

Follow-up

Date of randomisation

Please only report events that occurred from first randomisation until 28 days later on this form (except for Q2).

Patient's date of birth *

yyyy-mm-dd

1. Which of following treatment(s) did the patient **definitely** receive as part of their hospital admission after randomisation? *

(NB Include RECOVERY study-allocated drug, only if given, PLUS any of the other treatments if given as standard hospital care)

- No additional treatment
- Lopinavir-ritonavir
- Corticosteroid (dexamethasone, prednisolone, hydrocortisone or methylprednisolone)
- Hydroxychloroquine
- Azithromycin or other macrolide (eg, clarithromycin, erythromycin)
- Tocilizumab or sarilumab
- Remdesivir
- Intravenous immunoglobulin
- Synthetic monoclonal antibodies (REGN10933+REGN10987)
- Aspirin
- Colchicine
- Baricitinib
- Anakinra

Please select number of days the patient received lopinavir-ritonavir

1
 2
 3
 4
 5
 6
 7
 8
 9
 10

Please select number of days the patient received corticosteroid (dexamethasone, prednisolone, hydrocortisone or methylprednisolone)

1
 2
 3
 4
 5
 6
 7
 8
 9
 10

Please select number of days the patient received hydroxychloroquine

- 1 2 3 4 5 6 7 8 9 10

Please select number of days the patient received azithromycin

- 0 1 2 3 4 5 6 7 8 9 10

Please select number of days the patient received other macrolides (eg, clarithromycin, erythromycin)

- 0 1 2 3 4 5 6 7 8 9 10

Please select number of doses of tocilizumab or sarilumab the patient received

- 1 >1

Please select number of days the patient received remdesivir

- 1 2 3 4 5 6 7 8 9 10

Please select the proportion of days the patient received aspirin during the first 28 days after randomisation (or from randomisation to date of discharge if this is sooner)

- Most days ($\geq 90\%$) Some days ($\geq 50\%$ < 90%) Few days (< 50% of days, but not zero) None

Please select number of days the patient received colchicine

- 1 2 3 4 5 6 7 8 9 10

Please select number of days the patient received baricitinib

- 1 2 3 4 5 6 7 8 9 10

Please select number of days the patient received anakinra

- 1
 2
 3
 4
 5
 6
 7

» Convalescent Plasma

How many convalescent plasma infusions did the patient receive?

This is convalescent plasma (i.e. collected from people recovered from COVID-19), not any standard fresh frozen plasma or other blood products that the patient may have been given

- 0
 1
 2

Were any infusions stopped early for any reason ie, the patient did not receive the full amount?

- Yes
 No

How many were stopped early?

- 1
 2

» Health Status

2. Was a COVID-19 test done for this patient at any point during the admission? *

(If multiple tests were done, and the results were positive and negative, please tick Yes - positive result and Yes - negative result)

- Yes - positive result
 Yes - negative result
 Not done

3. What is the patient's vital status? *

- Alive
 Dead

3.1 What is the patient's current hospitalisation status? *

- Inpatient
 Discharged

The patient has been enrolled in the trial for **NaN** days

3.1.1 Date follow-up form completed

yyyy-mm-dd

3.1.1 What was the date of discharge? *

yyyy-mm-dd

3.1 What was the date of death? *

yyyy-mm-dd

3.2 What was the underlying cause of death? **This can be obtained from the last entry in part 1 of the death certificate*

- COVID-19
- Other infection
- Cardiovascular
- Other

Please give details**4. Did the patient require any form of assisted ventilation (ie, more than just supplementary oxygen) from day of randomisation until 28 days later?** *

- Yes
- No

Please answer the following questions:**4.1 For how many days did the patient require assisted ventilation?** ***4.2 What type of ventilation did the patient receive?**

Yes

No

Unknown

CPAP alone**Non-invasive ventilation (eg, BiPAP)****High-flow nasal oxygen (eg, AIRVO)****Mechanical ventilation (intubation/tracheostomy)****ECMO**

Total number of days the patient received invasive mechanical ventilation (intubation/tracheostomy) from randomisation until discharge/death/28 days after randomisation

5. Has the patient been documented to have a NEW cardiac arrhythmia at any point since the main randomisation until 28 days later? *

- Yes
- No
- Unknown

5.1 Please select all of the following which apply

- Atrial flutter or atrial fibrillation
- Supraventricular tachycardia
- Ventricular tachycardia (including torsades de pointes)
- Ventricular fibrillation
- Atrioventricular block requiring intervention (eg, cardiac pacing)

6. Did the patient require use of renal dialysis or haemofiltration from main randomisation until 28 days later? *

- Yes
- No

7. During the first 28 days after randomisation (or until discharge if sooner), did the participant have a thrombotic event? *

- Yes
- No
- Unknown

7.1 Please indicate the type of thrombotic event

Select all that apply

- Pulmonary embolism
- Deep-vein thrombosis
- Ischaemic stroke
- Myocardial infarction
- Systemic arterial embolism
- Other

8. During the first 28 days after randomisation (or until discharge if sooner), did the participant *

8. During the first 28 days after randomisation (or until discharge if sooner), did the participant experience clinically-significant bleeding ie, intra-cranial bleeding or bleeding that required intervention (eg, surgery, endoscopy or vasoactive drugs) or a blood transfusion?

- Yes
- No
- Unknown

8.1 Please indicate the site(s) of bleeding

*

Select all that apply

- Intra-cranial
- Gastrointestinal
- Other

8.2 Please indicate which interventions were required to manage the bleed

*

Select all that apply

- Blood transfusion
- Surgery
- Endoscopy
- Vasoactive drugs (e.g. inotropes on ICU)
- None of the above

9. Please indicate if the participant participated in any other COVID-19 trials

Select all that apply

- PRINCIPLE
- REMAP-CAP
- Other treatment trial(s)
- COVID-19 vaccine trial(s)

Please give name of other treatment trial(s)

Please give name of COVID-19 vaccine trial(s)

10. Please enter UKOSS case ID if known

*

Enter the full UKOSS case ID ie, COR_123

(select if you do not know the UKOSS case ID)

- Not known