**REMAP-CAP**

**Privacy notice**

This is a large global trial and we will follow the law by making sure your information is kept private and secure. UMC Utrecht is the sponsor for this study based in the Netherlands. We will be using information from you and your medical records in order to undertake this study and UMC Utrecht will act as the data controller for this study. This means that they are responsible for looking after your information and using it properly. UMC Utrecht will be storing study data, as outlined below, on servers based in Sydney, Australia. This information will be kept for 25 years after the study has finished.

Your rights to access, change or move your information are limited, as we need to manage your information in specific ways in order for the research to be reliable and accurate. If you withdraw from the study, we will keep the information about you that we have already obtained. To safeguard your rights, we will use the minimum personally-identifiable information possible.

You can find out more about how we use your information by contacting privacy@umcutrecht.nl.

Your hospital will collect information from you and your medical records for this research study in accordance with the sponsor’s instructions.

Your hospital will keep your name and contact details confidential and will not pass this information to UMC Utrecht. Your hospital will use this information as needed, to contact you about the research study, and make sure that relevant information about the study is recorded for your care, and to oversee the quality of the study. Certain individuals from UMC Utrecht and regulatory organisations may look at your medical and research records to check the accuracy of the research study. The people who analyse the information will not be able to identify you and will not be able to find out your name or contact details.

Minimal randomisation and clinical data will be collected on servers in Sydney, Australia which will collect some personal information about you for this global study. This information will include your initials, date of birth and gender, NHS number and basic eligibility health information. The information will be held securely with strict arrangements about who can access the information. With your permission, in order that we can contact you in 6 months and identify you in the Case Mix Programme database (as outlined above) your hospital will provide your name and telephone number to ICNARC (based in the UK), alongside some additional clinical data. Once you have been identified, the trial team will share your postcode and date of birth (held by the Case Mix Programme); along with your NHS number and name with NHS Digital. This will enable NHS Digital to provide us with information as described above.

Your hospital will keep identifiable information about you from this study for 25 years after the study has finished. When you agree to take part in a research study, the information about your health and care may be provided to researchers running other research studies in this organisation and in other organisations. These organisations may be universities, NHS organisations or companies involved in health and care research in this country or abroad. Your information will only be used by organisations and researchers to conduct research in accordance with the [UK Policy Framework for Health and Social Care Research](https://www.hra.nhs.uk/planning-and-improving-research/policies-standards-legislation/uk-policy-framework-health-social-care-research/)**.**

This information will not identify you and will not be combined with other information in a way that could identify you. The information will only be used for the purpose of health and care research, and cannot be used to contact you or to affect your care. It will not be used to make decisions about future services available to you, such as insurance. It is necessary for us to process your data as described to allow us to perform a task in the public interest (lawful basis).