



ISARIC WHO Clinical Characterisation Protocol for Severe Emerging Infections UK (CCP-UK) CASE REPORT FORM GUIDANCE

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v9.9 11NOV2020

DESIGN OF THE CCP-UK CASE REPORT FORM (CRF)

This CRF is divided into a “**ADMISSION**” form (4 pages), a “**DAILY**” form (2 pages) for daily clinical and laboratory and data, an “**OUTCOME**” form (4 pages) and a “**WITHDRAWAL**” form (1 page).

HOW TO USE THIS CRF

The CRF is designed to complement the **Tier** of activity that a site has capacity and capability to work to. This is likely to vary over the course of an outbreak. The decision on which **Tier** to use is up to the Local Principal Investigator. All high-quality data is valuable for analysis.

Data can be collected as Tier Zero activity without consent including retrospectively and from deceased cases.

IMPORTANT CHANGES FOR SECOND WAVE OF COVID-19

Tier Zero will only include proved (positive test) COVID-19/ SARS-COV-2 cases and ANY admission following proved COVID-19/ SARS-COV-2 in the past 21 days regardless of setting of test (community or hospital tests).

Tiers 1 and 2 for now will only apply to patients with *re-infection, co-infection (flu/RSV) or inflammation (MIS-A/MIS-C)*. Ideally, data and samples will be collected with consent using Tier 2 of the protocol schedule.

Consent must be obtained for any biological sampling at Tier 1 and Tier 2.

Tier Zero	<p>For collection of data without consent from any case; current, past and deceased.</p> <p>Please complete the ADMISSION CRF and DAILY CRF for the first day of hospital admission (day 1), the DAILY CRF again for the first day of any ICU admission, then the INTERIM OUTCOME CRF at day 28, discharge or death (whichever occurs first), and FINAL OUTCOME when known.</p> <p>N.B. For patients receiving Remdesivir (RDV), please complete an extra DAILY CRF for first day that the drug is dosed and for day 14 after drug initiation (if patient remains admitted). Collection of this data is requested by the CMOs in all nations.</p> <p>OR</p> <p>For sites where caseload or facilities limit research capacity to deliver planned Tier 1 or Tier 2 activity.</p>
Tier 1 & 2	<p>Tier 1- For sites where facilities limit research capacity to deliver Tier 2 activity or where consent is only for single timepoint biological sampling.</p> <p>Tier 2- For sites with available resources to deliver Tier 2 activity per the protocol schedule and then with consent for multiple timepoint biological sampling.</p> <p>For these tiers please complete the ADMISSION CRF and DAILY CRF for the first day of hospital admission (day 1), the DAILY CRF for the third (d3), sixth (d6) and ninth (d9) days, the DAILY CRF again for the first day of any ICU admission, and then the INTERIM OUTCOME CRF at day 28, discharge or death (whichever occurs first), and FINAL OUTCOME when known.</p> <p>N.B. For patients receiving Remdesivir (RDV), please complete an extra DAILY CRF for first day that such a drug is dosed and for day 14 after drug initiation (if patient remains admitted). Collection of this data is requested by the CMOs in all nations.</p>

CASE REPORT FORMS

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GENERAL GUIDANCE

- The CRF is designed to collect data obtained through examination, interview and review of hospital notes. Data may be collected retrospectively if the patient is enrolled after the admission date or deceased after admission.
- Participant Identification Numbers consist of a 5-digit CPMS / ODS site code and a 4-digit participant number. You should obtain a site code by contacting your local R&D office or CCP@liverpool.ac.uk
- Participant numbers should be assigned sequentially for each site beginning with 0001. In the case of a single site recruiting participants on different wards, or where it is otherwise difficult to assign sequential numbers, it is acceptable to assign numbers in blocks. E.g. Ward X will assign numbers from 0001 onwards and Ward Y will assign numbers from 5001 onwards. Enter the Participant Identification Number at the top of every page.
- **Please generate a new subject ID for each re-admission**
- CRF data should be entered to the central database at <https://ncov.medsci.ox.ac.uk>
- REDCap registration access is obtained by contacting CCP.REDCap@liverpool.ac.uk
- Please contact us at CCP.REDCap@liverpool.ac.uk for help with database problems.

RULES DEFINING DAYS

1. Day of Admission = Day of Admission regardless, e.g. even if admitted 2 months ago for a broken hip.
2. For Community Acquired COVID-19 i.e. admitted with symptoms consistent with COVID-19, day 1 = first 24 hours of admission.
3. For those who are already admitted for any other reason and who subsequently test positive, day 1 = day the positive COVID-19 test **was collected**.
4. Rules 2 and 3 are important but we recognise that start of biological sampling for Tier 1 and 2 may be deferred or delayed for several reasons, e.g. due to a delay in the COVID-19 result being reported. If this happens, please take the d1 sample set as soon as possible and then d3 and d9 according to schedule, or as close as possible.
5. For Tier Zero date of enrolment is date on which the act of data collection started (no consent). For Tier 1 & 2 date of enrolment is date of consent.

Patients with *confirmed Covid-19* with any of the following syndromes should be recruited to tiers 1 or 2:



- **Re-infection.** The patient had Covid-19 more than 21 days ago:
 1. See criteria for identifying suspected re-infection on page 4.
 2. If you think a patient has suspected re-infection, please call 0300 365 4423 to discuss.
- **Co-infection.** The patient has **confirmed co-infection** with:
 1. Influenza A or B virus; or,
 2. Respiratory syncytial virus (RSV).
- **Clinical suspicion of Multisystem Inflammatory Syndrome in Adults (MIS-A) or Children (MIS-C)**

CASE REPORT FORMS

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- Complete every line of every section, except for where the instructions say to skip a section based on certain responses.
- Selections with square boxes (□) are single selection answers (choose one answer only). Selections with circles (○) are multiple selection answers (choose as many answers as are applicable).
- Some fields are considered **URGENT AND ESSENTIAL**. These are marked **BOLD AND UNDERLINED IN ALL CIRCUMSTANCES PLEASE PRIORITISE THESE DATA POINTS FOR URGENT UPLOAD.**
- Mark 'N/K' for any results of laboratory values that are not known or not available.
- Avoid recording data outside of the dedicated areas. Sections are available for recording additional information.
- We recommend writing clearly in black ink, using BLOCK-CAPITAL LETTERS.
- Place an (X) when you choose the corresponding answer. To make corrections, strike through (-----) the data you wish to delete and write the correct data above it. Please initial and date all corrections.
- In the case of a participant transferring between study sites, such as to a Nightingale Hospital, or other surge facility, it is preferred to maintain the same Participant Identification Number across the sites. When this is not possible a new Participant Identification Number should be assigned, the transferred participant will be linked by their identifiable data.
- Please keep all of the sheets for a single participant together e.g. with a staple or participant-unique folder.
- These three FRONT PAGES do not need to be retained.
- **DO NOT SEND CRFs to anyone by email or post.**
- See the training guide on how to send consent to CCP@liverpool.ac.uk using [SECURE] encryption
- The Dalhousie University Clinical Frailty Score is provided below for your reference.

Clinical Frailty Scale*

 <p>1 Very Fit – People who are robust, active, energetic and motivated. These people commonly exercise regularly. They are among the fittest for their age.</p>	 <p>7 Severely Frail – Completely dependent for personal care, from whatever cause (physical or cognitive). Even so, they seem stable and not at high risk of dying (within ~ 6 months).</p>	
 <p>2 Well – People who have no active disease symptoms but are less fit than category 1. Often, they exercise or are very active occasionally, e.g. seasonally.</p>	 <p>8 Very Severely Frail – Completely dependent, approaching the end of life. Typically, they could not recover even from a minor illness.</p>	
 <p>3 Managing Well – People whose medical problems are well controlled, but are not regularly active beyond routine walking.</p>	 <p>9. Terminally Ill - Approaching the end of life. This category applies to people with a life expectancy <6 months, who are not otherwise evidently frail.</p>	
 <p>4 Vulnerable – While not dependent on others for daily help, often symptoms limit activities. A common complaint is being "slowed up", and/or being tired during the day.</p>	<p>Scoring frailty in people with dementia</p> <p>The degree of frailty corresponds to the degree of dementia. Common symptoms in mild dementia include forgetting the details of a recent event, though still remembering the event itself, repeating the same question/story and social withdrawal.</p> <p>In moderate dementia, recent memory is very impaired, even though they seemingly can remember their past life events well. They can do personal care with prompting.</p> <p>In severe dementia, they cannot do personal care without help.</p>	
 <p>5 Mildly Frail – These people often have more evident slowing, and need help in high order IADLs (finances, transportation, heavy housework, medications). Typically, mild frailty progressively impairs shopping and walking outside alone, meal preparation and housework.</p>		
 <p>6 Moderately Frail – People need help with all outside activities and with keeping house. Inside, they often have problems with stairs and need help with bathing and might need minimal assistance (cuing, standby) with dressing.</p>		

* 1. Canadian Study on Health & Aging, Revised 2008.
2. K. Rockwood et al. A global clinical measure of fitness and frailty in elderly people. CMAJ 2005;173:489-495.

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CASE REPORT FORMS

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GENERAL GUIDANCE

Definitions:

INFLAMMATION - Children and adolescents

WHO preliminary criteria Multisystem inflammatory syndrome in children and adolescents temporally related to COVID-19

Children and adolescents 0–19 years of age with fever ≥ 3 days

AND any two of the following:

1. Rash or bilateral non-purulent conjunctivitis or muco-cutaneous inflammation signs (oral, hands or feet).
2. Hypotension or shock.
3. Features of myocardial dysfunction, pericarditis, valvulitis, or coronary abnormalities (including ECHO findings or elevated Troponin/NT-proBNP),
4. Evidence of coagulopathy (by PT, PTT, elevated d-Dimers).
5. Acute gastrointestinal problems (diarrhoea, vomiting, or abdominal pain).

AND

Elevated markers of inflammation such as ESR, C-reactive protein, or procalcitonin.

AND

No other obvious microbial cause of inflammation, including bacterial sepsis, staphylococcal or streptococcal shock syndromes.

AND

Evidence of COVID-19 (RT-PCR, antigen test or serology positive), or likely contact with patients with COVID-19

INFLAMMATION - Adults

We deliberately do not give criteria to avoid selection bias. Adults with an inflammatory should to be identified at clinical discretion.

If you think a patient meets these criteria or wish to discuss, please call 0300 365 4423.

RE-INFECTION

To be considered a suspected Covid-19 re-infection the patient should meet one prior Covid-19 criterion and one timing criterion. If you think a patient meets these criteria or wish to discuss, please call 0300 365 4423.

Prior Covid-19 criteria

- A positive test for virus (PCR or antigen) or antibodies, in the community or in a hospital. Evidence of this can be from the patient's own recollection, or from medical records.
- Patient-reported symptoms strongly suggestive of Covid-19, including cough, fever and altered taste/smell

Timing criteria

- If the patient was previously hospitalised with Covid-19, they must be more than 21 days from discharge from acute hospital (not including rehabilitation hospital).
- If the patient was not hospitalised but had symptoms of Covid-19, they must be more than 21 days from last symptoms.
- If the patient did not have symptoms, they must be more than 21 days from their last positive Covid-19 test.

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ADMISSION FORM

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Date of enrolment | | | | | | | | | | | | | | | | | | | | | | Site Location

CLINICAL INCLUSION CRITERIA

Proven infection with pathogen of Public Health Interest: ☐ YES ☐ NO

N.B. For acute covid-19, please only collect data from proven (laboratory test-positive) people.

OR

Adult or child who meets Case Definition for Multisystem Inflammatory Syndrome (MIS-C/MIS-A): ☐ YES ☐ NO

N.B. This group should be recruited regardless of covid-19 test as this syndrome can occur after mild disease in the community which has gone untested.

DEMOGRAPHICS

Sex at Birth: ☐ Male ☐ Female ☐ Not specified **Date of birth** | | | | | | | | | | | | | | | | | |

If date of birth is Not Known (N/K) record Age: | | | | years OR | | | | months

Postcode: | | | | | | | | | |

England & Wales NHS number , Scotland CHI: | | | | | | | | | | | | | |

NB Northern Ireland Health & Care Number is not being collected at this time

Ethnic group (check all that apply):

☐ Arab ☐ Black ☐ East Asian ☐ South Asian ☐ West Asian ☐ Latin American ☐ White ☐ Aboriginal/First Nations

☐ Other: | | | | | | | | ☐ N/K

Employed as a Healthcare Worker? ☐ YES ☐ NO ☐ N/K

Pregnant? ☐ YES ☐ NO ☐ N/K **If YES: Gestational weeks assessment:** | | | | weeks

POST PARTUM (within six weeks of delivery)? ☐ YES ☐ NO or ☐ N/K (skip this section - go to INFANT)

Pregnancy Outcome: ☐ Live birth ☐ Still birth **Delivery date:** | | | | | | | | | | | | | |

Has infant(s) been tested for Mother's infection? ☐ YES ☐ NO ☐ N/K **If YES:** ☐ Positive ☐ Negative

IF POSITIVE PLEASE COMPLETE A SEPARATE CASE REPORT FORM FOR THE INFANT(s)
INFANT – Less than 1 year old? ☐ YES ☐ NO (skip this section) **Birth weight:** | | . | | kg ☐ N/K

Gestational: ☐ Term birth (≥37wk GA) ☐ Preterm birth (<37wk GA) **if <37wk Estimated gestation** | | | | weeks ☐ N/K

Breastfed? ☐ YES ☐ NO ☐ N/K **If YES:** ☐ Currently breastfed ☐ Breastfeeding discontinued ☐ N/K

VACCINATION STATUS

Has the patient received a Covid-19 vaccine (open label licenced product) ☐ YES ☐ NO ☐ N/K

date if known: | | | | | | | | | | | | | |

has the patient been involved in a vaccine COVID trial? ☐ YES ☐ NO ☐ N/K

date if known (first trial vaccination): | | | | | | | | | | | | | | (please complete study participation CRF page 3 of outcome CRF)

Has patient received a 2020/21 seasonal influenza vaccine ☐ YES ☐ NO ☐ N/K

date if known: | | | | | | | | | | | | | |

ONSET AND ADMISSION

Date of first/earliest symptom: [][]/[][]/[][][][] OR ☐ Asymptomatic

Admission date at this facility: [D][D]/[M][M]/[2][0][Y][Y]

Is the patient being readmitted with Covid-19? (Please only add re-admission episodes for COVID patients remaining positive or new positive COVID test- Please assign new subject ID) ☐YES ☐NO ☐N/K

Previous participant ID: I II II II II I--I II II II I ☐ NK

Please provide reason for readmission: ☐ N/K

Is this a suspected re-infection with COVID-19? Defined as proven (PCR or antibody test) or highly probable (clinical case definition met) more than 21 days prior to this new laboratory proven covid-19 infection ☐YES ☐NO ☐N/K
If yes, please complete REINFECTION FORM and seek consent for biological sampling, ideally at Tier 2)

Is this a NIGHTINGALE or other SURGE FACILITY ☐YES ☐NO ☐N/K

Transfer from other facility? ☐YES-other facility is a study site ☐YES-other facility is not a study site ☐NO ☐N/K

If YES: Name of prior facility: ☐ N/K

If YES: Admission date at previous facility (DD/MM/YYYY): [D][D]/[M][M]/[2][0][Y][Y] ☐ N/K

If YES-Study Site: Participant ID # at previous facility: I I I I I I -- I I I I I

OR ☐ Same as above

VITAL SIGNS AT HOSPITAL ADMISSION -first available data at presentation/Admission to the facility.
(This section should refer to data from the date of admission to this facility)

Temperature: [] [] [] [] °C HR: [] [] [] [] beats per minute RR: [] [] [] [] breaths per minute

Systolic BP: [] [] [] mmHg Diastolic BP: [] [] [] mmHg Severe dehydration: ☐ YES ☐ NO ☐ N/K

Sternal capillary refill time >2seconds ☐YES ☐NO ☐N/K

Oxygen saturation: [][][]% **On:** ☐ Room air ☐ Any Oxygen therapy ☐ N/K

SIGNS AND SYMPTOMS- *This section should refer to the start of the COVID episode*

<u>History of fever</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Lower chest wall indrawing</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Cough</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Headache</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>with sputum production</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Altered consciousness/confusion</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>bloody sputum/haemoptysis</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Seizures</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Sore throat</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Abdominal pain</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Runny nose (Rhinorrhoea)</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Vomiting / Nausea</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Ear pain</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Diarrhoea</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Wheezing</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Conjunctivitis</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Chest pain</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Skin rash</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Muscle aches (Myalgia)</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Skin ulcers</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Joint pain (Arthralgia)</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Lymphadenopathy</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Fatigue / Malaise</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Bleeding (Haemorrhage)</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Shortness of breath (Dyspnoea)</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>If Bleeding: specify site(s):</u>	
<u>Disturbance or loss of taste</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Disturbance or loss of smell</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>(Ageusia)</u>		<u>(Anosmia)</u>	
		<u>None</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K

CO-MORBIDITIES (existing prior to admission)			
<u>Chronic cardiac disease, including congenital heart disease. (not hypertension)</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Obesity (as defined by clinical staff)</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Hypertension (physician diagnosed)</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Diabetes and Type</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> N/K
<u>Chronic pulmonary disease (not asthma)</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Diabetes (any) with complications</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Asthma (physician diagnosed)</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Diabetes (any) without complications</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Chronic kidney disease</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Rheumatologic disorder</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Moderate / severe liver disease</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Dementia</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Mild liver disease</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Malnutrition</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
<u>Chronic neurological disorder</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Smoking</u> <input type="checkbox"/> YES <input type="checkbox"/> Never smoked <input type="checkbox"/> Former smoker <input type="checkbox"/> N/K	
<u>Malignant neoplasm</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<u>Other relevant risk factor</u>	
<u>Chronic hematologic disease</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	
<u>AIDS / HIV</u>	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	If yes, specify _____	

Is the patient thought to be a member of a CLINICALLY EXTREMELY VULNERABLE GROUP
<p>Solid organ transplant recipients: <input type="checkbox"/>YES <input type="checkbox"/>NO <input type="checkbox"/>N/K</p> <p>People with specific cancers: <input type="checkbox"/>YES <input type="checkbox"/>NO <input type="checkbox"/>N/K</p> <ul style="list-style-type: none"> • people with cancer who are undergoing active chemotherapy • people with lung cancer who are undergoing radical radiotherapy • people with cancers of the blood or bone marrow such as leukaemia, lymphoma or myeloma who are at any stage of treatment • people having immunotherapy or other continuing antibody treatments for cancer • people having other targeted cancer treatments which can affect the immune system, such as protein kinase inhibitors or PARP inhibitors • people who have had bone marrow or stem cell transplants in the last 6 months, or who are still taking immunosuppression drugs <p>People with <u>severe</u> respiratory conditions including all cystic fibrosis, severe asthma requiring daily oral steroid or injectable maintenance therapy and severe chronic obstructive pulmonary requiring oxygen (COPD): <input type="checkbox"/>YES <input type="checkbox"/>NO <input type="checkbox"/>N/K</p> <p>People with rare diseases and inborn errors of metabolism that significantly increase the risk of infections (such as Severe combined immunodeficiency (SCID), homozygous sickle cell): <input type="checkbox"/>YES <input type="checkbox"/>NO <input type="checkbox"/>N/K</p> <p>People on immunosuppression therapies sufficient to significantly increase risk of infection: <input type="checkbox"/>YES <input type="checkbox"/>NO <input type="checkbox"/>N/K</p> <p>Women who are pregnant with significant heart disease, congenital or acquired: <input type="checkbox"/>YES <input type="checkbox"/>NO <input type="checkbox"/>N/K</p>

ADMISSION FORM

PRE-ADMISSION MEDICATION Were any of the following taken within 14 days of admission?	
Immunosuppressant e.g. oral (not inhaled) corticosteroids (not low dose hydrocortisone) <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Angiotensin converting enzyme inhibitors (ACEI)? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Anti-infectives for this illness episode prior to admission? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K If yes, specify: _____	Angiotensin II receptor blockers (ARBs)? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K Non-steroidal anti-inflammatory (NSAID)? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K

<u>CLINICAL FRAILTY SCORE</u> With reference to the Dalhousie University Clinical Frailty Score (see guidance page 3 of complete CRF)	
<u>Clinical Frailty Score</u>	<input type="text"/> value 1 to 9 or <input type="checkbox"/> N/K

[illegible]

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REINFECTION FORM

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SUSPECTED RE-INFECTION WITH COVID-19: DETAILS OF PREVIOUS INFECTION	
Did the patient have a positive PCR (virus) test for SARS-CoV-2? If yes, enter date of positive test: [_] [_] / [_] [_] / [2] [0] [_] [_]	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Did the patient have a positive antigen (virus) test for SARS-CoV-2? If yes, enter date of positive test: [_] [_] / [_] [_] / [2] [0] [_] [_]	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Did the patient have a positive serology (antibody) test for SARS-CoV-2? If yes, enter date of positive test: [_] [_] / [_] [_] / [2] [0] [_] [_]	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Symptom onset date of first/earliest symptom for previous infection: [_] [_] / [_] [_] / [2] [0] [_] [_] OR <input type="checkbox"/> Asymptomatic	

SIGNS AND SYMPTOMS for PREVIOUS COVID-19 episode			
History of fever	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Lower chest wall indrawing	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Cough	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Headache	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
with sputum production	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Altered consciousness/confusion	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
bloody sputum/haemoptysis	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Seizures	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Sore throat	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Abdominal pain	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Runny nose (Rhinorrhoea)	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Vomiting / Nausea	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Ear pain	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Diarrhoea	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Wheezing	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Conjunctivitis	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Chest pain	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Skin rash	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Muscle aches (Myalgia)	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Skin ulcers	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Joint pain (Arthralgia)	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Lymphadenopathy	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Fatigue / Malaise	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Bleeding (Haemorrhage)	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Shortness of breath (Dyspnoea)	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	If Bleeding: specify site(s):	
Disturbance or loss of taste (Ageusia)	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Disturbance or loss of smell (Anosmia)	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
		None	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
TREATMENT: During the previous episode, was the patient:			
Admitted to hospital:	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Treated with:	
Treated with oxygen:	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Dexamethasone	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Admitted to HDU/ICU:	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Any other steroid	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Receive invasive ventilation:	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Tocilizumab	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Receive extracorporeal membrane oxygenation (ECMO)	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Remdesivir	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
		Convalescent plasma	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
		Lopinavir/Ritonavir	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
		Interferon	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
		Chloroquine/Hydroxychloroquine	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K

ISARIC WHO Clinical Characterisation Protocol for Severe Emerging Infections UK

DAILY FORM complete per Tier of activity AND if research samples are collected Page 1 of 1

DAILY TREATMENT <i>(complete every line):</i>	
DATE OF ASSESSMENT (DD/MM/YYYY): [_D_][_D_]/[_M_][_M_]/[_2_][_0_][_Y_][_Y_]	
Record the worst value between 00:00 to 24:00 on day of assessment <i>(if Not Available write 'N/K'):</i>	
Is the patient in a high-level care area i.e. admitted to ICU/ITU/IMC/HDU <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	
Highest Temperature: [_][_] . [_] °C	
Any Supplemental Oxygen <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K FiO ₂ (0.21-1.0) [_] . [_] or [_] [_] % or [_] [_] L/min (highest)	
Oxygen saturation <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K SpO ₂ [_] [_] % (lowest) RR: [_] [_] breaths per minute (highest)	
AVPU Alert [_] Verbal [_] Pain [_] Unresponsive [_] or <input type="checkbox"/> N/K Glasgow Coma Score (GCS / 15) [_] [_] or <input type="checkbox"/> N/K	
Is the patient currently receiving, or has received (from 00:00 to 24:00) on day of assessment:	
Non-invasive respiratory support (e.g. NIV, BIPAP, CPAP)? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K Invasive ventilation? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	
High-flow nasal canula? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K ECLS/ECMO? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	
DAILY LABORATORY RESULTS	
Record the values of laboratory results taken between 00:00 to 24:00 on day of assessment <i>(if Not Available write 'N/K, if multiple record the values for the blood draw taken closest to midday'):</i>	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K Haemoglobin _____ <input type="checkbox"/> g/L or <input type="checkbox"/> g/dL	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K WBC count _____ <input type="checkbox"/> x10 ⁹ /L or <input type="checkbox"/> x10 ³ /μL	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K Lymphocyte count _____ <input type="checkbox"/> cells/ μL or <input type="checkbox"/> x10 ⁹ /L or <input type="checkbox"/> x10 ³ /μL	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K Neutrophil count _____ <input type="checkbox"/> cells/ μL or <input type="checkbox"/> x10 ⁹ /L or <input type="checkbox"/> x10 ³ /μL	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K Platelets _____ <input type="checkbox"/> x10 ⁹ /L or <input type="checkbox"/> x10 ³ /μL Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K APTT/APTR _____	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K PT _____ seconds or Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K INR _____	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K ESR _____ mm/hr Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K AST/SGOT _____ U/L	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K Glucose _____ <input type="checkbox"/> mmol/L or <input type="checkbox"/> mg/dL	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K Blood Urea Nitrogen (urea) _____ <input type="checkbox"/> mmol/L or <input type="checkbox"/> mg/dL	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K Lactate _____ <input type="checkbox"/> mmol/L or <input type="checkbox"/> mg/dL	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K LDH [_] [_] [_] . [_] U/L Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K Procalcitonin [_] [_] . [_] [_] ng/mL	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K CRP [_] [_] [_] . [_] mg/L	
Done <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K eGFR _____ mL/min/1.73 m ² <input checked="" type="radio"/> CKD-EPI <input checked="" type="radio"/> MDRD <input checked="" type="radio"/> CG	
Most recent HbA1c _____ <input type="checkbox"/> N/K date of HbA1c [_D_][_D_]/[_M_][_M_]/[_2_][_0_][_Y_][_Y_]	
Chest X-Ray /CT performed? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K IF Yes: Were infiltrates present? <input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	

ISARIC CCP-UK RESEARCH SAMPLES	
Was a biological sample taken for research on this day?	<input type="checkbox"/> YES <input type="checkbox"/> NO
If yes, please record the KIT number:	KIT NUMBER [_C_][_C_][_P_][_][_][_][_][_]

ISARIC WHO Clinical Characterisation Protocol for Severe Emerging Infections UK

OUTCOME FORM

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PATHOGEN TESTING

Was pathogen testing done during this illness episode? ☐ YES ☐ NO ☐ N/K

(*NB Should be a YES as this is key eligibility criteria)

*please record the detail of any COVID-19 / SARS2-CoV-2 which may have been done in the community

Section 1: Pathogen Diagnosis Summary (Respiratory virus PCR or antigen tests -NOT serology/antibody tests)			
	Tested and POSITIVE (please tick)	Tested and NEGATIVE (Please tick)	NOT TESTED (please tick)
COVID-19 / SARS-CoV-2	Yes <input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Influenza virus NB: Please do not enter Haemophilus influenza or parainfluenza virus here – enter them under "other" below	Yes <input type="checkbox"/> Please confirm type: <input type="checkbox"/> A/H3N2 <input type="checkbox"/> A/H1N1pdm09 <input type="checkbox"/> A/H7N9 <input type="checkbox"/> A not typed other A <input type="checkbox"/> <input type="checkbox"/> B not typed <input type="checkbox"/> Other type (specify): _____	<input type="checkbox"/>	<input type="checkbox"/>
Respiratory syncytial virus (RSV)	Yes <input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Adenovirus	Yes <input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Other	Yes <input type="checkbox"/> please specify : _____		

Section 2: Pathogen Testing Details (Please record the details of all tests carried out during this illness episode below -including the details of the tests indicated above).			
	Select one:	Organism	Date sample obtained
Nasal and/ or throat swab	<input type="checkbox"/> Obtained: positive <input type="checkbox"/> Obtained: negative <input type="checkbox"/> Not obtained
Blood culture	<input type="checkbox"/> Obtained: positive <input type="checkbox"/> Obtained: negative <input type="checkbox"/> Not obtained
Sputum	<input type="checkbox"/> Obtained: positive <input type="checkbox"/> Obtained: negative <input type="checkbox"/> Not obtained
Deep respiratory sample (BAL/ETA)	<input type="checkbox"/> Obtained: positive <input type="checkbox"/> Obtained: negative <input type="checkbox"/> Not obtained

OUTCOME FORM

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MEDICATION: While hospitalised or at discharge, were any of the following administered?

Antiviral agent? ☐ YES ☐ NO ☐ N/K If YES, tick all the apply: **O**Ribavirin **O**Lopinavir/Ritonavir **O**Interferon alpha

OInterferon beta **O**Chloroquine / Hydroxychloroquine **O**Oseltamivir (Tamiflu®) **O**Zanamivir

OOther antiviral _____

ORemdesivir If YES: first dose: [_] [_] / [_] [_] / [_] [_] and last dose [_] [_] / [_] [_] / [_] [_]

OIL6 inhibitor If YES which ☐ Tocilizumab ☐ Other IL6 inhibitor _____

IL6 inhibitor first dose: [_] [_] / [_] [_] / [_] [_] and last dose [_] [_] / [_] [_] / [_] [_]

Antibiotic? ☐ YES ☐ NO ☐ N/K If YES: specify type(s): _____

Corticosteroid? ☐ YES ☐ NO ☐ N/K

If yes, please confirm type: ☐ Dexamethasone ☐ Methylprednisolone ☐ Prednisolone ☐ Other, please specify _____

Route: ☐ Oral ☐ Intravenous ☐ Inhaled, maximum daily dose: _____

If given Dexamethasone, was this given as 6mg once per day (od) ? ☐ YES ☐ NO ☐ N/K

If Other Dexamethasone dose, please confirm _____ mg

If Other Dexamethasone other frequency please confirm: **O**BD **O**TDS **O**QDS **O**Other _____

Antifungal agent? ☐ YES ☐ NO ☐ N/K If YES: which _____

Off-label / Compassionate Use medications? ☐ YES ☐ NO ☐ N/K If YES: which _____

Interleukin inhibitors ☐ YES ☐ NO ☐ N/K If YES: which _____

Convalescent plasma ☐ YES ☐ NO ☐ N/K

TREATMENT: At ANY time during hospitalisation, did the patient receive/undergo:

ICU or High Dependency Unit admission? ☐ YES ☐ NO ☐ N/K If YES, total duration: _____ days **O** still in ICU/HDU

If NO, ☐ Not Indicated ☐ Not appropriate*

(*Advanced care plan/discussion documented in notes regarding not for escalation of care beyond ward)

Date of ICU/HDU admission: [_] [_] / [_] [_] / [2] [0] [_] [_] ☐ N/K

ICU/HDU discharge date: [_] [_] / [_] [_] / [2] [0] [_] [_] ☐ N/K

Any Oxygen therapy? ☐ YES ☐ NO ☐ N/K

High-flow nasal canula? ☐ YES ☐ NO ☐ N/K

Non-invasive ventilation? (e.g. BIPAP, CPAP) ☐ YES ☐ NO ☐ N/K

Invasive ventilation (Any intubation)? ☐ YES ☐ NO ☐ N/K If YES, total duration: _____ days **O** still on

Prone Ventilation? ☐ YES ☐ NO ☐ N/K

Inhaled Nitric Oxide? ☐ YES ☐ NO ☐ N/K

Tracheostomy inserted? ☐ YES ☐ NO ☐ N/K

Extracorporeal (ECMO) support? ☐ YES ☐ NO ☐ N/K If YES, total duration: _____ days **O** still on

Renal replacement therapy (RRT) or dialysis? ☐ YES ☐ NO ☐ N/K If YES, total duration: _____ days **O** still on

Inotropes/vasopressors? ☐ YES ☐ NO ☐ N/K If YES, total duration: _____ days **O** still on

Blood Group (please check past as well as current medical record): **o**A **o**B **o**AB **o**O **o**N/K

OUTCOME FORM

COMPLICATIONS: At any time during hospitalisation did the patient experience:			
Viral pneumonia	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Cardiac ischemia	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Bacterial pneumonia	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Cardiac arrest	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Acute Respiratory Distress Syndrome	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Bacteraemia	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Cryptogenic organizing pneumonia (COP)	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Coagulation disorder / Disseminated Intravascular Coagulation	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Pneumothorax	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Deep vein thrombosis	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Pleural effusion	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Pulmonary thromboembolism	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Bronchiolitis	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Anaemia	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Meningitis / Encephalitis	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Rhabdomyolysis / Myositis	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Seizure	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Acute renal injury/acute renal failure	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Stroke / Cerebrovascular accident	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Gastrointestinal haemorrhage	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Other neurological complication	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Pancreatitis	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Congestive heart failure	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Liver dysfunction	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Endocarditis	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Hyperglycaemia	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Myocarditis/Pericarditis	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Hypoglycaemia	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Cardiomyopathy	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Other, if yes specify below	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K
Cardiac arrhythmia	<input type="checkbox"/> YES <input type="checkbox"/> NO <input type="checkbox"/> N/K	Other:	

STUDY PARTICIPATION
Is / Has the participant being/ been recruited to a trial or multi-centre study during the period of their current illness (including initiation in the community and hospital)? <input type="checkbox"/> YES <input type="checkbox"/> NO
IF YES , specify Name of study _____ Study Participant ID _____
Add another study? <input type="checkbox"/> YES <input type="checkbox"/> NO IF YES , specify Name of study _____ Study Participant ID _____
Add another study? <input type="checkbox"/> YES <input type="checkbox"/> NO IF YES , specify Name of study _____ Study Participant ID _____

OUTCOME FORM

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INTERIM OUTCOME: DAY 28 (i.e. 28 days from 'day 1' as per 'rules defining days')

Outcome: ☐ Discharged alive expected to survive

☐ Hospitalisation = Remains in Hospital \geq Day 28 after symptom onset

- if so

☐ Ongoing health care needs relating to this admission for COVID-19

OR

☐ Ongoing health care needs NOT related to COVID episode

OR

☐ Medically fit for discharge (COVID-19 resolved) but remains in hospital for other reason (e.g. awaiting suitable care in community, resident in long term health care or mental health facility)

☐ Transfer to other facility

☐ Palliative discharge

☐ Death

☐ N/K

Interim Outcome date: [][]/[][]/[][][][][][] ☐ N/K

If Discharged alive:

Ability to self-care at discharge versus before illness: ☐ Same as before illness ☐ Worse ☐ Better ☐ N/K

If Discharged alive: Post-discharge treatment:

Oxygen therapy? ☐ YES ☐ NO ☐ N/K

If Transferred: Facility name: _____ ☐ N/K

If Transferred: Is the transfer facility a study site? ☐ YES ☐ NO ☐ N/K

If a Study Site: Participant ID # at new facility: ☐ Same as above

☐ Different: [][][][][][]-[][][][] ☐ N/K

FINAL OUTCOME (If status has changed since day 28)

Outcome:

☐ Discharged alive expected to survive

☐ Palliative discharge

☐ Death

☐ Transfer to other facility

☐ N/K

Outcome date : [][]/[][]/[][][][][][] ☐ N/K

If Discharged alive:

Ability to self-care at discharge versus before illness: ☐ Same as before illness ☐ Worse ☐ Better ☐ N/K

If Discharged alive: Post-discharge treatment:

Oxygen therapy? ☐ YES ☐ NO ☐ N/K

If Transferred: Facility name: _____ ☐ N/K

If Transferred: Is the transfer facility a study site? ☐ YES ☐ NO ☐ N/K

If a Study Site: Participant ID # at new facility: ☐ Same as above

☐ Different: [][][][][][]-[][][][] ☐ N/K

**ISARIC WHO Clinical Characterisation Protocol for Severe Emerging Infections UK
WITHDRAWAL FORM****Page 1 of 1****WITHDRAWAL**Date of withdrawal: [_ D _] [_ D _] / [_ M _] [_ M _] / [_ 2 _] [_ 0 _] [_ Y _] [_ Y _] ☐ N/KType of withdrawal: ☐ Withdrawal from samples only ☐ Other Please specify: _____

Reason for withdrawal: _____