RANDOMISED EVALUATION OF COVID-19 THERAPY (RECOVERY)

Hospital Name: (use CAPITALS)	
Patient Name: (use CAPITALS)	
Study ID: (enter after randomisation)	

1. Information about the study has been provided to me: I confirm that I have read (or had read to me) and understood the Participant Information Leaflet (V19.1 19-Dec-2021) and I have had the opportunity to consider the information and ask questions. These have been answered satisfactorily.

2. Voluntary participation: I understand that my participation is voluntary and that I am free to withdraw at any time, without giving any reason, and without my medical care or legal rights being affected.

3. Access to study data about me: I give permission for relevant sections of my medical notes and information collected during the study to be looked at, in confidence, by authorised individuals from this hospital, the University of Oxford, and regulatory authorities to check that the study is being carried out correctly.

4. Access to my medical information: I agree that medical information collected by the doctors and hospitals which provide me with care and which may be located in local or national health and research organizations (including hospital admission, civil registration, audit and research data) may be provided to the study coordinating centre both during and for up to 10 years after my discharge. I understand that information that identifies me will be passed securely to such bodies to make this possible and that I can opt out of this at any time by writing to the coordinating centre team.

5. Data stored on computer: I understand that information about my progress in the study will be recorded on a computer database, and that this data will be stored on computers supervised by the University of Oxford. I understand that this information will be kept securely and confidentially.

6. GP: I understand that my GP may be informed of any issues relevant to my participation in the RECOVERY trial.

7. Samples: I am aware that a blood sample and nasal/mouth swabs may be sent to a central laboratory for measurement of coronavirus and antibodies against it and/or influenza virus.

8. Agreement to take part: I have read the information (or had it read to me), had an opportunity to ask questions and agree to take part in the above study.

……	…….	……../……../……
PRINTED name of participant	Signature	Today's date
……	…….	……../……../……
PRINTED name of person taking consent	Signature	Today's date

**1 copy for participant; 1 copy for researcher site file; 1 (original) to be kept in medical notes*

Hospital Name: (use CAPITALS)	
Patient Name: (use CAPITALS)	
Study ID: (enter after randomisation)	

If participant is not able to read the text and/or sign for themselves but has capacity to give consent

I witnessed accurate reading of the consent form to the potential participant, who could ask any questions and got satisfactory replies.

I confirm that they gave their consent freely.

……	…….	……../……../……..
PRINTED name of witness	Signature	Today's date
……	…….	……../……../……..
PRINTED name of person taking consent	Signature	Today's date

If participant lacks capacity to give consent due to the severity of their medical condition (e.g. acute respiratory failure or need for immediate ventilation) or prior disease:

I have read the information (or had it read to me) and had an opportunity to ask questions.

I have no other involvement in the RECOVERY trial.

I understand that the patient will be informed about the trial as soon as they have the capacity to do so and that if they wish, they will be able to withdraw from the study without it affecting their medical care.

I believe that if they were able to, the patient would wish to take part in this study.

……	…….	……../……../……..
PRINTED name of Legal Representative	Signature	Today's date

……

Relationship to participant *(or state 'professional' if clinician acting as legal representative)*

……	…….	……../……../……..
PRINTED name of person taking consent	Signature	Today's date

**1 copy for legal rep; 1 copy for researcher site file; 1 (original) to be kept in participant medical notes*

Invitation to participate

We are inviting people who have been admitted to hospital with COVID-19 and/or influenza pneumonia to consent to join this research study, which is comparing possible treatments. This form gives information about the study including the aims, risks and benefits of taking part.

WHAT YOU SHOULD KNOW ABOUT THIS RESEARCH STUDY:

1) Why is this research being done?

Your doctors have found that you have a lung disease called COVID-19 and/or influenza pneumonia ("flu"). COVID-19 is caused by a type of virus called SARS-CoV-2, or coronavirus for short. Influenza pneumonia is caused by a flu virus different to COVID-19. About 19 out of 20 patients who get these viruses get better without coming to hospital. Of those who are admitted to hospital, most also get better, but some may need oxygen or mechanical ventilation before they do so. However, a few percent do not get better.

This trial has already shown that low doses of a type of steroid, dexamethasone and other treatments reduce the risk of dying for some patients hospitalised with COVID-19. There are several other treatments which may turn out to be helpful (or possibly harmful) when added to the usual standard of care for either COVID-19 or influenza pneumonia. This study aims to find out whether any of these additional treatments are helpful.

2) What is the purpose of this study?

This study aims to compare several different treatments that may be useful for patients with COVID-19 and/or influenza pneumonia. These treatments have been recommended for testing by the expert panel that advises the Chief Medical Officer in England. Although these treatments show promise, nobody knows if any of them will help patients recover more effectively than the usual standard of care all patients at your hospital will receive.

The treatments for COVID-19, which may be given in addition to the usual care at your hospital, include a high dose steroid, dexamethasone (if you have low oxygen levels), a treatment for diabetes or heart failure called empagliflozin, a synthetic antibody treatment directed against the virus (called sotrovimab) and an antiviral drug called molnupiravir.

The treatments for influenza pneumonia, which may be given on top of your usual care, include two anti-viral treatments, oseltamivir and baloxavir and low-dose dexamethasone. At present, we don't know whether any of these will work. However, the side-effects are already well-known from other uses and so your doctor will be able to monitor you appropriately.

3) Who is doing the study?

The study is being conducted by researchers at the University of Oxford, which acts as the sponsor for the research, working with doctors at many hospitals across the UK.

4) Who is being included in the study?

Patients may be included in this study if they have COVID-19 and/or influenza pneumonia confirmed by a laboratory test, and are in hospital. Patients will not be included if the attending doctor thinks none of the study treatments are suitable for them. Patients may be included if they have previously been recruited into RECOVERY >6 months ago (although not into the same comparison more than once).

5) What happens next if I agree to be included in this study?

If you decide to join, you will be asked to sign the consent form. Next, brief details identifying you and answering a few questions about your health and medical conditions will be entered into a computer. If you are a woman of child-bearing potential, you will have a pregnancy test. If you might receive sotrovimab or molnupiravir a blood sample will be sent to a central laboratory for measurement of coronavirus and antibodies against it, and nasal and mouth swabs may be collected now and twice more in the next 5 days. If you have 'flu nasal and mouth swabs will be collected now and once more in 5 days. The results of all these tests will not be available to your medical team because they are for research and are not validated for clinical application, and the samples will be destroyed once testing is complete.

The computer will then allocate you at random (like rolling a dice) to one (or sometimes more) of the possible treatment options, depending on what illness you have and what your doctors think is suitable. In all cases this will include the usual standard of care for your hospital and it may also include additional treatment, which might be given by mouth or injection. Neither you nor your doctors can choose which of these treatments you will be allocated. Additional information about your health will be recorded and entered into the study computer. No additional visits will be required after you leave the hospital. Information about your health (before, during, and after the study) may be obtained from medical records or databases (including NHS Digital, Public Health England,

other equivalent bodies, and genetic or other research databases if you have provided samples to them) so that the study team can get more detailed or longer term information about the effects of the study treatments on your health for up to 10 years after your discharge. We may write to you to tell you about the trial periodically, but you will be able to opt-out of these communications if you prefer. Your GP may be informed of any issues relevant to your participation in the trial.

6) What are the possible benefits of being in the study?

We do not know if any of the treatments being tested will have additional benefits. Your study treatment may or may not help you personally, but this study should help future patients.

7) What are the possible risks of being in the study?

- Dexamethasone may also disturb sleep and increase the risk of infections. In people with diabetes it can raise blood sugar.
- Empagliflozin may cause urine or genital tract infections, like thrush. If you have diabetes, empagliflozin also lowers blood sugar in people taking insulin or some other diabetes treatments so your doctors may adjust the doses of those. It may also cause a condition called ketoacidosis (which rarely can be life-threatening), which is treated with a drip and insulin. You will be monitored for this with daily fingerprick blood or urine tests.
- Oseltamivir may cause headache, tummy upset and allergic reactions.
- Baloxavir rarely causes allergic reactions, but has no other known side effects.
- Sotrovimab is given by intravenous infusion and may cause allergic reactions during the infusion, but severe reactions have been rare.
- Molnupiravir may cause dizziness, headache, tummy upset and rashes. (Women taking molnupiravir should not get pregnant while taking the drug or for 4 days afterwards.)

There is also the unlikely possibility of a severe reaction to any study drug. Please ask your hospital doctor if you would like more information. Once you have been included in the study, you and your doctors will know which treatment the computer has allocated for you. Your doctors will be aware of whether there are any particular side effects that they should look out for.

Women who are pregnant may be included, however, the effect of some of the treatments on unborn babies is uncertain. Pregnant women will not receive empagliflozin or molnupiravir as it may be harmful in pregnancy or when breast-feeding. Dexamethasone and oseltamivir have previously been used in pregnancy for other medical conditions without safety concerns being raised. Baloxavir and sotrovimab are considered to have an acceptably low level of risk to use in pregnant women in this trial by a national expert panel; your medical team will discuss with you whether you would be willing to receive them.

8) Can I stop the study treatment or my participation early?

If you or your doctor want to stop the study treatment before the course has been completed, then you are free to do so. If you decide that you do not wish any more information to be collected about you, you are free to say so (although de-identified information that has been collected up to that point will continue to be analysed by the research team).

9) If I have any questions or problems, who can I call?

If you have any questions please speak to your hospital medical team. Further information about the study is available on the study website (www.recoverytrial.net).

10) What information do you hold about me and how do you keep it private?

All information about you and your health will be kept private. The only people allowed to look at the information will be the doctors who are running the study, authorised staff at Oxford University and your hospital, and the regulatory authorities who check that the study is being carried out correctly. A privacy notice is on the study website (https://www.recoverytrial.net/study-faq/data-privacy).

11) Do I have to take part?

Joining the study is voluntary. Your decision whether to take part will not affect the care you receive at this hospital.

12) Are there any financial costs or payments?

All trial treatments will be free. Neither you nor your medical staff will be paid for your participation in this study.

13) What else can you tell me?

The study has been approved by the Medicines and Healthcare products Regulatory Agency (MHRA) and by the Cambridge East Research Ethics Committee (Health Research Authority, ref 20/EE/0101). It is funded by UK Research and Innovation and the National Institute for Health Research, not the makers of any of the study treatments (who may provide the treatment free of charge to the trial). If we find out any new information that might affect your decision to stay in the study, we will give it to you. The University of Oxford, as Sponsor, has appropriate insurance in place in the unlikely event that you suffer any harm as a direct consequence of your participation in this study. NHS indemnity operates in respect of the clinical treatment that is provided.