WASHINGTON STATE BOARD OF HEALTH

U.S. Department of Health and Human Services COVID-19 Laboratory Data Reporting Guidance Comparison

COVID-19 Pandemic Response,	COVID-19 Pandemic Response,
Laboratory Data Reporting: CARES Act	Laboratory Data Reporting: CARES Act
Section 18115	Section 18115
(January 8, 2021)	(<u>March 8, 2022</u>)
Entities Required to Report	
All laboratories – including laboratories,	Specifies reporting requirements based on
testing locations operating as temporary	entity:
overflow or remote locations for a laboratory,	- Facilities licensed under CLIA to
and other facilities or locations performing testing at point of care or with at-home	perform moderate- or high-complexity tests conducting Nucleic Acid
specimen collection related to SARS-CoV-2	Amplification Test (NAAT) testing
specimen conection related to OANO-COV-Z	- Entities conducting all other SARS-
	CoV-2 testing, except antibody and
	self-administered testing
Reporting Test Results	
All test results must be reported	For facilities licensed under CLIA to perform
	moderate- or high-complexity tests: all
	positive, negative, and inconclusive results
	from NAAT testing
	For all other entities (except antibody and
	self-administered tests): positive test results
Reporting Timeframe	
Within 24 hours of results being known or	Within 24 hours of results being known or
determined, on a daily basis to the appropriate state or local public health	determined, on a daily basis to the appropriate state or local public health
department based on the individual's	department based on the individual's
residence	residence
Reportable Data Components	
Patient name (Last name, First name, Middle	Patient name (Last name, First name, Middle
Initial) (optional reporting)	Initial)
Patient street address (optional reporting)	Patient street address
Patient residence zip code	Patient residence zip code
Patient residence county	Patient residence county
Patient phone number with area code	Patient phone number with area code
(optional reporting)	
Patient date of birth (optional reporting)	Patient date of birth
Patient age	Patient age
Patient race	Patient race
Patient ethnicity	Patient ethnicity
Patient sex	Patient sex

Ordering provider name and NPI (as	Ordering organization or ordering provider
applicable), zip code	name and NPI (as applicable), address,
Ordering provider address and phone	phone number, zip code along with affiliated
number (optional reporting)	organization (specific facility)
Date test ordered (date format)	Data component not included
Test ordered – use harmonized LOINC codes	a) Test ordered and b) test resulted- use
provided by CDC	appropriate LOINC codes, as defined by the
	Laboratory In Vitro Diagnostics (LIVD) Test
	Code Mapping for SARS-CoV-2 Tests
	provided by CDC
Test result – use appropriate LOINC and	Test result (values) – use appropriate
SNOMED codes, as defined by the	SNOMED-CT codes, as defined by the
Laboratory In Vitro Diagnostics (LIVD) Test	Laboratory In Vitro Diagnostics (LIVD) Test
Code Mapping for SARS-CoV-2 Tests	Code Mapping for SARS-CoV-2 Tests
provided by CDC	provided by CDC
Test Result date (date format)	Test result date (date format)
Device Identifier	Device Identifier
Accession # / Specimen ID	Accession # / Specimen ID
Performing facility name and/or CLIA	Performing facility name and CLIA number,
number, if known; performing facility zip code	address, phone number, zip code
Specimen Source - use appropriate LOINC,	Specimen Source - use appropriate LOINC,
SNOMED-CT, or SPM4 codes, or	SNOMED-CT, or SPM4 codes, or
equivalently detailed alternative codes	equivalently detailed alternative codes
Date specimen collected (date format)	Date specimen collected (date format)
Data component not included	Reporting entity name and CLIA number (or
	appropriate ID), and address
Answers to ask-on-order entry questions:	Data component not included
- First test?	
 Employed in healthcare? 	
 Symptomatic as defined by CDC? 	
- Hospitalized?	
- ICU?	
 Resident in a congregate care 	
setting?	
- Pregnant?	