

COVID-19 VACCINE SCREENING AND CONSENT FORM

Adr	ministration	Facility Name/Fac	cility ID:						
SECTION 1: INFORMATION A	BOUT PATIFI	NT (PLEASE PRINT)							
Name: Last:	.50011711121	First:	Middle Initial:						
Date of Birth: Month	Day	Year	Mobile Phone Number (Patient or Guardian): ()						
Address:		Apt/Room #:							
City:	,	State: Zip:							
Name of Legal Guardian: Last:		First:		Middle Initial:					
Sex (Gender assigned at birth) ☐ Female ☐ Male	☐ Asian	n Indian or AlaskaNative African American	☐ Native Hawaiian or other☐ Pacific Islander☐ White	☐ Other Asian ☐ Unknown☐ Other Nonwhite☐ Other Pacific Islander	Ethnicity ☐ Hispanic or ☐ Not Hispani ☐ Unknown		no		
Primary Insurance Carrie	r ID #:		Grp #:		1				
Insurance Company:			Insu	rance Company Phone #					
Insured's Name:		R	elationship:	Insured's Date	e of Birth				
Secondary Insurance Car	rrier ID #:		Grp #:						
Insurance Company:	Insurance Company Phone #								
Insured's Name:	:Relationship:Insured's Date of Birth								
Designation of COVID-19	vaccination	dose number?	□ First Dose □ Sec	ond Dose ☐ Third Dose/Bo	nster Dose*				
Designation of COVID-13	Vaccination	dosc Hallisel :		0110 D030	00101 0000				
SECTION 2: COVID-19 SCREE	ENING QUEST	TIONS							
Please check YES or No for						Yes	No		
1. Do you have today or have you had at any time in the last 10 days a fever, chills, cough, shortness of breath, difficulty breathing,									
fatigue, muscle or body aches, headache, new loss of taste or smell, sore throat, congestion or runny nose, nausea, vomiting, or									
diarrhea? 2. Have you tested positive for and/or been diagnosed with COVID-19 infection within the last 10 days?									
Have you had a severe allergic reaction (e.g. needed epinephrine or hospital care) to a previous dose of this vaccine or to any of									
the ingredients of this vaccine?									
		erapy within the last 9	00 days (e.g. Regeneron,	COVID Convalescent Plasma, etc	c.)				
SECTION 3: IMMUNIZATION S			VID-19 VACCINE			V	NI.		
Please check YES or No for			via and/an have allemics a	u vanationa ta any mandinationa fa	- d-	Yes	No		
5. Do you carry an Epi-pen for emergency treatment of anaphylaxis and/or have allergies or reactions to any medications, foods, vaccines or latex?									
6. For women, are you pregna	ant or is there	a chance you could b	ecome pregnant?						
7. For women, are you currently breastfeeding?									
Are you immunocompromised or on a medication that affects your immune system?									
9. Do you have a bleeding disorder or are you on a blood thinner/blood-thinning medication?									
10. Are you a female age 18 to 49 years old receiving the Janssen (Johnson and Johnson) COVID-19 vaccine?									
11. If you are under the age of 18 are you and/or your guardian aware that you are only eligible to receive the Pfizer vaccine? 12. Have you received a previous dose of any COVID-19 vaccine? If yes, which manufacturer's vaccine did you receive:									
12. Have you received a prev	ious dose of a	ny COVID-19 vaccine	e? If yes, which manufactu	irer's vaccine did you receive:					
*13. If this is your third dose or	booster dose	of an mRNA (Pfizer-P	BioNTech or Moderna) CO	VID-19 vaccine or your second do	ose (booster)				
*13. If this is your third dose or booster dose of an mRNA (Pfizer-BioNTech or Moderna) COVID-19 vaccine or your second dose (booster) of Janssen (Johnson and Johnson) COVID-19 vaccine and you meet one or more of the following:									
Moderately to severely immunocompromised (e.g. solid organ transplant recipient, immunosuppressant medications,									
active treatment for cancer, etc.) and at least 28 days have passed from the completion of your mRNA COVID-19 primary									
series.			detion of on m-DNA COVIII	7. 40					
2) At least 6 n of the follow		assed since the comp	DIELION OF AN MRINA COVIL	D-19 vaccine primary series <u>and</u> y	rou meet one				

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- a. 65 years of age or older
- Reside in a long-term care facility
- Age 18-64 years of age with underlying medical condition(s) or
- Age 18-64 years of age with increased risk for COVID-19 exposure and transmission because of occupational
 or institutional setting
- 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.
- 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 12 years of age (for Pfizer vaccine consent only); or (c) legally authorized to consent for vaccination for the patient named above. Further, I hereby give my consent to the Florida Department of Health (DOH) or its agents to administer the COVID-19 vaccine.
- Currently, Pfizer is the only COVID-19 vaccine product that has been fully approved and licensed by FDA. This FDA approval and license is for use in individuals 16 years of age and older only. I understand that this product (other than Pfizer for usage in ages mentioned above only) 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 either 12-15 years of age (Pfizer only) or 18 years of age and older (Moderna and Johnson and Johnson); 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 COVID-19 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 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.
- On behalf of myself, my heirs and personal representatives, I hereby release and hold harmless the State of Florida, the Florida Department of Health (DOH),
 the Florida Division of Emergency Management (FDEM) and 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 Florida SHOTS, Florida's immunization registry and (b) DOH will include my personal
 immunization information in Florida SHOTS and my personal immunization information will be shared with the Centers for Disease Control (CDC) or other
 federal agencies.
- I further authorize DOH, FDEM, or its agents to submit a claim to my insurance provider or Medicare Part B without supplemental coverage payment for me for the above requested items and services. I assign and request payment of authorized benefits be made on my behalf to DOH, FDEM, or its agents with respect to the above requested items and services. I understand that any payment for which I am financially responsible is due at the time of service or if DOH invoices me after the time of service, upon receipt of such invoice.
- I acknowledge receipt of the DOH Notice of Privacy Practices.

Signature of Pa	atient or /	Authorized Representa	Date:									
Print Name of Representative and Relationship to Person Receiving Vaccine:												
Site (LD/RD)	Route	Manufacturer (MVX)		Lot # Unit of Use/ Unit of Sale	Expiration Date	Date of EUA Fact Sheet						
	IM											
Administered at location: facility name/ID												
Administered at location: Type												
Administration Address:												
CVX (product)												
Sending organization:												
Vaccinator Print Name:				Signature:		Date:						
Vaccine administering providersuffix:												

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