WDGPH COVID-19 Vaccine Backup Paper Record

All-In-One: Consent, Pre-Screening, Clinical Record (last updated 18Aug2021)

CONSENT

Last Name	First	Name		Identification (e.g., health card #)			
Gender: □ Female □ □ Prefer not to answer □	Primary Care Clinician (Family Physician or Nurse Practitioner)						
If Indigenous, please indice First Nations Métis (includes members Settlement) Inuk/ Inuit Other Indigenous, spect Prefer not to answer Unknown							
Home Phone	Mobile Pho	one					
Email Address							
Street Address			City	Province	Postal Code		
Date of Birth (month,day, year) Age			Has the client previously received one or more doses of a COVID-19 vaccine? If yes, please complete the information below for all doses of vaccine received. First Dose: • date:/ (month, day, year) • name: Second Dose: • date:/ (month, day, year) • name:				

Sources: Consent (MOH 17Aug2021);

Consent to Receive the Vaccine

I have read (or it has been read to me) and I understand the Immunization Prepackage, including the following documents: 'COVID-19 Vaccine Information Sheet' and 'What you need to know about your Covid-19 vaccine appointment'.

- I have had the opportunity to ask questions regarding the vaccine I am receiving and to have them answered to my satisfaction.

☐ I consent to receiving the vaccine, including all recommended doses in the series.

- I understand that I may withdraw this consent at any time.
- I understand that if I am withdrawing consent as a substitute decision maker of an individual, then I must contact the congregate setting that the individual resides in.

Note: Please contact the vaccination clinic where you are supposed to receive the Covid-19 vaccine if you change your mind and no longer consent to receiving the vaccine. This will allow someone else to take your spot. If consent has been withdrawn by a substitute decision maker of an individual who resides in a congregate setting, then the congregate setting must contact the local public health unit.

Acknowledgement of Collection, Use and Disclosure of Personal Health Information

The personal health information on this form is being collected for the purpose of providing care to you and creating an immunization record for you, and because it is necessary for the administration of Ontario's COVID-19 vaccination program. This information will be used and disclosed for these purposes, as well as other purposes authorized and required by law. For example,

- it will be disclosed to the Chief Medical Officer of Health and Ontario public health units where the disclosure is necessary for a purpose of the *Health Protection and Promotion Act*. And
- it may be disclosed, as part of your provincial electronic health record, to health care providers who are providing care to you.

The information will be stored in a health record system under the custody and control of the Ministry of Health.

Where a Clinic Site is administered by a hospital, the hospital will collect, use and disclose your information as an agent of the Ministry of Health.							
$\hfill I$ acknowledge that I have read and understand the above statement.							
You may be contacted by a hospital, local public health unit, or the Ministry of Health for purposes related to the COVID-19 vaccine (for example, to remind you of follow up appointments and to provide you with a record of immunization). If you consent to receiving these follow up communications by email, please indicate this using the box below.							
☐ I consent to receiving follow-up communications:							
☐ by email ☐ by text/SMS							
If selected by email, please provide your email address:							
Consent to Being Contacted About Research Studies							
You have the option of consenting to be contacted by researchers about participation in COVID-19 vaccine related research studies. If you consent to be contacted, your personal health information will be used to determine which studies may be relevant to you, and your name and contact information will be disclosed to researchers. Consenting to be contacted about research studies does not mean you have consented to participate in the research itself. Participating in research is voluntary. You may refuse to consent to be contacted about research studies without impacting your eligibility to receive the COVID-19 vaccine.							
If you do not wish to be contacte	ed about research studies, plea	ase indicate this below.					
If you consent to be contacted about research studies, and then change your mind, you may withdraw consent at any time by contacting the Ministry of Health at vaccine@ontario.ca .							
This will not impact your eligibility to receive the Covid-19 vaccine.							
I consent to be contacted about COVID-19 vaccine related research studies:							
□ by email □ by text/SMS □ by phone □ by mail							
If selected by email, please provide your email address:							
☐ I do not consent to be contacted about COVID-19 related research studies:							
Signature	Print Name	Date of Signature					

If signing for someone other than yourself, indicate your relationship to that other person:
$\hfill\Box$ If signing for someone other than myself, I confirm that I am the parent / legal guardian or substitute decision maker.
Specific Issues re: Long-Term Care Homes Act, 2007
The resident's consent to receive the vaccine may be withdrawn or revoked at any time.
Statement respecting section 83 of the Act:
Please note the following legal protection:
Every licensee of a long-term care home shall ensure that no person is told or led to believe that a prospective resident will be refused admission or that a resident will be discharged from the home because, a) a document has not been signed; b) an agreement has been voided; or c) a consent or directive with respect to treatment or care has been given, not given, withdrawn or revoked.
Notes

Pre-Screening

If the client is receiving the AstraZeneca/COVISHIELD or Janssen COVID-19 Vaccine:	If yes, please provide details
Have you experienced major venous and/or arterial thrombosis with thrombocytopenia following vaccination withany vaccine?	
□ No □ Yes	
Have you experienced a previous cerebral venous sinus thrombosis (CVST) with thrombocytopenia or a heparin- induced thrombocytopenia (HIT)?	
□ No □ Yes	If yes, please provide details
Have you experienced a pervious episode of capillary leak syndrome?	
□ No □ Yes	
Warning: The Astra Zeneca COVID-19 Vaccine/ COVISHIELD COVID-19 & Janssen COVID-19 Vaccines are contraindicated in individuals who have experienced:	
 A previous cerebral venous sinusthrombosis (CVST) with thrombocytopenia, and/or 	
Heparin-induced thrombocytopenia (HIT), and/or Size data of a city and leads through the second transfer of the second transfer	
 Episodes of capillary leak syndrome 	
Individuals who think they have experienced Heparin-induced thrombocytopenia (HIT) should not receive the vaccine.	
Has the COVID-19 Vaccine Information Sheet for individuals who received a first dose of Astra Zeneca COVID-19 Vaccine/COVID-19 COVISHIELD been reviewed with the client as part of the pre-assessment?	
□ No □ Yes	
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Ask all clients:

Have you been sick in the past few days? Do you have symptoms of COVID-19 or have a fever today?	If yes, please provide details		
□ No □ Yes			
Have you had a serious allergic reaction within 4 hours to the COVID-19 vaccine before?	If yes, please provide details		
□ No □ Yes			
Do you have allergies to polyethylene glycol, tromethamine(Moderna only) or polysorbate?	If yes, please provide details		
□ No □ Yes			
Have you had a serious allergic reaction to a vaccine or medication given by injection (e.g., IV, IM), needing medical care?	If yes, please provide details		
□ No □ Yes			
Have you received a vaccine inthe past 14 days?	If yes, please provide details		
□ No □ Yes			

Do you have a weakened immune system or are you taking anymedications that can weaken your immune system (e.g., high dose steroids, chemotherapy)?	If yes, please provide details		
□ No □ Yes			
If, are you receiving stem cell therapy, CAR-T therapy,yes chemotherapy, immune checkpoint inhibitors, monoclonal antibodies or other targeted agents?	If yes, please provide details		
□ No □ Yes			
If on one of the therapies listed, have you spoken with your treating health care provider about getting the vaccine?			
□ No □ Yes			
Do you have a bleeding disorder or are taking blood thinners? □ No □ Yes	If yes, please provide details		
Have you ever felt faint or fainted after receiving a vaccine or medical procedure?	If yes, please provide details		
□ No □ Yes			

Clinical Record

FOR CLINIC USE ONLY										
Agent	CO 19	VID-	Product Name			Lot # & Expiry				Dose Amount:
Anatom Site	ical	□ Le	Left deltoid □ Right deltoid		Route	Int	ramı	ıscular (IM)	Dose #:	
Date Giv	Date Given/		/ (m/d/	/ yyyy)	Time Given	: am pm	_	AEFI? (after receiving current dose)		□ Yes □ No
Given By (Name, Designation)				Location				Authorized By		
Reason for Immunization		Long Term Long Term Long Term Long Term Retiremen Retiremen Retiremen Retiremen ALC: Altern Hospitals Healthcare Indigenou Advanced	ing Term Care: Resident ag Term Care: Healthcare Wo ag Term Care: Other Employe ag Term Care: Other Non-Em ag Term Care: Essential Careg cirement Home: Healthcare W cirement Home: Resident cirement Home: Other Employ cirement Home: Other Employ cirement Home: Other Patis spitals althcare Worker igenous community conic Home Care wanced Age: Community Dwe con with Priority Health Cond				 □ Caregiver of Person with Priority Health Condition □ Congregate Living: Resident □ Congregate Living: Staff □ Congregate Living: Essential Caregiver □ Agriculture: Temporary Foreign Worker-Congregate Setting □ Community at Greater Risk □ Agriculture or Farm Worker (not temp foreign worker) □ Food Manufacturing Worker □ Education Worker □ Child Care Worker □ Child Care Worker □ Child Care Worker □ Age Eligible Population 			
Reason Immunization NotGiven			tion	munizati actitione actitione edically I	on is contrain r recommend r decision to t	dicated Is immunization emporarily def			TIENT consent	