You may 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
- A blood sample collected upon entering the study
- An MRI scan upon entering the study and at 1 year follow up
- 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: