

## COVID-19 Vaccine Acknowledgement and Consent Form Recipient Information (Please Print Clearly)

**Section 1: Demographic Information** 

Last Name:		First Name:				
Date of Birth (MM/DD/YYYY)		Age (years):	Weight (pounds):			
Home Address:			Phone:			
City:		State:	Zip:			
Pediatrion based up Adult: For Booster waned oo Addition protective	Ferminology for COVID-19 Vaccine dosing:  Pediatrics: For the purposes of this initiative, pediatrics is defined as persons aged 0-17 years of age*, but there is a distinct dosing pased upon age.  *Of note, there are different dosages for pediatrics ages 5-11, and pediatrics ages 12+  Adult: For the purposes of this initiative, adults are defined as persons aged 18+.  Booster dose: a subsequent dose of vaccine administered to people in whom protection from primary vaccination is likely to have evaned over time.  Additional dose after an initial primary series: a subsequent dose of vaccine administered to people who likely did not mount a protective immune response after primary vaccination in order to optimize vaccine-induced protection. An additional mRNA COVID-19 vaccine dose is recommended for moderately and severely immunocompromised people who received an mRNA vaccine primary series.					
Section	2: COVID-19 Vaccination History					
1.	Which dose of the COVID-19 vaccine are you r  First Dose Second Dose Booster I  *If this is your First Dose, skip to Section 3:	se (Immunocompromised) Dose	on Screening below*			
2.	Please indicate the manufacturer(s) of your previous COVID-19 vaccination doses:					
	<u>First Dose</u> : □ Pfizer □ Moderna □ Janssen (	Johnson & Johnson)				
	Second Dose: ☐ Pfizer ☐ Moderna ☐ I have n	not received a Second Dose				
	Third Dose (immunocompromised): ☐ Pfizer ☐	□ Moderna				
3.	FOR IMMUNOCOMPROMISED PERSONS AC immunocompromised and seeking a third dose Pfizer-BioNtech or Moderna vaccines and has i second dose in your primary series?  ☐ Yes ☐ No ☐ N/A	, have you been fully vaccinat	ted with either the			
4.	FOR PERSONS AGED 18 YEARS AND OLDE scenario applies to you:  I have been fully vaccinated with the Pfizer-E than 6 months since I received the second dose my primary series.  I am at least 18 years of age or older and received the second dose my primary series.	BioNtech or Moderna vaccine, e (or third dose for immunoco ceived a single dose of Janss	and it has been more mpromised patients) in en (Johnson & Johnson)			

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## Section 3: Pre-Vaccination Screening

The following questions will help us determine whether you can receive the COVID-19 vaccine today. If you answer "yes" to any question, it does not necessarily mean you should not be vaccinated. It just means additional questions must be asked. If a question is not clear, please ask a staff member for further explanation:

		Yes	No	
1. How old	are you? □ 5-11 years □12 years or older □ Other			
2. Are you	feeling sick today?			
Have you ever had a severe allergic reaction* to any of the following:				
• A 0	component of the COVID-19 vaccine			
• Po	olysorbate			
• A p	previous dose of COVID-19 vaccine			
• An	nother vaccine (other than COVID-19 vaccine) or an injectable medication			
	omething other than a vaccine or injectable therapy such as food, pet, nom, environmental or oral medication allergies			
4. Check all	I that apply to you:			
• Ha	ave a history of myocarditis or pericarditis			
	ad COVID-19 and was treated with monoclonal antibodies or convalescent rum			
	agnosed with Multisystem Inflammatory Syndrome (MIS-C or MIS-A) after a DVID-19 infection			
• Ha	ave a bleeding disorder and/or take a blood thinner			
• Ha	ave a history of heparin-induced thrombocytopenia (HIT)			
• An	n currently pregnant or breastfeeding			
• His	story of Guillain-Barré Syndrome (GBS)			

\*An allergic reaction includes a severe allergic reaction (e.g., anaphylaxis) that required treatment with epinephrine or EpiPen® or that caused you to go to the hospital. It would also include an allergic reaction that occurred within 4 hours that caused hives, swelling, or respiratory distress, including wheezing.

I consent to administration of a COVID-19 vaccination and acknowledge and agree with the following statements:

- The U.S. Food and Drug Administration (FDA) has provided Emergency Use Authorization (EUA) of the Moderna and Janssen (Johnson & Johnson) COVID-19 vaccines. On August 23rd, 2021 the FDA gave full approval of the Pfizer-BioNTech COVID-19 vaccine to go from EUA to full FDA approval for ages 16 and above. Use of the Pfizer-BioNTech COVID-19 vaccine for patients aged 5-15 years remains available under EUA at this time.
- I have received the Fact Sheet for Recipients and Caregivers for the vaccine that I am receiving, and have read it or have it read to me.
- Some versions of the COVID-19 vaccine require additional doses to be effective. I understand that I will be informed at the time of vaccination whether I will need additional dose(s). If additional dose(s) are required, I understand that I am responsible for scheduling an appointment for future doses in accordance with the timeframe outlined in the Fact Sheet.
- I understand the known and potential risks and benefits to the COVID-19 vaccine and the extent to which such benefits and risks are unknown.
- I acknowledge that I have the option to refuse vaccination and have been informed of any available alternatives to the COVID-19 vaccine and the risks and benefits of available alternatives.
- Recipients who are Pregnant or Breastfeeding: Evidence about the safety and effectiveness of COVID-19 vaccination during
  pregnancy and while breastfeeding, although limited, has been growing. CDC recommends COVID-19 vaccination to persons

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who are pregnant, breastfeeding, trying to get pregnant now, or might become pregnant in the future. I have discussed the potential risks of COVID-19 infection versus the risk of vaccination with my healthcare provider and have made the informed decision to receive a COVID-19 vaccine.

- Recipients of additional doses: By receiving a third dose, you represent that you are immunocompromised.
- Recipients of booster doses: By receiving a booster dose, you represent that you meet one of the following qualifications: you are 18 years or older and received your mRNA (Moderna or Pfizer-BioNTech) COVID-19 primary vaccine series at least 6 months ago; or 18 years or older and received your primary vaccine series (single dose) with Janssen (Johnson & Johnson).
- I understand that it is recommended that I remain at the vaccination clinic for fifteen (15) minutes following administration of the vaccine for observation (the "Monitoring Period") to ensure I do not experience an adverse reaction. Recipients that have a history of severe allergic reactions should be monitored for thirty (30) minutes post vaccination.
- I acknowledge that I have received information on V-safe, a voluntary smartphone based tool operated by the Centers for
  Disease Control and Prevention (CDC). Through V-safe, vaccine recipients can report any side effects of the COVID-19
  vaccine to the CDC. This information helps CDC monitor the safety of COVID-19 vaccines in near real time.
- I authorize Ascension or its agents to submit a claim to my insurance provider for administration of the COVID-19 vaccine. I understand that I will have no out of pocket cost or cost sharing associated with receiving the vaccine. I acknowledge I was offered the Notice of Privacy Practices, which is also available at healthcare.ascension.org/NPP.
- I have had the opportunity to ask questions which have been answered to my satisfaction.

If you experience an adverse reaction to the COVID-19 vaccine, please contact your primary care provider or present to the nearest emergency department. If you are experiencing a medical emergency, call 911.

Signature of Recipient/Authori	Date:				
Print:					
If signed by Authorized Repres	sentative, please state relationshi	p to Recipient:			
	FOR CLINIC US	SE ONLY			
Vaccine Administrator (Signatu	ure):				
Vaccine Administrator (Print N	ame):				
Administration Date/Date Fact	Sheet Provided:				
Manufacturer	Lot Number	Expiration Date Sit	e of Administration		
☐ Monitoring Period completed and no adverse reaction noted. ☐ Recipient declined Monitoring Period. Waiver completed.					
Signature of Observer: Date/time:		_ Print Name:			
		ng Period Waiver (if applicable) up	loaded to PureOHS		
	on associates, contractors, or med	dical staff members only) or the na			

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