

# COVID-19

- 1) Hospitalized patients with acute COVID-19 (i.e., microbiologically confirmed SARS-CoV-2)
- 2) Hospitalized patients with paediatric inflammatory multisystem syndrome/Kawasaki disease temporally associated with COVID-19
- 3) Non-hospitalized patients with acute COVID-19 (i.e., microbiologically confirmed SARS-CoV-2) AND chronic comorbid conditions

## CANADIAN PAEDIATRIC SURVEILLANCE PROGRAM

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## REPORTING INFORMATION

Report number:

Month of reporting:

Province:

Today's date:

Please complete the following sections for the case identified above. If the information asked below is not readily available, please leave it blank. Strict confidentiality of information will be assured.

### CASE DEFINITION FOR COVID-19 STUDY:

Report any new patient less than 18 years of age (up to the 18<sup>th</sup> birthday) who meets one of the following three case definitions:

- (1) HOSPITALIZED with acute COVID-19 (i.e., microbiologically confirmed SARS-CoV-2)
- (2) HOSPITALIZED with paediatric inflammatory multisystem syndrome (PIMS)/Kawasaki disease temporally associated with COVID-19, defined as:

- Persistent fever (>38 degrees Celsius for 3 or more days) and elevated inflammatory markers (C-reactive protein [CRP], erythrocyte sedimentation rate [ESR], or ferritin)

#### AND one or both of the following:

- Features of Kawasaki disease (complete or incomplete)
- Toxic shock syndrome (typical or atypical)

#### AND

- No alternative etiology to explain the clinical presentation

**IMPORTANT NOTE: Patients should be reported regardless of SARS-CoV-2 status**

- (3) NON-HOSPITALIZED with acute COVID-19 (i.e., microbiologically confirmed SARS-CoV-2) AND at least one of the following chronic comorbid conditions:

< 12 months of age	Asthma
Obesity	Chronic lung disease
Congenital heart disease	Chronic renal disease
Immunocompromising medications (high-dose steroids,* chemotherapy, biologics, immunomodulators)	Solid tumor or hematologic malignancy
Solid organ transplant	Bone marrow transplant
Primary or secondary immunodeficiency	Chronic neurologic or neurodevelopmental condition
Sickle cell disease or other chronic hematologic condition	Diabetes
Tracheostomy	Chronic rheumatologic or autoimmune disease
Inflammatory bowel disease or other chronic gastrointestinal or liver disease	Genetic/metabolic disease

\* Equivalent to at least 2 mg/kg or 20 mg/day of prednisone for at least two weeks

**PLEASE INDICATE CASE DEFINITION(S) BEING REPORTED (select all that apply):**

- ☐ Patient hospitalized with acute COVID-19
- ☐ Patient hospitalized with paediatric inflammatory multisystem syndrome (PIMS)/Kawasaki disease temporally associated with COVID-19
- ☐ Non-hospitalized patient with acute COVID-19 AND at least one chronic comorbid condition or < 12 months of age

**SECTION 1 – PATIENT INFORMATION**

1.1 Date of birth: \_\_\_\_ / \_\_\_\_ / \_\_\_\_  
DD MM YYYY

1.2 Sex: ☐ Male ☐ Female ☐ Intersex

1.3 First 3 digits of patient's postal code: \_\_\_\_ \_\_\_\_ \_\_\_\_

1.4 Population groups (**select all that apply**):

- |   |   |  |  |
|---|---|--|--|
| <input type="checkbox"/> Arab   | <input type="checkbox"/> Black  | <input type="checkbox"/> Chinese                               | <input type="checkbox"/> Filipino              |
| <input type="checkbox"/> Japanese   | <input type="checkbox"/> Korean   | <input type="checkbox"/> Latin American                        | <input type="checkbox"/> White                 |
| <input type="checkbox"/> First Nations  | <input type="checkbox"/> Inuit  | <input type="checkbox"/> Métis                                 | <input type="checkbox"/> Unknown               |
| <input type="checkbox"/> Southeast Asian<br>(e.g., Vietnamese,<br>Cambodian, Laotian) | <input type="checkbox"/> South Asian<br>(e.g., East Indian,<br>Pakistani, Sri Lankan) | <input type="checkbox"/> West Asian<br>(e.g., Iranian, Afghan) | <input type="checkbox"/> Other, specify: _____ |

**ASSOCIATION WITH COVID-19**

1.5 Has the patient tested positive for SARS-CoV-2? ☐ Yes ☐ No ☐ Pending

*If yes, specify date:* \_\_\_\_ / \_\_\_\_ / \_\_\_\_  
DD MM YYYY

***If no or pending, proceed to question 1.6.***

1.5.1 *If yes, what was the method of microbiologic diagnosis (select all that apply):*

- ☐ Polymerase chain reaction (PCR) or nucleic acid amplification ☐ Serology ☐ Rapid point-of-care test
- ☐ Other, specify: \_\_\_\_\_
- ☐ Unknown

1.5.2 *If yes, what type(s) of samples tested positive for SARS-CoV-2 (select all that apply):*

- ☐ Nasal-pharyngeal swab ☐ Nose swab ☐ Throat swab ☐ Endotracheal tube (ETT) aspirate
- ☐ Blood (PCR) ☐ Blood (serology- prior to any IVIG) ☐ Blood (serology- following IVIG) ☐ Stool
- ☐ Other, specify: \_\_\_\_\_
- ☐ Unknown

1.6 Has the patient travelled (outside of the province, or internationally) in the 12 weeks prior to symptom onset?  
☐ Yes ☐ No ☐ Unknown

1.6.1 *If yes, where?* \_\_\_\_\_

1.6.2 Date of return (if known): \_\_\_\_ / \_\_\_\_ / \_\_\_\_  
DD MM YYYY

1.7 Did the patient have any **known close contact** (household or non-household) with a microbiologically **confirmed** case of COVID-19 in the 8 weeks prior to symptom onset? ☐ Yes ☐ No ☐ Unknown

1.7.1 *If yes, state relationship to patient (e.g., parent, teacher, friend):* \_\_\_\_\_

1.8 Did **any members in the household** become unwell with symptoms consistent with COVID-19 (**NOT** microbiologically confirmed) in the 8 weeks before the onset of symptoms in the patient?

☐ Yes ☐ No ☐ Unknown

1.8.1 *If yes, state who (e.g., sibling, parent):* \_\_\_\_\_

1.9 Is it suspected that the patient contracted COVID-19 while in hospital or long-term care facility (nosocomial infection)?

☐ Yes, in hospital ☐ Yes, in long-term care ☐ No ☐ Unknown

## SECTION 2 – POTENTIAL RISK FACTORS

2.1 Was the patient born prematurely? ☐ Yes ☐ No ☐ Unknown

2.1.1 If yes, gestational age at birth: \_\_\_\_\_ weeks \_\_\_\_\_ days

2.2 Does this patient vape? ☐ Yes ☐ No ☐ Unknown

2.3 Does this patient smoke? ☐ Yes ☐ No ☐ Unknown

2.3.1 If yes, specify (e.g., cigarettes, cannabis, other): \_\_\_\_\_

2.4 Has the patient experienced housing insecurity in last 4 weeks? ☐ Yes ☐ No ☐ Unknown

2.5 Are this patient's routine immunizations up to date for age? ☐ Yes ☐ No ☐ Unknown

2.5.1 If no, list vaccines that are not up to date for age: \_\_\_\_\_

2.6 Did this patient receive the current year's influenza vaccine? ☐ Yes ☐ No ☐ Unknown

2.7 Did this patient have any other epidemiologic risk factors for acquiring SARS-CoV-2? ☐ Yes ☐ No ☐ Unknown

2.7.1 If yes, specify: \_\_\_\_\_

## SECTION 3 – MEDICAL COMORBIDITIES AND TREATMENT

3.1 Does this patient have any chronic comorbid conditions? ☐ Yes ☐ No ☐ Unknown

**If no or unknown, please proceed to Section 4.**

**If yes, select all that apply:**

<input type="checkbox"/> Asthma <input type="radio"/> Not requiring daily controller medication <input type="radio"/> Requiring daily controller medication	<input type="checkbox"/> Chronic lung disease <input type="checkbox"/> Cystic fibrosis <input type="checkbox"/> Bronchopulmonary dysplasia/chronic lung disease of prematurity <input type="checkbox"/> Other, specify: _____
<input type="checkbox"/> Obesity Indicate height _____ cm and weight _____ kg	<input type="checkbox"/> Diabetes mellitus (DM) <input type="radio"/> Insulin-dependant DM <input type="radio"/> Non-insulin DM
<input type="checkbox"/> Congenital heart disease Specify type if known: _____ <input type="checkbox"/> Cardiomyopathy	<input type="checkbox"/> Chronic renal disease Specify type if known: _____ Dialysis dependent? <input type="radio"/> Yes <input type="radio"/> No
<input type="checkbox"/> Immunocompromising medications in last 4 weeks List medication(s): _____	<input type="checkbox"/> Malignancy (active and/or undergoing therapy) Specify type if known: _____
<input type="checkbox"/> Solid organ transplant List organ and year of transplant: _____	<input type="checkbox"/> Bone marrow transplant List type and year of transplant: _____
<input type="checkbox"/> Primary immunodeficiency Specify diagnosis if known: _____	<input type="checkbox"/> Secondary immunodeficiency Specify: _____
<input type="checkbox"/> Sickle cell disease or other chronic hematologic condition Specify: _____	<input type="checkbox"/> Neurologic and/or neurodevelopmental condition(s) <input type="checkbox"/> Epilepsy <input type="checkbox"/> Cerebral palsy <input type="checkbox"/> Genetic disorder, specify: _____ <input type="checkbox"/> Other, specify: _____
<input type="checkbox"/> Tracheostomy	<input type="checkbox"/> Chronic rheumatologic or autoimmune disease Specify: _____
<input type="checkbox"/> Inflammatory bowel disease Specify type if known: _____ <input type="checkbox"/> Other chronic gastrointestinal or liver condition Specify type if known: _____	<input type="checkbox"/> Genetic/metabolic disease Specify type if known: _____
<input type="checkbox"/> Other, specify: _____	

**SECTION 4 – CLINICAL PRESENTATION – COMPLETE ONLY FOR PATIENTS WITH ACUTE COVID-19 (MICROBIOLOGICALLY CONFIRMED SARS-COV-2 INFECTION)**

**\*\* If you are reporting a case of a HOSPITALIZED PATIENT WITH PIMS/KAWASAKI DISEASE, PLEASE PROCEED TO SECTION 5.**

4.1 Was the patient hospitalized? ☐ Yes ☐ No

**If no, proceed to question 4.2.**

4.1.1 If yes, date of admission: \_\_\_\_ / \_\_\_\_ / \_\_\_\_  
DD MM YYYY

4.1.2 If yes, was this patient admitted primarily for:

- ☐ Care related to symptoms consistent with acute COVID-19
- ☐ Care unrelated to symptoms consistent with acute COVID-19 (i.e., an incidental finding in a patient admitted for another reason)  
If yes, describe: \_\_\_\_\_
- ☐ Care required for purposes of isolation/infection control after testing positive for SARS-CoV-2 (i.e., supportive care for patients living in long-term care facilities and/or group homes)  
If yes, describe: \_\_\_\_\_
- ☐ Humanitarian/compassionate reasons  
If yes, describe: \_\_\_\_\_

4.2 Symptoms and signs at presentation (**select all that apply**):

- |                                      |   |   |  |
|--------------------------------------|---|---|--|
| <input type="checkbox"/> Fever       | <input type="checkbox"/> Respiratory distress | <input type="checkbox"/> Muscle aches     | <input type="checkbox"/> Conjunctivitis        |
| <input type="checkbox"/> Cough       | <input type="checkbox"/> Lethargy             | <input type="checkbox"/> Rash             | <input type="checkbox"/> Headache              |
| <input type="checkbox"/> Sore throat | <input type="checkbox"/> Seizures             | <input type="checkbox"/> Vomiting         | <input type="checkbox"/> Loss of smell         |
| <input type="checkbox"/> Runny nose  | <input type="checkbox"/> Coma                 | <input type="checkbox"/> Diarrhea         | <input type="checkbox"/> Loss of taste         |
| <input type="checkbox"/> Sneezing    | <input type="checkbox"/> Skin manifestations, | <input type="checkbox"/> Loss of appetite | <input type="checkbox"/> Other, specify: _____ |
- specify: \_\_\_\_\_

4.3 Total duration of symptoms prior to diagnosis of COVID-19, if known \_\_\_\_ days

4.4 During the course of COVID-19 illness, did the patient experience any of the following? (**Select all that apply**):

<b><u>Respiratory:</u></b> <input type="checkbox"/> Pneumonia <input type="checkbox"/> Bronchiolitis <input type="checkbox"/> Acute respiratory distress syndrome	<b><u>Inflammatory:</u></b> <input type="checkbox"/> Cytokine storm/macrophage activating syndrome	<b><u>Neurologic:</u></b> <input type="checkbox"/> Seizures <input type="checkbox"/> Stroke <input type="checkbox"/> Encephalitis <input type="checkbox"/> Encephalopathy <input type="checkbox"/> Acute necrotizing encephalopathy <input type="checkbox"/> Coma
<b><u>Hematologic:</u></b> <input type="checkbox"/> Coagulation dysfunction, Specify: - Prothrombin time (PT)/international normalized ratio (INR) _____ - Partial thromboplastin time (PTT) _____ - D-dimer _____ <input type="checkbox"/> Anemia <input type="checkbox"/> Lymphopenia <input type="checkbox"/> Neutropenia <input type="checkbox"/> Thrombocytosis	<b><u>Cardiac:</u></b> <input type="checkbox"/> Hypotension <input type="checkbox"/> Acute cardiac dysfunction (myocarditis, pericarditis), specify: _____	<b><u>Other:</u></b> <input type="checkbox"/> Skin changes, specify: _____ <input type="checkbox"/> Gastrointestinal symptoms (abdominal pain, vomiting, diarrhea) specify: _____ <input type="checkbox"/> Hepatitis, specify: _____ <input type="checkbox"/> Renal dysfunction, specify: _____ <input type="checkbox"/> Other, specify: _____

# SECTION 5 – CLINICAL PRESENTATION – COMPLETE ONLY FOR PATIENTS HOSPITALIZED WITH PIMS/KAWASAKI DISEASE TEMPORALLY ASSOCIATED WITH COVID-19

**\*\* If you are reporting a case of a PATIENT WITH ACUTE COVID 19 (MICROBIOLOGICALLY CONFIRMED SARS-COV-2 INFECTION, PLEASE PROCEED TO SECTION 6.**

5.1 Date of hospital admission: \_\_\_\_ / \_\_\_\_ / \_\_\_\_  
DD MM YYYY

5.2 Fever

5.2.1 Number of days of fever at diagnosis: \_\_\_\_ ☐ Unknown

5.2.2 Total number of days of fever (START of fever until complete resolution): \_\_\_\_ ☐ Unknown

5.3 Did this patient require hospitalization for acute COVID-19 in the past 8 weeks? ☐ Yes ☐ No ☐ Unknown

5.4 Is this a Kawasaki disease re-occurrence? ☐ Yes ☐ No ☐ Unknown

5.4.1 If yes, what was the month and year of the most recent previous diagnosis of Kawasaki disease?  
\_\_\_\_ month \_\_\_\_ year

5.5 **Clinical features**

Yes No Unknown

5.5.1 Changes in the peripheral extremities (e.g., redness, swelling)

☐ ☐ ☐

5.5.2 Rash

☐ ☐ ☐

If yes, specify: \_\_\_\_\_

5.5.3 Bilateral bulbar conjunctival injection without exudate

☐ ☐ ☐

5.5.4 Changes in the lips or oral cavity

☐ ☐ ☐

5.5.5 Cervical lymphadenopathy >1.5 cm diameter

☐ ☐ ☐

5.5.6 Abdominal pain/vomiting/diarrhea

☐ ☐ ☐

5.5.7 Coagulation dysfunction

If yes, specify prothrombin time (PT)/international normalized ratio (INR),  
partial thromboplastin time (PTT): \_\_\_\_\_

5.5.8 Periungual desquamation

☐ ☐ ☐

5.6 Did the patient present with clinical features consistent with shock/hypotension?

☐ Yes ☐ No ☐ Unknown

5.7 **Laboratory features**

5.7.1 Highest C-reactive protein (CRP) \_\_\_\_\_

5.7.2 Highest erythrocyte sedimentation rate (ESR) \_\_\_\_\_

5.7.3 Highest ferritin \_\_\_\_\_

5.7.4 Highest D-dimer \_\_\_\_\_

5.7.5 Highest troponin \_\_\_\_\_

5.7.6 Highest liver enzymes (aspartate aminotransferase test [AST], alanine aminotransferase test [ALT], lactate dehydrogenase [LDH], bilirubin) \_\_\_\_\_

5.7.7 Lowest sodium \_\_\_\_\_

5.7.8 Lowest albumin \_\_\_\_\_

5.7.9 Lowest platelet count (while an inpatient) \_\_\_\_\_

5.7.10 Highest platelet count (if patient seen in follow-up) \_\_\_\_\_

5.8 Was an echocardiogram performed? ☐ Yes ☐ No ☐ Unknown

**If no or unknown, proceed to section 6.**

5.8.1 If yes:

(Note: If multiple echocardiograms were performed, please provide results from the study with the most severe findings)

Was coronary ectasia present? ☐ Yes ☐ No Maximum size (by z-score): \_\_\_\_\_

Was coronary dilation present? ☐ Yes ☐ No Maximum size (by z-score): \_\_\_\_\_

Were one or several aneurysms present? ☐ Yes ☐ No Maximum size (by z-score): \_\_\_\_\_

Was decreased heart function present?

○ Yes ○ No *If yes, specify ejection fraction:* \_\_\_\_\_

Were other cardiac findings present?

☐ Myocarditis ☐ Pericardial effusion☐ Valvular insufficiency ☐ Other, specify: \_\_\_\_\_**SECTION 6 – CONCURRENT INFECTIONS AND IMAGING**6.1 Were there any other confirmed **concurrent viral** infections? ○ Yes ○ No ○ Unknown6.1.1 *If yes, select all that apply:* ☐ Respiratory syncytial virus ☐ Influenza ☐ Other, specify: \_\_\_\_\_6.2. Were there any **other concurrent** microbiologically confirmed infections? ○ Yes ○ No ○ Unknown6.2.1 *If yes, specify:* \_\_\_\_\_6.3 Were there any other concurrent **clinically diagnosed** infections? ○ Yes ○ No ○ Unknown6.3.1 *If yes, specify:* \_\_\_\_\_6.4 Did the patient have any of the following imaging? (**Select all that apply**):☐ Chest X-ray: ○ Yes ○ No *If yes, describe main findings:* \_\_\_\_\_☐ CT scan: ○ Yes ○ No *If yes, describe anatomical location(s) and main findings:* \_\_\_\_\_☐ MRI scan: ○ Yes ○ No *If yes, describe anatomical location(s) and main findings:* \_\_\_\_\_☐ Other, specify: \_\_\_\_\_**SECTION 7 – TREATMENT**

## 7.1 Treatment

	Yes	No	Name of product(s), if applicable
Bronchodilators	<input type="radio"/>	<input type="radio"/>	
Azithromycin	<input type="radio"/>	<input type="radio"/>	N/A
Other antibiotics	<input type="radio"/>	<input type="radio"/>	1 2 3
Remdesivir			
Other antivirals	<input type="radio"/>	<input type="radio"/>	1 2 3
Steroids	<input type="radio"/>	<input type="radio"/>	
Hydroxychloroquine	<input type="radio"/>	<input type="radio"/>	
Chloroquine	<input type="radio"/>	<input type="radio"/>	
Immunoglobulin (IVIG) <i>If yes, specify number of doses:</i> _____	<input type="radio"/>	<input type="radio"/>	
Aspirin	<input type="radio"/>	<input type="radio"/>	
Anti-TNF	<input type="radio"/>	<input type="radio"/>	
Anti-IL-1 (e.g., anakinra, canakinumab)	<input type="radio"/>	<input type="radio"/>	
Anti-IL-6 (e.g., tocilizumab)	<input type="radio"/>	<input type="radio"/>	
Prophylactic anticoagulation	<input type="radio"/>	<input type="radio"/>	
Therapeutic anticoagulation	<input type="radio"/>	<input type="radio"/>	
Mechanical anticoagulation (e.g., antiembolism stockings, pneumatic boots)	<input type="radio"/>	<input type="radio"/>	
Other (e.g., blood products, other antiplatelet therapies)	<input type="radio"/>	<input type="radio"/>	

- 7.2 Is this patient enrolled in a clinical trial and receiving novel therapy (e.g., convalescent plasma, antiretrovirals)? ☐ Yes ☐ No ☐ Unknown  
 7.2.1 If yes, provide details: \_\_\_\_\_
- 7.3 What was the highest level of care required: ☐ Managed at home ☐ Inpatient ward ☐ ICU  
 7.3.1 If admitted to ICU, what was the total duration of ICU admission? \_\_\_\_\_ days  
 Is the patient still in ICU? ☐ Yes ☐ No
- 7.4 Did the patient require any of the following forms of respiratory/other support? **(Select all that apply):**
- ☐ Increased baseline home oxygen
  - ☐ Low-flow oxygen
  - ☐ High-flow nasal canula
  - ☐ Non-invasive ventilation (e.g., CPAP or BiPAP)
  - ☐ Conventional mechanical ventilation
  - ☐ High-frequency oscillatory ventilation
  - ☐ Nitric oxide (NO)
  - ☐ Extracorporeal membrane oxygenation (ECMO)
  - ☐ Vasopressors
  - ☐ Surgical thrombectomy
  - ☐ Hemofiltration
- 7.5 Did the patient experience any other complications over the course of their illness? ☐ Yes ☐ No ☐ Unknown  
 7.5.1 If yes, specify: \_\_\_\_\_
- 7.6 What was the final disposition of the patient? ☐ Remains at home ☐ Remains in hospital  
☐ Discharged home from hospital ☐ Transferred to another facility ☐ Died ☐ Unknown  
 7.6.1 If patient was transferred to another facility, specify reason: \_\_\_\_\_  
 7.6.2 If patient died, specify cause of death: \_\_\_\_\_

## SECTION 8 – FOLLOW-UP

- 8.1 Additional details: \_\_\_\_\_  
 \_\_\_\_\_  
 \_\_\_\_\_  
 \_\_\_\_\_

- ☐ I agree to be contacted by the CPSP for further information on this questionnaire.  
☐ I do not wish to be contacted by the CPSP for further for information on this questionnaire.

## LONG-TERM IMPACT

A separate study may be conducted to better understand the long-term impact and epidemiology of COVID-19 in children and youth. Do you agree to be contacted by the study team with follow-up questions about this case in the future? ☐ Yes ☐ No

(If yes, you agree to the CPSP releasing your contact information to the study team (led by Dr. Shaun Morris) for potential follow-up.)

**SECTION 9 – REPORTING PHYSICIAN**

First name \_\_\_\_\_ Surname \_\_\_\_\_  
Address \_\_\_\_\_  
City \_\_\_\_\_ Province or territory \_\_\_\_\_ Postal code \_\_\_\_\_  
Telephone number \_\_\_\_\_ Fax number \_\_\_\_\_  
E-mail \_\_\_\_\_ Date completed \_\_\_\_\_

**THANK YOU FOR COMPLETING THIS QUESTIONNAIRE. YOUR PARTICIPATION IS VITAL.**

COVID-19 06/2020