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Fast-track assessment and rehabilitation for Stroke (FASTAR)

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	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
03/12/2008	Circulatory System	

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
RHC28102

Study information

Scientific Title

Acronym
FASTAR

Study objectives
To evaluate the costs and benefits for patients with stroke, their carers and the NHS of an acute assessment team and a home care rehabilitation team.

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. Provision of an acute assessment team and a home care rehabilitation team
2. Standard care

Intervention Type
Other

Phase
Not Specified

Primary outcome measure

1. All cause mortality
2. Functional Status
3. Length of hospital stay
4. Cause specific mortality
5. Morbidity
6. Time from admission to fit for discharge
7. Quality of life (patients and carers)

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/11/1999

Completion date
01/11/2002

Eligibility

Key inclusion criteria
Patients admitted to North Manchester General Hospital or Stepping Hill within 48 hrs of onset of a new clinical diagnosis of stroke

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/11/1999

Date of final enrolment
01/11/2002

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Manchester Health Authority
Manchester
United Kingdom
M60 9PL

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
Not defined

Funder Name
NHS Executive North West (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/07/2005		Yes	No