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A randomised trial to evaluate a standing frame to hasten recovery from acute stroke

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
09/12/2008	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
rctc98 Hudson

Study information

Scientific Title

Study objectives
The general research question is whether a more intensive early stroke physiotherapy intervention facilitated by the Oswestry standing frame will promote improved and/or faster stroke recovery.
Specifically, is the Oswestry standing frame associated with improved clinical outcome for patients in the acute phase after stroke and what are the resource implications of using the standing frame?

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: Stroke

Interventions
1. A minimum treatment schedule of one treatment session each week day for two weeks using the Oswestry standing frame (intervention group)
2. No use of the standing frame (control group)

In all other respects the rehabilitation care, both content and organisation, will be the same and delivered by the same ward staff.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Blind assessments will be completed by a research physiotherapist at baseline, ten weeks, and six months post stroke.

Secondary outcome measures
Not provided at time of registration

Overall study start date
09/01/1999

Completion date
10/01/2002

Eligibility

Key inclusion criteria
Patients from the Bradford Stroke Unit with a new hemiplegic stroke who are medically stable and alert but who need support from two or more rehabilitation staff to achieve standing will be approached for recruitment to the trial.

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
09/01/1999

Date of final enrolment
10/01/2002

Locations

Countries of recruitment
United Kingdom

Study participating centre
St Luke's Hospital
Bradford
United Kingdom
BD5 ONA

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/2005		Yes	No