

Stop Stroke: The South London Secondary Prevention Programme

Submission date 19/07/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 07/09/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 07/12/2010	Condition category Circulatory System	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Prof Charles Wolfe

Contact details
Division of Health & Social Care Research
King's College London
7th Floor Capital House
42 Weston Street
London
United Kingdom
SE1 3QD
+44 (0)20 7848 6608
charles.wolfe@kcl.ac.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
1824/1845

Study information

Scientific Title

Acronym

Stop Stroke

Study objectives

To test the efficacy of a complex secondary prevention intervention on improving post stroke risk factor control. The study uses a cluster randomised controlled trial design with 120 general practices randomised at the start of the trial. All participants at a given practice (patients, carers and primary care staff) receive either the intervention or usual care.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration.

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Prevention

Participant information sheet

Health condition(s) or problem(s) studied

Stroke

Interventions

Intervention: Computer algorithms are used to systematically transform data collected by an ongoing stroke register (SLSR) into individually tailored secondary prevention plans for patients, carers and the primary care team. The plans include details of the patient's current risk factor control and evidence based advice on how to improve risk factor management. Plans are updated and distributed at three time points.

Control: Usual care.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Prevalence of uncontrolled key modifiable risk factors for stroke (hypertension, smoking, aspirin adherence) measured at one year post stroke.

Secondary outcome measures

Prevalence of other uncontrolled risk factors for stroke (cholesterol, diabetes, atrial fibrillation, obesity, heavy drinking); stroke recurrence rate.

Overall study start date

21/07/2003

Completion date

01/01/2007

Eligibility

Key inclusion criteria

All patients with a diagnosis of first-in-a-lifetime stroke living in Lambeth and Southwark.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

450 patients

Key exclusion criteria

Patients must be alive six weeks post stroke, identified by the study team within six months of stroke and registered with a general practitioner.

Date of first enrolment

21/07/2003

Date of final enrolment

01/01/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Division of Health & Social Care Research
London
United Kingdom
SE1 3QD

Sponsor information

Organisation
King's College London (UK)

Sponsor details
Division of Health and Social Care Research
7th Floor Capital House
42 Weston Street
London
England
United Kingdom
SE1 3QD
+44 (0)207848 6651/4113
lesley.higgins@kcl.ac.uk

Sponsor type
University/education

Website
<http://www.kcl.ac.uk>

ROR
<https://ror.org/0220mzb33>

Funder(s)

Funder type
Not defined

Funder Name
The Health Foundation 1824/1845,

Funder Name

The Stroke Association TSA G1,

Funder Name

Medical Research Council Special Training Fellowship Award G106/1105, Department of Health Career Scientist Award PHCSA02/07.

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/11/2010		Yes	No