Appendix 1: Case report form

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

100-2305 St. Laurent Blvd. Ottawa, ON. K1G 4J8 Tel: 613-526-9397, ext. 239

Fax: 613-526-3332 cpsp@cps.ca www.cpsp.cps.ca

	INIMATION
Report number:	
Month of reporting:	
Province:	

REPORTING INFORMATION

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 18th 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 <u>at least one</u> 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: /

1.2	Sex: O Male	> Female	O Intersex		
1.3	First 3 digits of	patient's po	ostal code:		
1.4	Population grou	ups (select	all that apply):		
	☐ Arab ☐ Japanese ☐ First Nations ☐ Southeast A (e.g., Vietnames Cambodian, Lac	s (sian (☐ Black ☐ Korean ☐ Inuit ☐ South Asian (e.g., East Indian, Pakistani, Sri Lankan)	☐ Chinese ☐ Latin American ☐ Métis ☐ West Asian (e.g., Iranian, Afghan)	☐ Filipino☐ White☐ Unknown☐ Other, specify:
ASS	OCIATION WITH	H COVID-1	9		
1.5	Has the patient t	tested posit	tive for SARS-CoV-2?	O Yes O No O Pend	ing
	If yes, specify da	ate:/_	/		
	If no or pending	g, proceed	to question 1.6.		
	□ Polymorare test □ Other □ Unknot 1.5.2 If yes, where □ Nasal	nerase cha -, specify: _ own hat type(s) I-pharyngea	in reaction (PCR) or nu of samples tested posi al swab □ Nose swal	tive for SARS-CoV-2 (sele ct) Throat swab	Rapid point-of- ct all that apply): dotracheal tube (ETT) aspirate
Sto		I(PCR)	i Blood (serology- prior	to any IVIG) UBlood (se	rology- following IVIG)
		, specify: _			
	☐ Unkno	own			
1.6	onset? O Yes O No 1.6.1 If yes, w 1.6.2 Date of r	O Unknowhere?	own nown): / / / / / / /_		
			·	or to symptom onset? O Y arent, teacher, friend):	
1.8	microbiologicall O Yes O No	ly confirme O Unkno	d) in the 8 weeks before own	well with symptoms consise the onset of symptoms in	•
1.9	(nosocomial infe	ection)?	tient contracted COVIDs, in long-term care	-19 while in hospital or long No Ͻ Unknown	g-term care facility

SEC	TION 2 – POTENTIAL RISK FACTORS						
2.1	Was the patient born prematurely? • Yes • No 2.1.1 <i>If yes</i> , gestational age at birth:wee						
2.2	Does this patient vape? • O Yes O No O Unknown	own					
2.3 Does this patient smoke? • Yes • No • Unknown 2.3.1 <i>If yes</i> , specify (e.g., cigarettes, cannabis, other):							
2.4 Has the patient experienced housing insecurity in last 4 weeks? O Yes O No O Unknown							
2.5	Are this patient's routine immunizations up to date for a 2.5.1 <i>If no</i> , list vaccines that are not up to date for a	•					
2.6	Did this patient receive the current year's influenza	vaccine? O Yes O No O Unknown					
	Did this patient have any other epidemiologic risk fa nown 2.7.1 <i>If yes,</i> specify:	ctors for acquiring SARS-CoV-2? • Yes • No •					
SEC	TION 3 – MEDICAL COMORBIDITIES AND TREAT	rment .					
	Does this patient have any chronic comorbid conditi						
0.1	If no or unknown, please proceed to Section 4.	ons: 9 100 9 tto 9 onknown					
	If yes, select all that apply:						
	☐ Asthma ☐ Not requiring daily controller medication	☐ Chronic lung disease ☐ Cystic fibrosis					
	Requiring daily controller medication	 □ Bronchopulmonary dysplasia/chronic lung disease of prematurity □ Other, specify: 					
	Obesity Indicate height cm and weight kg	☐ Diabetes mellitus (DM) ☐ Insulin-dependant DM ☐ Non-insulin DM					
	☐ Congenital heart disease Specify type if known: ☐ Cardiomyopathy	☐ Chronic renal disease Specify type if known:					
		Dialysis dependent? O Yes O No					
	☐ Immunocompromising medications in last 4 weeks List medication(s):	☐ Malignancy (active and/or undergoing therapy) Specify type if known: ———					
	Solid organ transplant List organ and year of transplant:	☐ Bone marrow transplant List type and year of transplant:					
	Primary immunodeficiency Specify diagnosis if known:	Secondary immunodeficiency Specify:					
	Sickle cell disease or other chronic hematologic condition Specify:	 □ Neurologic and/or neurodevelopmental condition(s) □ Epilepsy □ Cerebral palsy 					

				☐ Genetic	disorder, specify:	
				☐ Other, sp	pecify:	
	☐ Tracheostomy			Chronic rhe Specify:	eumatologic or autoimmu	ne disease
	Other chronic g Specify type if k	known: astrointestinal or liver cond known:	dition _	Genetic/me Specify typ		
	Other, specify:					
(MIC ** If	ROBIOLOGICALLY	PRESENTATION - COM CONFIRMED SARS-CO case of a HOSPITALIZE I 5.	V-2 INFE	CTION)		
4.1	Was the patient hos	spitalized? O Yes O No)			
	If no, proceed to o	uestion 4.2.				
	4.1.1 <i>If yes</i> , date of	admission://_MM	YYYY			
	•	s patient admitted primarily				
		ed to symptoms consisten				
	patient	ated to symptoms consist	ent with a	cute COVID-	-19 (i.e., an incidental fin	ding in a
		or another reason)				
		scribe: ired for purposes of isolation		n control aft	er testing positive for SA	PS CoV 2
	(i.e.,	ned for purposes or isolation	OH/HHECKIC	ni control art	er testing positive for SA	N3-00V-2
	supportive	e care for patients living in scribe:	_		es and/or group homes)	
	Humanitar	ian/compassionate reasor	าร			
4.2	Symptoms and sign	s at presentation (select a	all that ap	ply):		
	□ Fever □ Cough □ Sore throat □ Runny nose □ Sneezing	□ Respiratory distress □ Lethargy □ Seizures □ Coma □ Skin manifestations, specify:	☐ Musc ☐ Rash ☐ Vomi	le aches ting	☐ Conjunctivitis☐ Headache☐ Loss of smell☐ Loss of taste☐ Other, specify:	
4.3	Total duration of syr	nptoms prior to diagnosis	of COVID	-19, if knowr	n days	
	During the course o	f COVID-19 illness, did the			 •	ect all that

	Respiratory:	Inflammatory:	Neurologic:
	□ Pneumonia□ Bronchiolitis□ Acute respiratory distress syndrome	☐ Cytokine storm/macrophage activating syndrome	□ Seizures □ Stroke □ Encephalitis □ Encephalopathy □ Acute necrotizing encephalopathy □ Coma
	Hematologic:	Cardiac:	Other:
	□ Coagulation dysfunction, Specify: - Prothrombin time (PT)/international normalized ratio (INR) - Partial thromboplastin time (PTT) - D-dimer	☐ Hypotension☐ Acute cardiac dysfunction (myocarditis, pericarditis), specify:	☐ Skin changes, specify:☐ ☐ Gastrointestinal symptoms (abdominal pain, vomiting, diarrhea) specify: ☐ Hepatitis, specify:
	■ Anemia		☐ Renal dysfunction, specify:
	□ Lymphopenia□ Neutropenia□ Thrombocytosis		☐ Other, specify:
PIMS ** If y	KAWASAKI DISEASE TEMPORA	ON – COMPLETE ONLY FOR PATIEI ALLY ASSOCIATED WITH COVID-19 IENT WITH ACUTE COVID 19 (MICR ROCEED TO SECTION 6.	
5.1	Date of hospital admission:/	·/	
5.2	Fever 5.2.1 Number of days of fever at		olution): O Unknown
5.3	Did this patient require hospitalizate Unknown	tion for acute COVID-19 in the past 8 v	veeks? O Yes O No O
5.4		urrence? O Yes O No O Unknown nd year of the most recent previous dia	agnosis of Kawasaki disease?
5.5	Clinical features		Yes No
			Unknown
	5.5.1 Changes in the peripheral e5.5.2 Rash	extremities (e.g., redness, swelling)	0 0 0
	If yes, specify:		
	5.5.3 Rilateral hulbar conjunctiva		\circ

	5.5.4 Changes in the lips or oral cavity		O	•	O
	5.5.5 Cervical lymphadenopathy >1.5 cm diameter		O	•	O
	5.5.6 Abdominal pain/vomiting/diarrhea		•	•	0
5.6	 5.5.7 Coagulation dysfunction If yes, specify prothrombin time (PT)/internation partial thromboplastin time (PTT): 5.5.8 Periungual desquamation Did the patient present with clinical features consistent 		O	•	•
0.0	O Yes O No O Unknown	The Will Shook hypotonision:			
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 aminotransf [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	erase test [AST], alanine amir	notransfe	erase test	
5.8	Was an echocardiogram performed? O Yes O No	O Unknown			
	If no or unknown, proceed to section 6. 5.8.1 If yes: (Note: If multiple echocardiograms were performed, severe findings) Was coronary ectasia present?	please provide results from th			nost
	Was coronary dilation present?	O Yes O No Maximum si	ze (by z-	score):	
	Were one or several aneurysms present?	O Yes O No Maximum si	ze (by z-	score):	
	Was decreased heart function present?	O Yes O No If yes, speci	fy ejectic	n fractio	n:
_	Were other cardiac findings present?	☐ Myocarditis ☐ Valvular insufficiency ☐		dial effusi specify:	ion

SECTION 6 – CONCURRENT INFECTIONS AND IMAGING

6.1	•			□ Respiratory syncytial virus □ Influenza □ Other, specify:
6.2.	•			microbiologically confirmed infections? O Yes O No O Unknown
6.3	-			linically diagnosed infections? O Yes O No O Unknown
6.4	•	•		owing imaging? (Select all that apply): If yes, describe main findings:
	☐ CT scan:	O Yes	O No	If yes, describe anatomical location(s) and main findings:
	☐ MRI scan:	O Yes	O 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	O	0	
Azithromycin	O	O	N/A
Other antibiotics	0	0	1 2 3
Remdesivir			
Other antivirals	0	0	1 2 3
Steroids	0	0	
Hydroxychloroquine	0	0	
Chloroquine	0	O	
Immunoglobulin (IVIG) If yes, specify number of doses:	•	0	
Aspirin	0	0	
Anti-TNF	0	0	
Anti-IL-1 (e.g., anakinra, canakinumab)	0	•	
Anti-IL-6 (e.g., tocilizumab)	0	0	
Prophylactic anticoagulation	0	0	
Therapeutic anticoagulation	0	O	

	Mechanical anticoagulation	0	O					
	(e.g., antiembolism stockings, pneumatic boots)							
	Other (e.g., blood products,	0	O					
	other antiplatelet therapies)							
7.2	Is this patient enrolled in a clinic antiretrovirals)? O Yes O No 7.2.1 <i>If yes</i> , provide details:	O U	nknowi					
7.0								
.3	<u>-</u>		•	: O Managed at home O Inpatient ward O ICU				
				duration of ICU admission? days				
	Is the patient still in ICU?	O Ye	es On	NO				
7.4	Did the patient require any of the	e foll	owing f	forms of respiratory/other support? (Select all that apply):				
	☐ Increased baseline home ox	ygen						
	□ Low-flow oxygen							
	☐ High-flow nasal canula		_					
	□ Non-invasive ventilation (e.g			BIPAP)				
	☐ Conventional mechanical ve							
	☐ High-frequency oscillatory ve	entiiati	on					
	□ Nitric oxide (NO) □ Extracorporeal membrane oxygenation (ECMO)							
	□ Vasopressors	vygen	ation (i					
	☐ Surgical thrombectomy							
	☐ Hemofiltration							
	Did the patient experience any o	other o	complic	cations over the course of their illness? O Yes O No O				
	7.5.1 <i>If yes</i> , specify:							
7.6	•	ital () Tran	sferred to another facility O Died O Unknown				
	7.6.1 If patient was transferred	to an	other fa	acility, specify reason:				
	7.6.2 If patient died, specify ca	use o	f death	:				
SEC	TION 8 – FOLLOW-UP							
8.1	Additional details:							
<i>,</i>	, idalieriai detaile.							

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? O Yes O 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.

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