The Second Bradford Community Stroke Trial -A qualitative evaluation

Submission date	Recruitment status	Prospectively registered
23/01/2004 N	No longer recruiting	[_] Protocol
Registration date	Overall study status	[] Statistical analysis plan
23/01/2004	Completed Condition category	[_] Results
Last Edited		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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Contact details

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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

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