WASHINGTON STATE COVID-19 POINT OF CARE TEST RESULT REPORT FORM Complete one form per result. Submit by fax to the Washington State Department of Health at (206) 512-2126.			
Submitter name: Submitted date (MN			
Section 1: Testing Facility and Ordering Provider Information	///////////////////////////////////////		
Facility name: License or CLIA num	her (if applicable):		
	ьет (п аррпсавіе).		
Facility address:	City:		
State: Zip code: County:	Phone:		
Type of facility: ☐ Airport/Transit station ☐ Hospital	□ Homeless shelter		
☐ Assisted Living/Adult Family Home ☐ Inpatient b	•		
☐ Childcare or daycare health care	□ K-12 School		
□ College/University □ Nursing Ho			
	care (including Other (specify):		
□ Correctional setting freestanding emergency			
☐ Drive-/walk-through testing site department,			
Ordering provider name (first and last): Phone:	NPI (if applicable):		
Ordering provider street address:			
Ordering provider city: Zip code:	County:		
Section 2: Patient Information			
Last name: First name:	Middle name:		
Sex at birth: □ Female □ Neither/Other Is the patient: □ Pregnant	□ Postpartum □ Unknown		
☐ Male ☐ Unknown ☐ Neither pregnant nor postpartum			
What is the patient's affiliation to the facility? Date of birth (MM/DD/YYYY):			
□ Resident □ Staff □ Visitor □ Patient □ Student □ Client □Inmate □ / /			
Age: years Did the patient die? Yes No Date of death (MM/DD/YYYY): //			
Patient's address:	City:		
State: Zip code: County:	Phone:		
Race (select all that Unknown Asian American Indian or Alaska Native Asian			
apply):			
□ Other race (specify):			
Ethnicity: Hispanic or Latino Did the patient have symptoms at time of testing?			
□ Not Hispanic or Latino □ Unknown □ Yes □ No □ Unknown Patient identifier (if applicable): □ □ N/A			
☐ Medical Record Number ☐ Patient Internal ID ☐ Public Health Case ID			
	r (specify):		
Section 3: Test Information	. (65-66.77)		
	OW COVID-19		
☐ Access Bio CareStart COVID-19 Antigen Test ☐ BD Veritor System for Rapid Detection of SARS-CoV-2			
☐ BioFire Diagnostics Respiratory Panel 2.1-EZ ☐ Cepheid Xpert Xpress SARS-CoV-2 test			
☐ Cue Health Cue COVID-19 Test ☐ Luminostics Clip COVID Rapid Antigen Test			
☐ LumiraDx SARS-CoV-2 Ag Test ☐ Roche cobas SARS-CoV-2 & Influenza A/B Nucleic Acid			
☐ Quidel Sofia 2 Flu + SARS Antigen FIA Test for use on the cobas Liat System			
☐ Quidel Sofia SARS Antigen FIA ☐ Other (specify):			
Specimen type: Test result:	Specimen collection date		
□ Nasal swab □ Detected/Positive	(MM/DD/YYYY):		
□ NP (nasopharyngeal swab) □ Not detected/Negative	/		
□ Other (specify): □ Inconclusive/Undetermined/Invalid/Equivocal Device identifier: Specimen ID:			

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POC Report Form Field Descriptions

A description for each field in the Report Form is provided below. These explanations are intended to help you fill out the form completely. Please read them before contacting doh-surv@doh.wa.gov with questions on how to fill out the Report Form.

WASHINGTON STATE COVID-19 POINT OF CARE TEST RESULT REPORT FORM

Submitter name	The name of the person filling out the form	
Submitted date	The date this form was sent to the Washington State Department of	
	Health	
Section 1: Testing Facility and Ordering Provider Information		
Facility name	The facility's name	
License number or CLIA number (if	The facility's state license number or CLIA number. If the facility doesn't	
applicable)	have either number, put "N/A".	
Facility address (including city, state,	The facility's physical address. Use only five-digit zip codes.	
and zip code)		
County	The county where the facility is located	
Phone	The facility's phone number that DOH can call if there are questions	
	about results. Use 10-digit phone numbers.	
Type of facility	Check only one. Check the best option that describes the facility. If the	
	facility type isn't listed, check "Other" and provide additional details.	
Ordering provider name (first and last)	For health care providers or facilities, the full name of the medical	
	provider who ordered the POC test. Other facilities can put "N/A".	
Phone	The ordering provider's phone number. Use 10-digit phone numbers. If	
	there is not an ordering provider, put "N/A".	
NPI (if applicable)	The order provider's or health care facility's National Provider Identifier	
	(NPI). If there is not an NPI, put "N/A".	
Ordering provider street address	The ordering provider's physical address where they work. Use only five-	
(includes city and zip code)	digit zip codes. If there is not an ordering provider, put "N/A".	
Section 2: Patient Information		
Last name, First name, and Middle name	Provide the full name of the patient	
Sex at birth	Check the option that best describes the patient	
Is the patient pregnant?	Check the option that best describes the patient	
What is the patient's affiliation to the	How the patient is related to the facility where he or she was tested	
facility?		
Date of birth	The patient's date of birth	
Age	The patient's age in years at time of testing. If the patient is a child under	
	1 year of age, enter 0.	
Patient's address (includes city, state,	The patient's physical address. Use only five-digit zip codes.	
and zip code)		
County	The county where the patient lives	
Phone	The best phone number to reach the patient. Use 10-digit phone	
	numbers; if area code is unknown, enter 999 (example: (999) 555-1234).	
Did the patient die?	Check the option that best describes the patient	
Date of death	If the patient died, indicate the date the patient died	
Race	Check the option(s) with which the patient identifies	
Ethnicity	Check only one. Check the option with which the patient identifies	
Did the patient have symptoms at the	Indicate if the patient had symptoms of COVID-19 disease. This includes	
time of testing?	cough, shortness of breath or difficulty breathing, fever, chills, muscle	

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	pain, sore throat, and new loss of taste or smell. Other less common
	symptoms include nausea, vomiting, or diarrhea.
Patient identifier	Check only one. If your facility uses or assigns identifiers to patients,
	check the option used and provider the identifier of the patient. If your
	facility does not use or assign identifies, check "N/A".
Section 3: Test Information	
Test name	Check only one. Indicate the brand and name of the test the facility used
	to test this patient.
Specimen type	Check only one. Indicate the type of specimen used for this test. A nasal
	swab specimen is obtained by inserting an absorbent tip into both
	nostrils, just around the inside of the nostrils (also referred to as "nares").
	A NP (nasopharyngeal swab) specimen is obtained from "deep" in the
	nose. If the specimen type isn't listed, check "Other" and provide
	additional details.
Test result	Check only one. Indicate the option that identifies the patient's test
	result.
Specimen collection date	The date the patient's specimen was collected and tested
Device identifier (DI)	The DI for some tests can be found in the National Institute of Health's
,	Access GUDID Database. The Device Model is also acceptable here, or the
	full human readable form of the barcode. If the DI is unknown, put
	"Unknown."
Specimen ID	If the facility uses or assigns unique identifiers to specimens, provide that
-	ID. Many facilities using POC testing may not use specimen IDs because
	specimens are not stored. In that case, put "N/A".

