Michigan Interim COVID-19 Person Under Investigation (PUI) Case Report Form

As the COVID-19 situation in the State of Michigan evolves, MDHHS continues to adapt resource and capacity planning to support the varied needs of our partners in healthcare and local public health organizations. MDHHS recently ordered that all health professionals should conduct testing for the Novel Coronavirus in accordance with the COVID-19 prioritization criteria published by MDHHS.

1. <u>Expansion of COVID-19 Testing Prioritization Criteria to Include All Critical Infrastructure</u> Workers with Potential Exposure, Whether Symptomatic or Asymptomatic

Given the continued expansion of COVID-19 testing capacity in Michigan, MDHHS is expanding the COVID-19 testing prioritization criteria to broaden the populations eligible for testing to include individuals with mild symptoms in certain circumstances. Specifically, health care providers should test any healthcare facility worker or first responder (even if they do not have symptoms). Critical infrastructure workers (i.e., any worker still leaving the home for in-person work) with potential COVID-19 exposure, whether symptomatic or asymptomatic, should be tested as well, so long as adequate specimen collection and test processing capacity remains after serving all known patients in higher-priority testing categories.

The U.S. Centers for Disease Control and Prevention (CDC) have issued clinical guidance to help prioritize COVID-19 testing resources that, unfortunately, remain too scare nationwide. These guidelines (https://www.cdc.gov/coronavirus/2019-ncov/hcp/clinical-criteria.html) group patients into Priority One, Priority Two, and Priority Three categories, reflecting risk of severe illness and other considerations like integrity of the healthcare system.

Providers must continue to follow MDHHS prioritization criteria and must prioritize test capacity for populations from Priority One, Priority Two, and Priority Three patients; these priorities are reproduced at the end of this memorandum. If capacity remains after serving patients from those priority populations, providers should test all critical infrastructure workers with potential COVID-19 exposure, whether symptomatic or asymptomatic. This population is not prioritized by the CDC guidelines.

MDHHS recognizes that population health needs, patient characteristics, and testing capacity vary significantly across the state, and this system seeks to broaden eligibility for testing to fully take advantage of available test capacity in the state, while still ensuring that the highest-risk patients can access testing resources. It is also important to note that Michigan is seeing alarming racial disparities in COVID-19 cases and deaths, with African Americans consisting of 14% of the state's population, but 33% of cases and 40% of deaths. Clinicians should be mindful of this disparity and have heightened awareness when considering testing and treatment strategies in this patient population.

Health care providers should assess available testing resources on a periodic basis (e.g., weekly) and determine if resources are sufficient to serve asymptomatic critical infrastructure workers, alongside other priority populations. Please note that testing asymptomatic critical infrastructure workers does not change precautions that should be taken to decrease COVID-19 spread, including quarantine of exposed individuals, and that a negative test result reflects infection status at a point in time. An individual may receive a negative test result soon after exposure but later develop an infection.

As a reminder, per the March 24, 2020 MDHHS Emergency Order, all CLIA-certified laboratories in Michigan are required to comply with prioritization criteria as promulgated by MDHHS. This includes Public Health, commercial, and healthcare facility laboratories. We believe that these clarifications and this expansion of prioritization criteria will help to improve access to COVID-19 testing.

This expanded prioritization criteria will take effect at 8:00 AM on April 21, 2020.

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2. Full COVID-19 Testing Prioritization Criteria Currently in Effect

As a reminder, the current MDHHS COVID-19 testing prioritization criteria are as follows:

Priority One

- Hospitalized Patients
- Healthcare facility workers with symptoms

Note: MDHHS interprets this to include all workers within a healthcare facility, not just providers of direct healthcare services.

Priority Two

- Patients in long-term care facilities with symptoms
 - Note: MDHHS interprets this to include any resident with symptoms in congregate living arrangements, not only long-term care facilities
- Patients over age 65 years with symptoms
- Patients with underlying conditions with symptoms
- First responders with symptoms

Priority Three

- Critical infrastructure workers with symptoms
- Individuals who do not meet any of the above categories with symptoms
- (Newly added) Asymptomatic health care workers and first responders
- Individuals with mild symptoms in communities experiencing high COVID-19 hospitalizations
 - o Note: MDHHS interprets the full state of Michigan to be a community with high COVID-19 hospitalizations

Newly Added Priority

- Critical infrastructure workers, including asymptomatic workers
 - O Note: these individuals may be tested only if specimen collection and testing capacity remains after serving all patient groups above

To streamline access to testing, MDHHS does not require healthcare providers to seek prior approval from MDHHS or submit a Person Under Investigation form when ordering COVID-19 lab testing that does not leverage MDHHS Bureau of Laboratories (BOL) testing. However, a medical provider must still order COVID-19 testing in line with the MDHHS COVID-19 Specimen Collection and Testing Prioritization Criteria for any test requisition submitted to any laboratory. If a COVID-19 test result is positive, the findings must be reported to the Michigan Disease Surveillance System to facilitate public health investigation. Medical providers, facilities, or laboratories must still obtain a Person Under Investigation number if submitting a specimen for testing to the MDHHS Bureau of Laboratories. To obtain testing at the MDHHS BOL, MDHHS will continue operation of the Mi-CLERN healthcare provider hotline, used for PUI identification number issuance. The Mi-CLERN hotline will operate twelve hours per day, from 8:00 AM to 8:00 PM, seven (7) days per week.

Please remember that, when submitting specimens to BOL, healthcare providers must include the PUI identification number on all of the following:

- The PUI Case Report Form. This form must be submitted to the patient's local health department when leveraging BOL testing facilities.
- All BOL laboratory testing requisition documents
- The specimen container

BOL will not prioritize specimens that arrive without a corresponding PUI identifier.

MDHHS is making these changes as part of its efforts to increase testing access to as many Michiganders as feasible while ensuring that statewide testing capacity is sustained. We will continue to monitor test availability and adjust this protocol, as necessary. For the latest information on Michigan's response to COVID-19, please visit www.michigan.gov/coronavirus. You may also email our Community Health Emergency Coordination Center at: checcdeptcoor@michigan.gov. Thank you for all you do to serve the residents of Michigan at this difficult time.

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Michigan Interim COVID-19 Person Under Investigation (PUI) Case Report Form Patient Information:

First name	e:				Last name:			
Date of bi	irth:	/	/	Age:	Sex:	Female	Male	
Parent/Gu	uardian nan	ne(s) if	patient is a	a minor:		/		
Patient re	sidence str	eet add	ress:			City:		
	County: _			S	tate:	Zip C	Code:	
Patient ph	none numbe	er(s):						
ls the pati	•		•	Asymptomatic		tient deceased		No
Submitting	g Facility I	nformat	tion:					
Reporting	healthcare	facility	:					
					le:			
	Health	care fac	cility conta	ct phone number:				
Reason fo	or Testing:							
	res optimal rity of the U.				ents, lessen the risk	of healthcare-as	ssociated infect	ions, and maintain the
	Hospitaliz	ed Patie	ent					
	Healthcare healthcare			ith symptoms (inc	ludes all workers wi	thin a healthca	re facility, not j	ust providers of direct
2.) Ensur	res those at	highest	risk of co	mplication of infect	ion are rapidly ident	ified and appro	priately triaged:	
					congregate living a sing facilities, etc.) <u>w</u>		., dormitories, ja	ails/prisons, camps,
	Patient ov	er 65 ye	ears of age	with symptoms				
	Patient wi	th unde	lying cond	litions <u>with sympto</u>	<u>oms</u>			
	First resp	onder <u>w</u>	ith sympto	<u>ms</u>				
3.) Ensur	res the heal	th of es	sential woı	kers in Michigan C	Communities:			
•	Critical Inf	rastruct	ure workei	with symptoms				
	Any indivi	dual <u>wit</u>	h mild syn	nptoms consistent v	with COVID-19			
	Asymptor	<u>natic</u> he	alth care w	orkers or first resp	oonders			
4.) Addit		<u>matic</u> cr	itical infras	structure workers. ter serving all patio		uals may be te	sted only if spe	cimen collection and
<u>Specimen</u>	Being Sub	mitted t	to:	Spe	cimen collection d	ate:/		_ (mm/dd/yyyy)
					(Rovider hotline at: (8		ned by case en	try into MDSS by
	Clinical or	Comm	ercial lab.	PUI (nCoV) ID is	not required			

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	CD	OC 2019-nCoV ID:	Form A	approved: OMB: 0920-1011 Exp. 4/23/2020
	PATIENT IDE	NTIFIER INFORMATIO	ON IS NOT TRANSMITTED TO CDC	
Patient first	name	Patient last name	Date of birth (MN	M/DD/YYYY):/
	PATIENT IDE	NTIFIER INFORMATIO	N IS NOT TRANSMITTED TO CDC	
GENTICES TO THE STATE OF THE ST	Human I	nfection with	2019 Novel Coronavi n (PUI) and Case Repo	rus
Reporting jurisdiction: Reporting health depart Contact ID a:		CD:	se state/local ID: C 2019-nCoV ID: IDSS loc. rec. ID/Case ID ^b :	
		r source case-patient. Assign Contact porters, use GenV2 or NETSS patient i	ID using CDC 2019-nCoV ID and sequential contact ID, e.g identifier.	ر., Confirmed case CA102034567 has contacts
Interviewer i	nformation			
Name of interviewer: L	ast	First		
Affiliation/Organization	1:	Telephor	ne Email	
Basic informa	ation			
What is the current status Patient under invest Laboratory-confirme Report date of PUI to CDC	of this person? igation (PUI) ed case	Ethnicity: Hispanic/Latino Non-Hispanic/ Latino Not specified	Date of first positive specimen collection (MM/DD/YYYY)://	Was the patient hospitalized? Yes No Unknown If yes, admission date 1 //(MM/DD/YYYY)
Report date of case to CD	C (MM/DD/YYYY):	Sex: Male Female	Did the patient develop pneumonia? Yes Unknown No	If yes, discharge date 1/ (MM/DD/YYYY) Was the patient admitted to an
County of residence:		Unknown Other	Did the patient have acute respiratory distress syndrome? Yes Unknown	intensive care unit (ICU)? Yes No Unknown
Race (check all that apply Asian Black White	American Ind	lian/Alaska Native iian/Other Pacific Islander	No Did the patient have another diagnosis/etiology for their illness? Yes Unknown	Did the patient receive mechanical ventilation (MV)/intubation? Yes No Unknown If yes, total days with MV (days)
Other, specify:	YY):/		No Did the patient have an abnormal chest X-ray? Yes Unknown No	Did the patient receive ECMO? Yes No Unknown Did the patient die as a result of this illness?
Symptoms present during course of illness: Symptomatic Asymptomatic Unknown	If symptomatic, onset date (MM/DD/YYYY): /	If symptomatic, date of s /	symptom resolution (MM/DD/YYYY): Unknown symptom status , unknown date	Pes No Unknown Date of death (MM/DD/YYYY): /
Is the patient a health car Does the patient have a h	ess onset, did the patient ha C la China A US country la vith another lab Ar	re facility (as a patient, work	osures (check all that apply): other Exposure to a cluster of e-patient respiratory distress of un another Other, specify:	Unknown patients with severe acute lower nknown etiology
If the patient had contact	with another COVID-19 case he PUI or case first identifie		See? Yes, nCoV ID of source case: Clinical evaluation leading to PUI determon of travelers: if checked, DGMOID	No Unknown N/A

Symptoms, clinical course, past medical history and social history

Collected from (check all that apply): Patient interview Medical record review

Unknown

Other, specify:

Public reporting burden of this collection of information is estimated to average 30 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information including suggestions for reducing this burden to CDC/ATSDR Reports Clearance Officer; 1600 Clifton Road NE, MS D-74 Atlanta, Georgia 30333; ATTN: PRA (0920-1011).



CDC 2019-nCoV ID:

Human Infection with 2019 Novel Coronavirus Person Under Investigation (PUI) and Case Report Form

During this illness, did the patient experience any of the following symptoms?	Sympton	n Present	?	
Fever >100.4F (38C) ^c	Yes	No	Unk	
Subjective fever (felt feverish)	Yes	No	Unk	
Chills	Yes	No	Unk	
Muscle aches (myalgia)	Yes	No	Unk	
Runny nose (rhinorrhea)	Yes	No	Unk	
Sore throat	Yes	No	Unk	
Cough (new onset or worsening of chronic cough)	Yes	No	Unk	
Shortness of breath (dyspnea)	Yes	No	Unk	
Nausea or vomiting	Yes	No	Unk	
Headache	Yes	No	Unk	
Abdominal pain	Yes	No	Unk	
Diarrhea (≥3 loose/looser than normal stools/24hr period)	Yes	No	Unk	
Other, specify:		•		•

Pre-existing medical conditions?					Yes	No	Unknown
Chronic Lung Disease (asthma/emphysema/COPD)	Yes	No	Unknown				
Diabetes Mellitus	Yes	No	Unknown				
Cardiovascular disease	Yes	No	Unknown				
Chronic Renal disease	Yes	No	Unknown				
Chronic Liver disease	Yes	No	Unknown				
Immunocompromised Condition	Yes	No	Unknown				
Neurologic/neurodevelopmental	Yes	No	Unknown	(If YES, specify)			
Other chronic diseases	Yes	No	Unknown	(If YES, specify)			
If female, currently pregnant	Yes	No	Unknown				
Current smoker	Yes	No	Unknown				
Former smoker	Yes	No	Unknown			•	_

Respiratory Diagnostic Testing

Test	Pos	Neg	Pend.	Not done				
Influenza rapid Ag A B								
Influenza PCR A B								
RSV								
H. metapneumovirus								
Parainfluenza (1-4)								
Adenovirus								
Rhinovirus/enterovirus								
Coronavirus (OC43, 229E, HKU1, NL63)								
M. pneumoniae								
C. pneumoniae								
Other, Specify:	•							

Specimens for COVID-19 Testing

Specimen	Specimen	Date	Sent to	State Lab
Type	ID	Collected	CDC	Tested
NP Swab				
OP Swab				
Sputum				
Other,				
Specify:				