

A randomised controlled trial (RCT) to evaluate the cost-effectiveness and patient and carer satisfaction associated with different levels of intensity of community rehabilitation

Submission date 23/01/2004	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 23/01/2004	Overall study status Completed	☐ Statistical analysis plan ☐ Results
Last Edited 28/10/2016	Condition category Other	☐ Individual participant data ☐ 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

Protocol serial number

RBG 99XX7

Study information

Scientific Title

A randomised controlled trial (RCT) to evaluate the cost-effectiveness and patient and carer satisfaction associated with different levels of intensity of community rehabilitation

Study objectives

Evaluate and compare the levels of impairment, disability, handicap and well-being among elderly patients receiving either a supplemented CRT service or non-intensive CRT service following stroke or fractured neck of femur. Evaluate and compare the costs, both direct and indirect, of providing intensive community rehabilitation and non-intensive community rehabilitation for elderly patients following stroke or fractured neck of femur. Evaluate and compare levels of patient and informal carer satisfaction with intensive community rehabilitation and non-intensive community rehabilitation. Estimate the cost effectiveness of intensive community rehabilitation intervention versus non-intensive community rehabilitation for elderly patients following stroke or fractured neck of femur.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

Stroke or fractured neck of femur

Interventions

- i. Intensive community rehabilitation service
- ii. Standard care

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Death at 3 and 12 months. Institutionalisation at 3 and 12 months. Length of stay. Re-admission rates. Use of GP services. Use of other health services. Use of social services.

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/12/2002

Eligibility

Key inclusion criteria

Patients aged 65 plus, admitted from their own home to either the Central Sheffield University Hospital or Northern General Hospital following a 'first' stroke or fractured neck of femur.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/2000

Date of final enrolment

31/12/2002

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Centre for Ageing and Rehabilitation Studies

Sheffield

United Kingdom

S5 7AU

Sponsor information

Organisation

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

Funder(s)**Funder type**

Government

Funder Name

NHS Executive Trent (UK)

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration