IRAS ID: 288479 Consent Nurse v4 Dated 2021-01-12



COLLEGE OF MEDICINE AND HEALTH

Research title: COVID-NURSE: evaluation of the effects of a COVID-specific fundamental nursing care protocol compared to care as usual on experience of care for non-invasively ventilated patients in hospital with the SARS-CoV-2 virus: a randomised controlled trial.

CONSENT FORM FOR NURSE PARTICIPANTS

| | Participant Identification Number: | | | | | | | | | | |
|---------------------|---|--|--|--|--|--|--|--|------|----|-----------------------|
| Name of Researcher: | | | | | | | | | | | |
| | | | | | | | | | | | Please initial box |
| 1. | I confirm that I have read the information sheet dated [2021- 01-13] version no [4] for the above project. I have had the opportunity to consider the information, ask questions and have had these answered satisfactorily. | | | | | | | | | | |
| 2. | understand that my participation in providing my data is voluntary and that I am free to withdraw at any time without giving any reason and without my legal rights being affected. | | | | | | | | | | |
| 3. | I understand that I may also be asked to take part in a short online interview with follow up questions, which will be recorded for later transcription, at the end of the 6 week trial period by a researcher. | | | | | | | | | | |
| 4. | If interviewed, the transcription will be anonymized to protect participant confidentiality. | | | | | | | | | | |
| 5. | I understand that relevant sections of the data collected during the study may be looked at by members of the research team, individuals from the University of Exeter, Exeter Clinical Trials Unit or regulatory authorities, where it is relevant to my taking part in this research. I give permission for these individuals to have access to my data. | | | | | | | | | | |
| 6. | I understand that data I provide for the study will be stored on servers all of which are hosted within the Europe Economic Area | | | | | | | | | | |
| 7. | I understand that taking part involves que the purposes of research | lerstand that taking part involves questionnaire responses to be used for ourposes of research | | | | | | | | | |
| 8. | I understand that taking part involves inclute to 10 years. Data will be stored securely a | | | | | | | | d of | up | |

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| 9. | | nderstand that taking part involves anonymized data being shared with other searchers for use in future research projects; | | | | | | | | |
|--------|---|--|---|--------|--|--|--|--|--|--|
| 10. | I understand that if my Trust has be condition I will be asked to undertal FutureLearn platform. If I choose to General Data Protection Regulation read on the FutureLearn course we 'Learning Manager' at the Universit about the course. I understand that name and any other personally idea before collating and sending anony research team. I understand that no researcher at my hospital. | ke some online training on the oundertake this training, I will comprise privacy policies of FutureLearn, ebsite. I understand that a specificity of Exeter may support and intest the Learning Manager will remove the training to be an emissed records of training to be an | sent to the which I can course ract with me we my email, rese records, halysed by the | | | | | | | |
| 11. | I understand the research team would like my contact details so they may contact me to participate in an interview and let me know the results of the study [optional item] | | | | | | | | | |
| 12. | I understand that taking part involves anonymized data being used in reports published in academic publication, project website and media publication; | | | | | | | | | |
| 13. | I understand that taking part involves anonymized data being used in public engagement activities. | | | | | | | | | |
| 14. | I understand that there may be circumstances in which confidential information may need to be disclosed if information shared suggests a risk to myself or others. | | | | | | | | | |
| 15. | I agree to take part in the above pro | oject. | | | | | | | | |
| (F | Printed name of participant) | (Signature of participant) | (Date) | | | | | | | |
| • | Printed name of researcher aking consent) | (Signature of researcher) | (Date) | (Date) | | | | | | |
| , | When completed: 1 copy for participa | ant; 1 copy for researcher/project | file | | | | | | | |

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