ARKANSAS DEPARTMENT HEALTH (ADH) COVID-19 IMMUNIZATION CONSENT FORM

		7
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:		
Address:City:County:		
State: Zip Code: Date of Service:		
Person Receiving Vaccine:		1
(Legal) First Name: MI: Last Name:		
Date of Birth:		
. MEDICAL HISTORY: Complete the following questions for the individual receiving the vaccine. If you answer "YES	S" vou	_
may not be able to receive the COVID-19 vaccine.	3 you	
ADH staff: *If YES and further guidance is needed, notify your local Communicable Disease Nurse Specialist (CDNS).	*YES	NO
Have you had a previous COVID-19 vaccine? If yes, what type, number of doses and date ? Have you had a (Mpox) JYNNEOS vaccine in last 4 weeks?		
Do you have a fever today? Are you sick today? Do you have COVID-19 infection and are currently in isolation?		
Have you ever had an allergic reaction to a COVID-19 vaccine, or COVID-19 vaccine component (including polyethylene glycol [PEG], found in some medications, or laxatives, and preparations for colonoscopy; or polysorbate found in some vaccines, coated tablets, or IV steroids)?		
Have you ever had an immediate allergic reaction that caused hives, swelling, respiratory distress (including wheezing) or anaphylaxis to a vaccine other than COVID-19 vaccine or an injectable medication that required treatment with epinephrine (EpiPen) or treatment at a hospital? Severe reaction or anaphylaxis to food, pet, venom, environmental, or oral medication allergies are not contraindications or precautions to vaccination with any COVID-19 vaccine.		
Have you received a hematopoietic cell transplant (HCT) or CAR-T-cell therapy since receiving COVID-19 vaccine? Follow the COVID-19 vaccine schedule for unvaccinated people. Revaccinate starting at least 12 weeks after transplant or CAR-T-cell therapy with an age-appropriate bivalent Pfizer, Moderna, or monovalent Novavax COVID-19 vaccine.		
Did you develop myocarditis or pericarditis after any dose of COVID-19 vaccine? You should not receive a subsequent dose		
of any COVID-19 vaccine. If you have developed myocarditis or pericarditis unrelated to an mRNA COVID vaccination, you may receive age-appropriate bivalent Pfizer, Moderna, or monovalent Novavax COVID-19 vaccine after the episode has completely resolved.		
Are you immunocompromised or receiving immunosuppressive therapy? Do you have a condition that weakens your immune system? You are eligible to receive age-appropriate bivalent Pfizer, Moderna, or monovalent Novavax COVID-19 vaccine unless you have a contraindication to COVID-19 vaccine for some other reason.		
Have you had history of Heparin-Induced Thrombocytopenia (HIT) or Thrombosis with Thrombocytopenia Syndrome (TTS)? You may receive age-appropriate bivalent Pfizer, Moderna, or monovalent Novavax COVID-19 vaccine.		
Have you had history of Thrombosis with Thrombocytopenia Syndrome (TTS) following Janssen or any other adenovirus-vector (AstraZeneca) COVID-19 vaccine? Those who developed TTS after the initial Janssen vaccine should not receive a Janssen or any other adenovirus-vector COVID-19 vaccine. You may receive age-appropriate bivalent Pfizer, Moderna, or monovalent Novavax COVID-19 vaccine.		
Have you received monoclonal antibodies or convalescent plasma as part of COVID-19 treatment or for post-exposure prophylaxis (PEP)? You may receive a COVID-19 vaccine. No delay to receive a COVID-19 vaccine is necessary.		
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)? You may receive age-appropriate bivalent Pfizer, Moderna, or monovalent Novavax COVID-19 vaccine. People who had GBS after receiving Janssen vaccine should receive an age-		
appropriate bivalent Pfizer, or Moderna COVID-19 or monovalent Novavax vaccine at least 8 weeks after the Janssen dose.	<u>L</u>	
Note: At the time of initial vaccination, depending on vaccine product, children ages 6mo-4 years are recommended to receive 2 or 3 bivalent children aged 5 years are recommended to receive 1 or 2 bivalent doses. People ages 6 years and older who are unvaccinated or previously remonovalent vaccine doses are recommended to receive 1 bivalent dose. People 65 years and older may receive 1 additional bivalent dose. People of the dose are recommended to receive and older who received Janssen COVID-19 Vaccine and have not received a bivalent booster dose are recommended to receive an age-approance. Note: At the time of initial vaccination, depending on vaccine 2 or 3 bivalent ages 6 mo-4 years are recommended to receive 2 or 3 bivalent ages 6 mo-4 years are recommended to receive 2 or 3 bivalent ages 6 mo-4 years are recommended to receive 2 or 3 bivalent ages 6 years and older who are unvaccinated or previously remonovalent booster additional bivalent dose. People 65 years and older may receive 1 additional bivalent dose. People 65 years and older may receive 1 additional bivalent dose. People 65 years and older may receive 1 additional bivalent dose. People 65 years and older may receive 1 additional bivalent dose. People 65 years and older may receive 1 additional bivalent dose. People 65 years and older may receive 1 additional bivalent dose.	eceived only ersons 18 year	ars
2. RELEASE AND ASSIGNMENT: Please read the section on the reverse side of this form. The Providers Privacy Notice is available at the clinic site or accompanies this form. Then sign in the box at right. My signature below indicates I have read, understand, and agr 2. Release and Assignment of the COVID-19 Immunization Form and Vaccine Recipient Emergency Use of Authorization Fact Sheet.	n Consent	i
Please sign here.		İ
Date		

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	ENT INFORMATION) First Name:			MI: Last Na	ame:				
Date of	f Birth: /	/	Gender:	☐ Male ☐ Female P	Phone #:				
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☐ Me	dicare Number:		 						
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REQUIRED POLICY HOLDER INFORMATION:									
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COVII Co-adn admini Monke	D-19 VACCINE ADM ninistration of COVID- istered without regard ypox Poxvirus CDC	INISTRATION (Comp. 19 vaccines and other vactorial to timing (same visit) was Refer Interim Clinical Control of VID-19 vaccines to clarify	leted by secones inclicate the exconsideration	taff only) ADH Immuruding flu vaccine. COV ception of JYNNEOS vacants for Use of COVID-	nization Section @ /ID-19 vaccines and accine. Refer to JY 19 Vaccines CDC	501-537-8969. If other vaccines many NNEOS Vaccine Refer to the Pre-	·		
Pfizer 0 Pfizer 0 Pfizer 0		Maroon Cap) Gray Bivalent Label) p/Orange Bivalent Label)	☐ Moder ☐ Moder ☐ Moder ☐ Moder	COVID-19 Vaccine: Labe na 0.2mL Bivalent 6mo-5 na 0.5mL ≥ 12yrs (Gray B na 0.25mL 6-11yrs (Gray na 0.25mL 6mo-5yrs (Gray	yrs (Yellow Label) ivalent Label) Bivalent Label) y Bivalent Label)	Refrigerated COV Vaccine Novavax-Matri Other COVID-	ix-M1 19 Vaccine		
Route	Site Code	Dosage mL	MFG Code	Lot Number	≥ 6 mo4 or	≥6 years Bivalent Dose #	Novavax Dose #		
			Code		5years Bivalent Dose #	□One	One		
					□One □Two	□Two	□Two		
□IM					☐Three ☐ Four		☐ Booster Dose for ≥ 18 yrs only		

MFG Codes: PFR=Pfizer-BioNTech, MOD=Moderna, ASZ=AstraZeneca, NVX=Novavax, MSD=Merck Site Codes: Right Deltoid = RD, Left Deltoid = LD, Right Leg = RL, Left Leg = LL, Right Arm = RA, Left Arm = LA

Date Vaccine Administered: __/_/__/ FORM 4133 Revised 05/18/2023 Signature and Title of Vaccine Administrator: __