You may be eligible to take part in our research study. If so you might be eligible to take part in our research study.

The main procedures in the trial are as follows:

- Questionnaires at the beginning of the study, at 3 months and 1 year
- Where possible we will obtain an MRI brain scan; a blood sample will also be collected
- You would be assigned to usual care or receive a telemetric blood pressure monitor to measure BP 3 times daily, and accordingly receive adjusted BP treatment to a target of 120mmHg systolic
- A 24-hour blood pressure monitoring at the beginning of the study, 3 months and 1 year

If you would like more information about taking part in the PROHIBIT-ICH study, please contact our research team

Research team contact name:

Research team contact telephone number: