

COVID-19 VACCINE SCREENING AND CONSENT FORM

Name: Last:		PATIENT (PLEASE PRINT) First: Middle Initial:							
Date of Birth: Month	Day	Year Mobile Phone Number (Patient or Guardian): ()							
Address: Apt/Room #:									
City:			State:	Zip	<u> </u>				
Name of Legal Guardian	: Last:		First: Middle Initial:						
Sex (Gender assigned at birth) Female Male	Race ☐ America ☐ Asian	Ethnic nerican Indian or AlaskaNative					ispanic orLatino ot Hispanic orLatino		
Primary Insurance Carri	er ID#:								
Insurance Company:	-		Insu	rance Company	Phone #				
Insured's Name:		R	elationship:		Insured's Date	of Birth			
Secondary Insurance Ca	arrier ID #:		Grp #:						
Insurance Company:	nsurance Company:Insurance Company Phone # nsured's Name:Insured's Date of Birth					6D: ()			
Insured's Name:		K	elationship:		_Insured's Date	of Birth			
Designation of COVID-19	9 vaccination	dose number?	□First Dose □ Sec	ond Dose \square	Third Dose/Roo	ster Dose*			
Doorgination of Corns it	, raddination	- dood Hambol I		0110 0000 🗀	771114 2000/200	0101 0000			
ECTION 2: COVID-19 SCRE	ENING QUES	TIONS							
Please check YES or No fo							Yes	No	
1. Do you have today or have									
fatigue, muscle or body ac diarrhea?	hes, headache	, new loss of taste or	smell, sore throat, conges	tion or runny nose	e, nausea, vomitin	g, or			
2. Have you tested positive for	or and/or been	diagnosed with COVI	D-19 infection within the I	ast 10 days?					
3. Have you had a severe all					nis vaccine or to a	nv of			
the ingredients of this vacc		,gp				,			
4. Have you had any COVID-	-19 Antibody th	erapy within the last 9	90 days (e.g. Regeneron,	COVID Convaleso	cent Plasma, etc.)				
	0005511110	0111D 4 N 05 50D 00	VID 40 VA 00 NIE						
SECTION 3: IMMUNIZATION Please check YES or No fo			VID-19 VACCINE				Yes	No	
Please check YES or No for each question. 5. Do you carry an Epi-pen for emergency treatment of anaphylaxis and/or have allergies or reactions to any medications, foods,								140	
vaccines or latex?	or ciriorgonoy t	realment of anaphyla	Alo alla, or riave allergice c	or redections to drift	medications, ioo	uo,			
6. For women, are you pregr			ecome pregnant?						
7. For women, are you curre				-	-				
8. Are you immunocomprom				.0					
9. Do you have a bleeding di					202			-	
10. Are you a female age 1811. If you are under the age									
12. Have you received a pre									
12. Have you loodivou a pie	*1000 0000 01 C	ing COVID-10 Vaccing	5. II yoo, willon manufacti	aror o vaccino dia ;	, ou 10001VG.				
13. If this is your third dose or					r your second dos	se (booster)			
of Janssen (Johnson and John	,	•		•					
activetreat days have	mentfor cancer passed from t	, etc.), are at least 12 he completion of your	e.g. solid organ transplan years of age (Pfizer-BioN mRNA COVID-19 primar	Tech COVID-19 v yseries.	accine only) and a				
2) At least 6	months have p	assed since the comp	letion of an mRNA COVID	0-19 vaccine prima	ary series.				

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;	3) At least 2 months have passed since the initial dose of your Janssen (Johnson and Johnson) COVID-19 vaccination and you are 18 years of age or older.								
	t I am: (a)	the patient and at leas	t 18 years of age; (b) th		the patient and confirm that the				
					patient named above. Further, I	hereby give my conser	nt to the Florid	ak	
		(DOH) or its agents to			iconcod by FDA. This FDA cons	aval and license is for ve	ما المانية المانية ما	مام	
16 years of			e product that has been	iully approved and i	icensed by FDA. This FDA appr	oval and license is for us	e in individua	IIS	
•	•	•	er for usage in ages me	entioned above only) has not been approved or lice	nsed by FDA, but has b	een authoriz	ed	
	ency use by FDA, under an EUA to prevent Coronavirus Disease 2019 (COVID-19) for use in individuals either 5-15 years of age (Pfizer only) or 18 ge and older (Moderna and Johnson and Johnson) ; and the emergency use of this product is only authorized for the duration of the declaration that								
		ustifying the authorizati zation revoked sooner.	on of emergency use o	t the medical produ	ct under Section 564(b)(1) of the	ne FD&C Act unless the	eclaration	IS	
			all nossible side effects	or complications as	sociated with receiving vaccine	(s) Lunderstand the ris	ks and hene	fits	
associated	with the a	bove vaccine and have	received, read and/or ha	ad explained to me t	he Emergency Use Authorization individual that such questions were an	n Fact Sheet on the CC	VID-19 vacci		
		-		·	-	•		for	
observation	I acknowledge that I have been advised to remain near the vaccination location for approximately 15 minutes (or more in specific cases) after administration for observation. If I experience a severe reaction, I will call 9-1-1 or go to the nearest hospital.								
					armless the State of Florida, the sors, divisions, affiliates, subsidi				
and employ	ees from a	any and all liabilities or c			t of, in connection with, or in any				
the vaccine			/bfit	CHOTO Flavidada		O. I			
immunizati	age that: (a	a) i understand the purp ition in Florida SHOTS a	oses/benetits of Florida nd my nersonal immuni	i SHOTS, Florida ST zation information w	mmunization registry and (b) Do ill be shared with the Centers fo	JH WIII INCIUGE MY PERS Ir Disease Control (CDC	onai :) or other		
federal age			na my porochai mimam	zation information w	in bo ondrod with the contere to	i bioodoo control (obc	y or ourior		
		H, FDEM, or its agents	to submit a claim to my	insurance provider	or Medicare Part B without sup	plemental coverage pay	ment for me	for	
					nefits be made on my behalf to				
		ed items and services. I ervice, upon receipt of s		yment for which I ar	n financially responsible is due	at the time of service or	if DOH invoid	:es	
		t of the DOH Notice of F							
1 acknowled	age receip	tor the Dorn Notice of t	iivacy i ractices.						
Signature of P	atient or A	Authorized Representa	tive		Date:		=		
Duint Name of	Danuara	tativa and Dalatianahi	n ta Dawaan Daasiisina	Vassina					
Print Name of	Represen	tative and Relationshi	p to Person Receiving	vaccine:			_		
						Date of EUA Fact St	and the same		
Site	Route	Manufact	rer (MVX)	Lot #	Everiredian Data	Dale of EUA raci 3i	ieei		
(LD/RD)				Unit of Use/ Unit of Sale	Expiration Date				
	IM			om or saic					
Administer	red at la	ocation: facility							
name/ID		,							
Administered at location: Type									
Administra	ıtion Ad	ldress:							
7 (
CVX (prod	luct)								
Sending o	rganiza	tion:							
	<u> </u>								
Vaccinator Prir	nt Name:			Signature:		Date:			
Vaccina or mi Vaccine admi		providorauffix		Jigilalole.		baie			
vaccine aami	instering	providersumx:							

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