ISRCTN82122880 https://doi.org/10.1186/ISRCTN82122880

The Second Bradford Community Stroke Trial - A qualitative evaluation

Submission date	Recruitment status	[ ] Prospectively registered
23/01/2004	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
23/01/2004	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
23/10/2019	Circulatory System	[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
PCC115

Study information

Scientific Title
The Second Bradford Community Stroke Trial - A qualitative evaluation

Study objectives
The Second Bradford Community Stroke Trial seeks to assess the neglected area of psychosocial functioning in stroke patients and their families. Patients are randomly allocated to a control or intervention group. The latter group receive regular visits from one of the geriatric department liaison nurses who provide information, advice and support focusing on emotional and social recovery rather than physical function. Quantitative assessments are undertaken at 3, 6 and 12 months. However, there is a concern that the quantitative measures used may not be sufficiently comprehensive to capture the range of outcome experiences for the patient and carer. An additional qualitative evaluation would assess more fully the meanings and experience of their strokes for the patients in the two trial groups and thus facilitate:
1. A more detailed picture of the psycho-social difficulties experienced by stroke patients and their carers;
2. A better understanding of the nature of the intervention provided by the liaison nurses;
3. A more comprehensive evaluation of the trial outcome with a balance between quantitative and qualitative outcome assessments.

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)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Cardiovascular diseases: Cerebrovascular disease

Interventions

1. Regular visits from one of the geriatric department liaison nurses who provide information, advice and support focusing on emotional and social recovery rather than physical function
2. Standard care

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/07/1994

Completion date
09/06/1995

Eligibility

Key inclusion criteria
Patients undergoing trabeculectomy

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
01/07/1994

Date of final enrolment
09/06/1995

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Bradford Hospitals NHS Trust
Bradford
United Kingdom
BD5 0NA

Sponsor information

Organisation
NHS R&D Regional Programme Register - Department of Health (UK)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
NHS Executive Northern and Yorkshire (UK)

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