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The treatment of urinary incontinence in stroke patients

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
19/04/2012	Urological and Genital Diseases	[_] 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
MS9

Study information

Scientific Title

Study objectives
1. To determine the prevalence of urinary incontinence and other lower urinary tract symptoms (frequency, nocturia, urgency) in community-dwelling stroke survivors.
2. To assess the impact that urinary incontinence and other urinary symptoms have on the lives of community-dwelling stroke survivors.
3. To evaluate a new continence service centred on the Continence Nurse Practitioner working in the community and compare this with the existing General Practitioner led service in a randomised controlled trial.

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)
GP practice

Study type(s)
Treatment

Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied
Cardiovascular diseases: Cerebrovascular disease; Urological and genital diseases: Incontinence

Interventions
1. Continence service provided by a nurse practitioner
2. Existing general practice care (standard care)

Intervention Type
Other

Phase

Not Specified

Primary outcome measure
Prevalence
Self-reported presence of urinary incontinence and other lower urinary tract symptoms (frequency, nocturia, urgency) and impact.
Randomised controlled trial
Nursing assessment of urinary symptoms and a structured home interview carried out at baseline, 3 months and 6 months after entering the service.

Secondary outcome measures
Not provided at time of registration

Overall study start date
31/08/1997

Completion date
30/06/2000

Eligibility

Key inclusion criteria
Stroke survivors living in the community who reported experiencing urinary symptoms.

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
31/08/1997

Date of final enrolment
30/06/2000

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Division of Medicine for the Elderly
Leicester
United Kingdom
LE3 9QP

Sponsor information

Organisation
Record Provided by the NHS R&D 'Time-Limited' National Programme Register - Department of Health (UK)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
NHS Cardiovascular Disease and Stroke National Research and Development Programme (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?
Other publications	prevalence and management	01/10/1999		Yes	No