### RECORD OF TELEPHONE CONSENT

[affix\_barcode]

Version X.X DD/MM/YYYY

Local Lead Investigator: [local\_lead\_investigator\_name]

**Please sign this form to indicate that the consenting party understands and agrees to the statements below:**

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| * I have read the information sheet (vX.X DD/MM/YYYY) for this study (or it has been read to me). I understand it and have had the opportunity to ask questions. * In my opinion the patient would have no objection to providing a DNA sample and for this sample to be analysed to look for genetic factors important in critical illness. * I can withdraw the patient from the study at any time without giving any reason. * There is a possibility that findings which are relevant to the patient will arise through this research. There is a process through which the patient can be informed of this. * The patient’s DNA, and data derived from their DNA, including the whole sequence of their genome, may be stored and used for future research. Researchers may include national or international scientists, companies and healthcare staff. To access the data, researchers must all be approved by an independent committee of experts, including clinicians, scientists and patients. There will be no access to the data by personal insurers or marketing companies. * Different aspects of the patient’s health data will be collected by the GenOMICC investigators, the study sponsor (enter sponsor’s name), and partner organisations. * I agree that the investigators of this study may contact the patient in the future to participate in future research studies, including clinical trials and studies unrelated to critical illness. |
| **Name of patient:** \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ |

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