CKiD Chronic Kidney Disease in Children Cohort Study SECTION A: GENERAL INFORMATION

A1.	P/	ARTICIPANT ID: ENTER NUMBER (ONLY IF LABEL IS NOT AVAILABLE
			_ - _ -
A2.	Cł	KID VISIT #:	
A3.	FC	ORM VERSION:	0 3 / 1 5 / 2 1
A4.	DA	ATE OF THIS REPORT:	$\overline{M} \overline{M}' \overline{D} \overline{D}' \overline{Y} \overline{Y} \overline{Y} \overline{Y}$
A5.	FC	ORM COMPLETED BY (INITIALS)	
A6.	Pr	otocol type:	Regular Study Visit
A7.		this study visit an irregular ccelerated) visit?	Yes
A8.		d the participant receive a laboratory of the participant receive a laboratory	confirmed diagnosis of COVID and/or contact with a
			Yes
A9.	Sc	ource of Information	Participant/Family
		SECTION B: COVI	D-19 ILLNESS INFORMATION
B1.	a.	Did the participant receive a laborate Yes	• •
	b.	COVID-19?	hcare provider tell them that they had a suspected case of
	C.		
		Date:///	-
	d.	Did the participant have contact with Yes	1
	e.		he participant's suspected or confirmed COVID-19 illness? 1 2

	f.	At the time of the participant's suspected or and an individual with a confirmed case of C Yes				
		No				
B2.	a.	Is the participant currently sick with COVID-	19?			
		Yes 1				
		No	(Skip to B2c)			
	b.	Number of days since symptom onset				
	c.	(Skip to B3a) Don' Total length of illness (if recovered)	't know8	(Skip to B	3a)	
			month(s) don't know			
B3a.	Syr	mptoms present during COVID-19 illness (Sele	ect all that apply)			
				<u>Yes</u>	<u>No</u>	Don't know
	a.	Cough		1	2	-8
	b.	Rhinitis		1	2	-8
	C.	Fever		1	2	-8
	d.	Diarrhea		1	2	-8
	e.	Shortness of breath		1	2	-8
	f.	High temperature (greater than 38.0°C/100.4	°F)	1	2	-8
	g.	Myalgias (muscle aches)		1	2	-8
	h.	Fatigue or malaise		1	2	-8
	i.	Loss of taste or loss of smell		1	2	-8
	j.	Headache		1	2	-8
	k.	Pink eye		1	2	-8
	I.	Sore throat		1	2	-8
	m.	Runny nose		1	2	-8
	n.	Chills		1	2	-8
	0.	Loss of appetite		1	2	-8
	p.	Discomfort tightness or pressure in chest		1	2	-8
	q.	Vomiting		1	2	-8
	r.	Nausea		1	2	-8
	S.	Joint aches		1	2	-8
	t.	Seizure		1	2	-8
	u.	Dizziness		1	2	-8
	٧.	Altered consciousness or feeling like it was d	ifficult to stay awa	ke 1	2	-8
	W.	Abdominal pain		1	2	-8
	I.	Other		1	2	(Skip to B3b)
		1. Please specify:				

33b.	Presence of inflammatory syndrome in participant	(Select all that apply)
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		<u>Yes</u>	<u>No</u>	Don't know
a.	Diagnosis of multisystem inflammatory syndrome (MIS-C)	1	2	-8
b.	Kawasaki disease	1	2	-8
c.	Toxic shock syndrome	1	2	-8
l.	Other	1	2	(Skip to B4)
	Please specify:			

B4. Which of the following medications was the participant taking or prescribed prior to the COVID-19 illness?

VVhi	ch of the following medications was the participant	taking o	r preso <u>No</u>	Don't know	(
a.	Angiotensin-converting-enzyme Inhibitor (ACEi)	1	2	-8	
b.	Angiotensin II receptor blockers (ARB)	1	2	-8	
c.	Chloroquine or Hydroxychloriquine	1	2	-8	
d.	Steroids (IV/PO)	1	2	-8	
e.	Infliximabe/ Remicade	1	2	-8	
f.	Cyclophosphamide (IV)	1	2	-8	
g.	Cyclophosphamide (PO)	1	2	-8	
h.	Azathioprine	1	2	-8	
i.	Mycophenolate mofetil (MMF)	1	2	-8	
j.	Methotrexate	1	2	-8	
k.	Cyclosporin A	1	2	-8	
I.	Tacrolimus	1	2	-8	
m.	Everolimus	1	2	-8	
n.	Sirolimus	1	2	-8	
Ο.	Rituximab in last 6 months	1	2	-8	
p.	Basiliximab in last 6 months	1	2	-8	
q.	Alemtuzumab in last 6 months	1	2	-8	
r.	Other:	1	2	(Skip to B5)	
	1. Please specify:				

The next set of questions ask about the participant's laboratory results measured upon diagnosis, during treatment and at recovery.

B5.	White blood cell count (cells/uL)	
	a. Upon COVID-19 diagnosis:	8 Don't know8
	b. Peak value:	8 Don't know8
	c. Upon recovery:	8 Don't know8
		Not applicable1
B6.	C-reactive protein level (mg/L)	
	 a. Upon COVID-19 diagnosis: 	8 Don't know8
	b. Peak value:	8 Don't know8
	c. Upon recovery:	8 Don't know8
		Not applicable1
B7.	Serum creatinine (umol/L or mg/dL)	
	a. Upon COVID-19 diagnosis:	8 Don't know8
	b. Peak value:	B Don't know8
	c. Most recent value:	8 Don't know8
	d. Upon recovery:	8 Don't know8
	•	Not applicable1
d	uring the COVID-19 illness.	cute kidney infection (AKI) and kidney replacement therapy
B8.	, ,	sis of AKI as part of the COVID-19 disease episode?
	Yes	
	No	2
B9.	Did the participant receive renal rep treatment of the COVID-19 disease Yes	<u>.</u> 1
	e next set of questions ask about the ess.	treatment the participant received during the COVID-19
B10.	Was respiratory support (options liste Yes	ed in B11) needed as part of the treatment of the COVID-19 illness?
	No	2 (Skip to B12)
	Don't know	8 (Skip to B12)

B11. Level of respiratory support needed by participant at peak of COVID-19 illness (Select all that apply)

		<u>Yes</u>	<u>No</u>	Don't know	
a.	Supplemental oxygen	1	2	-8	
b.	High flow nasal cannula	1	2	-8	
c.	CPAP (Continuous Positive Airway Pressure)	1	2	-8	
d.	BiPAP (Bilevel Positive Airway Pressure)	1	2	-8	
e.	Conventional invasive ventilation	1	2	-8	
f.	Oscillatory invasive ventilation	1	2	-8	
g.	ECMO (Extracorporeal Membrane Oxygen)	1	2	-8	
h.	Other	1	2	(Skip to B12)	
	1. Please specify:				

B12. Specific therapies administered to participant to treat COVID-19 during the illness (Select all that apply)

		<u>Yes</u>	<u>No</u>	Don't know
a.	Remdesivir	1	2	-8
b.	ACEi	1	2	-8
C.	ARB	1	2	-8
d.	Chloroquine or hydroxychloroquine	1	2	-8
e.	Convalescent plasma	1	2	-8
f.	Azithromycin	1	2	-8
g.	Lopinavir/Ritonavir (Kaletra)	1	2	-8
h.	Ribavirin (Rebetol, Ribasphere, RibaPak, Copegus, Virazole, Moderiba)	1	2	-8
i.	Vitamin C	1	2	-8
j.	Zinc	1	2	-8
k.	Decadron	1	2	-8
l.	Other	1	2	(End Form)
	1. Please specify:			

END FORM