## **Randomisation Program**

Call Freefone 0800 138 5451 to contact the RECOVERY team for URGENT problems using the Randomisation Program or for medical advice. All NON-URGENT queries should be emailed to recoverytrial@ndph.ox.ac.uk

	Logged in as: RECOVERY Site
	Section A: Baseline and Eligibility
	Date and time of randomisation: 30 Dec 2021 14:00
Treating clinician  A1. Name of treating clinician	
Patient details A2. Patient surname	
Patient forename	
A3. NHS number	☐Tick if not available
A4. What is the patient's Nepali date of birth?	
What is the patient's date of birth?	□ Itick if estimated
A5. What is the patient's sex?	
Inclusion criteria  A6. Has consent been taken in line with the protoco?  If answer is No patient cannot be enrolled in the study	<u> </u>
A6.0 How was consent obtained?	V
A6.5 Does this patient have viral pneumonia? See protocol for typical features. If answer is No patient cannot be enroted in the study	Yes v
A7.0 Does the patient have proven SARS-CoV-2 infection?	Yes v
A7.0.1 What was lateral flow test result?	
A7.0.2 What was PCR test result?	
AS. Does the patient have any medical history that might, in the opinion of the attending clinician, put the patient at significant risk if they were to participate in the trial?	
A9. Symptom onset date:	~/ ~/ ~
A10. Date of hospitalisation:	
A11. Does the patient require oxygen?	
A12. Please select one of the following to describe the current level of ventilation support	
current level of ventilation support  A12.1 Enter latest oxygen saturation measurement (%)	
A12.2 Enter latest CRP measurement since admission to hospital (mg/L)	□Tick if not measured □Tick if greater than limit of measurement
Enter 0 if below the limit of measurement  A12.3 Enter latest creatinine measurement since	mg/dL ✓ □Tick if not measured
admission to hospital Please select correct units for the measurement (mg/dL or µmol/L).	
A12.4 Enter latest D-dimer measurement since admission to hospital Please select correct units for the measurement (mg/L or ng/mL). Enter 0 if below the limit of measurement	
A12.5 Has the patient received a COVID-19 vaccine?	<b>V</b>
A12.6 Has the patient received an influenza vaccine in the	
last 12 months?  Does the patient have any CURRENT comorbidities or of	
A13.1 Diabetes	
A13.2 Heart disease	<u> </u>
A13.3 Chronic lung disease	v
A13.4 Tuberculosis	v
A13.5 HIV	v
A13.6 Severe liver disease	v
A13.7 Severe kidney impairment (eGFR<30 or on dialysis)	
A13.7.1 Is the patient on dialysis or haemofiltration?	v
A13.9.0 Does their clinician consider the patient to be severely immunocompromised?	
A13.12 Has the patient received tocilizumab or sarilumab therapy during this admission?	~
A13.14 Current or planned treatment with neuraminidase inhibitor eg. osetamivir, zanamivir	
A13.15 Has the patient received	
casirivimab+imdevimab (Ronapreve) during this illness?  A13.16 Has the patient received sotrovimab during this	
illness?	
A13.17 Has the patient received molnupiravir during this illness?	V V
Are the following treatments UNSUITABLE for the pa If you answer Yes it means you think this patient shou	
A14E.1 High-dose corticosteroids	
A14F.1 Empagliflozin  Empagliflozin is NOT suitable if patient (i) has type 1 or post- pannreatectomy diabetes militus; or (ii) has a history of ketoacidosis; or (iii) has blood ketonos 21,5 mmol/L or urine ketones 2±2+; or (iv) is perganent or breastleeding Empagliflozin cannot be given via an enteral feeding tube.	
Are the following treatments available?  A15E.1 High-dose corticosteroids	•
A15F.1 Empagliflozin	
Current medication	
A16.1 Is the patient currently prescribed remdesivir?	
A16.2 Is the patient currently prescribed systemic corticosteroids (dexamethasone, prednisolone, hydrocortisione, methylprednisolone)? Please do not include topical or inhaled treatments	<u> </u>
A16.5 What venous thromboembolism prophylaxis is the patient receiving?  Standard — usual for hospitalised patients (not increased due to COVID-19); Higher dose — treatment dose or increased prophylaxis due to COVID-19	<b></b>
Please sign off this form once complete Surname:	
Forename:	
Professional email:	
	Continue Cancal