVID-					
Then sign in the box at right.	19 Immunization Consent Form and Vaccine Recipient					

Please sign here

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19 Immunization Consent Form and Vaccine Recipient Emergency Use of Authorization Fact Sheet (EUA). Signature of Patient/Parent/Guardian:

Patient Information:

(Legal) First Na	me	MI			
		Date of Birth			
Age Ge	ender: M	F	_ Phone		
Street Address		P.O. Box			
City		_Zip	Stat	te	
			n Native A der White	merican / Alaska Native Other	
Ethnicity: Hispa	nic / Latino _	Nor	n-Hispanic		
Social Security I	Number				
Signature of Pat	ient / Parent	/ Guar	dian		
For Children! N	ame of Prim	arv Car	e Provider		

I request the vaccine to be given to me or to the person named above, a minor for whom I represent, and I am authorized to sign this consent form. I understand the benefits and risks of the COVID-19 vaccine as described in the Emergency Use Authorization (EUA) Fact Sheet (https://www.cdc.gov/vaccines/covid-19/eua/index.html or https://lawsdrugstore.com/forms) a copy of which I was provided with this consent form (online or in print). I have had a chance to ask questions that were answered to my satisfaction. I agree to stay in the vaccine administration area for fifteen (15) minutes or longer if indicated by the vaccine administrator after receiving my vaccine to ensure that no immediate adverse reactions occur. I understand that I will be receiving the vaccination at no cost to me. If insured, I authorize the pharmacy to bill my insurance on my behalf for the immunization – understanding that I will not incur any costs. If uninsured, I authorize the pharmacy to use my social security number, state identification number, or driver's license number to bill the United States Health Resources & Services Administration's COVID-19 Program on my behalf for the immunization – understanding that I will not incur any costs. I understand that at this time, some COVID-19 vaccines require 2 doses given 21-28 days apart dependent on the manufacturer. If this is my first dose of the COVID-19 vaccine and a second dose is required (Pfizer and Moderna only), I intend to receive a second dose of the same vaccine in accordance with the timeframe specified in the Fact Sheet to complete the series. I understand that information about this vaccination will be included in (WebIZ) Arkansa Immunization Information System. I acknowledge offer of the Laws Drug Store Notice of Privacy Practices (online at https://lawsdrugstore.com/forms or in print).

Vaccine Given (Please Check One)							
Pfizer-BioNTech Adult (12 & up) Pfizer-BioNTech Kids (5-11)			Moderna Janssen (Johnson & Johnson)				
Route	Site Code	Dosage mL	MFG Code		Lot Number		
IM							

Signature of Vaccine Administrator

Pharm/Tech

Date Vaccine Administered ____ / ____ / ____