Michigan Interim 2019 Novel Coronavirus (COVID-19) Person Under Investigation (PUI)/Case Report Form Cover Sheet

Please use the attached Human Infection with 2019 Novel Coronavirus Person Under Investigation (PUI) and Case Report Form. If you are a healthcare provider with a suspect COVID-19 PUI, contact your local health department (LHD). The LHD can assist in determining whether the individual meets the current CDC PUI criteria for testing (see below). If you are unable to reach an LHD contact, please call the Michigan Department of Health and Human Services (MDHHS) Communicable Disease (CD) Division at the numbers below.

For patients meeting the CDC PUI criteria for testing, the MDHHS CD Division will assign an nCoV ID number. An nCoV ID number **MUST** be assigned prior to submitting specimens to the MDHHS Bureau of Laboratories (BOL). The MDHHS CD Division can be contacted at: **(517) 335-8165** during business hours, or at: (517) 335-9030 after-hours and on holidays.

Evaluating and Reporting Persons Under Investigation (PUI): Local/State health departments, in consultation with clinicians, should determine whether a patient is a PUI for COVID-2019. The CDC clinical criteria for COVID-19 PUIs have been developed based on available information about this novel virus, as well as what is known about Severe Acute Respiratory Syndrome (SARS) and Middle East Respiratory Syndrome (MERS). These criteria are subject to change as additional information becomes available.

For more information, please see: https://www.cdc.gov/coronavirus/2019-ncov/hcp/clinical-criteria.html

Clinical features and epidemiologic risk				
Clinical Features	&	Epidemiologic Risk		
Fever or signs/symptoms of lower respiratory illness (e.g. cough or shortness of breath)	AND	Any person, including health care workers ² , who has had close contact ³ with a laboratory-confirmed ⁴ COVID-19 patient within 14 days of symptom onset		
Fever ¹ and signs/symptoms of a lower respiratory illness (e.g., cough or shortness of breath) requiring hospitalization		A history of travel from affected geographic areas ⁵ (see below) within 14 days of symptom onset		
Fever ¹ with severe acute lower respiratory illness (e.g., pneumonia, ARDS) requiring hospitalization ⁴ and without alternative explanatory diagnosis (e.g., influenza) ⁶	AND	No source of exposure has been identified		

The criteria are intended to serve as guidance for evaluation. In consultation with public health departments, patients should be evaluated on a case-by-case basis to determine the need for testing. Testing may be considered for deceased persons who would otherwise meet the PUI criteria.

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Footnotes:

¹Fever may be subjective or confirmed.

²For healthcare personnel, testing may be considered if there has been exposure to a person with suspected COVID-19 without laboratory confirmation.

³Close contact is defined as—a) being within approximately 6 feet (2 meters) of a COVID-19 case for a prolonged period of time; close contact can occur while caring for, living with, visiting, or sharing a healthcare waiting area or room with a COVID-19 case—or—b) having direct contact with infectious secretions of a COVID-19 case (e.g., being coughed on). If such contact occurs while not wearing recommended personal protective equipment or PPE (e.g., gowns, gloves, NIOSH-certified disposable N95 respirator, eye protection), criteria for PUI consideration are met. See CDC's updated <u>Interim Infection Prevention and Control Recommendations for Patients with Confirmed COVID-19 or Persons Under Investigation for COVID-19 in Healthcare Settings</u>. Data to inform the definition of close contact are limited. Considerations when assessing close contact include the duration of exposure (e.g., longer exposure time likely increases exposure risk) and the clinical symptoms of the person with COVID-19 (e.g., coughing likely increases exposure risk as does exposure to a severely ill patient). Special consideration should be given to healthcare personnel exposed in healthcare settings as described in CDC's <u>Interim U.S. Guidance for Risk Assessment and Public Health Management of Healthcare Personnel with Potential Exposure in a Healthcare Setting to Patients with COVID-19.</u>

⁴Documentation of laboratory-confirmation of COVID-19 may not be possible for travelers or persons caring for COVID-19 patients in other countries.

⁵Affected areas are defined as geographic regions where sustained community transmission has been identified. Relevant affected areas will be defined as a country with <u>at least</u> a CDC Level 2 Travel Health Notice. See all <u>COVID-19 Travel Health Notices</u>.

⁶ Category includes single or clusters of patients with severe acute lower respiratory illness (e.g., pneumonia, ARDS) of unknown etiology in which COVID-19 is being considered.

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After approval, the completed PUI/Case Report form with intact cover sheet (with patient identifiers below) should be faxed to the MDHHS Communicable Disease Division at (517) 335-8263 or uploaded to the Michigan Disease Surveillance System (MDSS) if the patient has been entered into the MDSS.

Patient Information:
irst name: Last name:
CoV ID#: MI-
Note: An nCoV ID # must be obtained from MDHHS prior to specimens being submitted from the healthcare facility he MDHHS BOL for COVID-19 testing)
MDSS ID number (MDHHS/LHD use):
Date of birth:/ Age:Sex: Female Male
Patient residence street address:City:
County: State: Zip Code:
Patient phone number(s):
Patient hospital ID (Medical Record) number:
Reporting healthcare facility:
Reporting healthcare facility contact name and title:
Healthcare facility contact phone number:

For additional information about 2019 Novel Coronavirus (COVID-19), please see:

https://www.cdc.gov/coronavirus/2019-ncov/index.html https://www.michigan.gov/coronavirus

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CDC	2019-nCoV ID:	Form App	proved: OMB: 0920-1011 Exp. 4/23/2020
PATIENT IDENT	TIFIER INFORMATION	I IS NOT TRANSMITTED TO CDC	
Patient first name F	Patient last name	Date of birth (MM/	/DD/YYYY):/
PATIENT IDENT	IFIER INFORMATION	I IS NOT TRANSMITTED TO CDC	
Human Inf		2019 Novel Coronaviru (PUI) and Case Repo	
Reporting jurisdiction: Reporting health department: Contact ID a: a. Only complete if case-patient is a known contact of prior sour CA102034567 -01 and CA102034567 -02. PFOR NNDSS reported.	CDC NND		Confirmed case CA102034567 has contacts
Interviewer information Name of interviewer: Last	First		
Affiliation/Organization:			
Basic information			
What is the current status of this person? Patient under investigation (PUI) Laboratory-confirmed case Report date of PUI to CDC (MM/DD/YYYY): Report date of case to CDC (MM/DD/YYYY): County of residence: State of residence: Race (check all that apply): Asian Black Native Hawaiia White Other, specify: Date of birth (MM/DD/YYYY): Age: Age units(yr/mo/day): Symptoms present If symptomatic, onset	n/Other Pacific Islander	Date of first positive specimen collection (MM/DD/YYYY):	Was the patient hospitalized? Yes No Unknown If yes, admission date 1
during course of illness: date (MM/DD/YYYY): Symptomatic Asymptomatic Unknown	Still symptomatic Symptoms resolved,	☐ Unknown symptom status , unknown date	Date of death (MM/DD/YYYY):/ Unknown date of death
Travel to Hubei lab Travel to mainland China And Travel to other non-US country lab specify:	e facility (as a patient, worker any of the following exportment of the following exportment of the following exportment of the following exportment of the following exposure of the facility of the following exposure of the facility of th	ker or visitor) in China?	

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).



Other, Specify:

Additional State/local Specimen IDs:

CDC 2019-nCoV ID:	
CDC ZUIS-HCUV ID.	

Form Approved: OMB: 0920-1011 Exp. 4/23/2020

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

Symptoms, clinical course, past medical history and social history Collected from (check all that apply): Patient interview Medical record review During this illness, did the patient experience any of the following symptoms? **Symptom Present?** Fever >100.4F (38C)c □Yes □No \Box Unk Unk Subjective fever (felt feverish) Yes No Chills Yes Νo Unk Muscle aches (myalgia) Yes Πo \Box Unk Runny nose (rhinorrhea) No Yes Unk Unk Sore throat Yes No Cough (new onset or worsening of chronic cough) ☐Yes No Unk Shortness of breath (dyspnea) Yes ΠNο Unk Yes No Unk Nausea or vomiting Yes No Unk Headache Yes No Unk Abdominal pain 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 Yes Пио Unknown Diabetes Mellitus Cardiovascular disease Yes □No Unknown Chronic Renal disease Yes ∏No Unknown Yes ПNо Unknown Chronic Liver disease □Yes □No Unknown Immunocompromised Condition Neurologic/neurodevelopmental □No Yes Unknown (If YES, specify) Yes No (If YES, specify) Other chronic diseases Unknown If female, currently pregnant Yes Пио Unknown No Yes Unknown Current smoker Yes No Unknown Former smoker Respiratory Diagnostic Testing Specimens for COVID-19 Testing Pos Pend. Not done Specimen Date State Lab Test Neg Specimen Sent to Type ID Collected CDC Tested NP Swab Influenza rapid Ag □ A □ B Influenza PCR □ A □ B **OP Swab** RSV Sputum H. metapneumovirus Other, Parainfluenza (1-4) Specify: Adenovirus Rhinovirus/enterovirus Coronavirus (OC43, 229E, HKU1, NL63) M. pneumoniae C. pneumoniae

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