ISRCTN66554171 https://doi.org/10.1186/ISRCTN66554171

# Pilot RCT to assess the impact of additional supported standing practice on functional ability post stroke

| | | |
|---|---|---|
| **Submission date**<br>29/09/2006 | **Recruitment status**<br>No longer recruiting | [_] Prospectively registered<br>[_] Protocol<br>[_] Statistical analysis plan<br>[X] Results<br>[_] Individual participant data |
| **Registration date**<br>29/09/2006 | **Overall study status**<br>Completed | |
| **Last Edited**<br>14/11/2022 | **Condition category**<br>Circulatory System | |

**Plain English summary of protocol**
Not provided at time of registration

## Contact information

**Type(s)**
Scientific

**Contact name**
Mrs Rhoda Allison

**Contact details**
Teignbridge PCT
Newton Abbot Hospital
East Street
Newton Abbot
United Kingdom
TQ12 4PT
+44 01626 357201
rhoda.allison@nhs.net

## Additional identifiers

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**

N0620164288

# Study information

### Scientific Title
Pilot RCT to assess the impact of additional supported standing practice on functional ability post stroke

### Study objectives
To assess the effect of providing extra sessions of standing practice on functional recovery after stroke.

### Ethics approval required
Old ethics approval format

### Ethics approval(s)
Not provided at time of registration

### Study design
Pilot randomised controlled trial

### Primary study design
Interventional

### Secondary study design
Randomised controlled trial

### Study setting(s)
Hospital

### Study type(s)
Treatment

### Participant information sheet

### Health condition(s) or problem(s) studied
Cardiovascular: Stroke

### Interventions
People who fill the criteria for inclusion will be invited to participate and after giving informed consent will be randomly allocated to one of the two treatment groups. The pilot will admit a total of twenty participants who will be randomly allocated to either the group for conventional physiotherapy or the group with conventional physiotherapy plus additional standing practice.

### Intervention Type
Other

### Phase
Not Specified

### Primary outcome measure
A battery of outcome measures will be recorded throughout the study. These are measures when the participant is asked to perform a number of tasks such as rolling over in bed, standing up, etc. The outcome measurement typically takes about 30-45 minutes to complete. Outcome measures will be taken on a weekly basis during the inpatient stay, and then at 12 weeks after the initial assessment.

### Secondary outcome measures
No secondary outcome measures

### Overall study start date
21/02/2005

### Completion date
30/04/2006

## Eligibility

### Key inclusion criteria
1. People who are received rehabilitation on the Stroke Rehabilitation Ward, Newton Abbot Hospital
2. Confirmed diagnosis of stroke
3. People who are able to participate in a rehabilitation programme (of 45 minutes of physiotherapy each working day)

### Participant type(s)
Patient

### Age group
Adult

### Sex
Not Specified

### Target number of participants
20 participants

### Total final enrolment
17

### Key exclusion criteria
The ward in question will not admit people who are unconscious or who have a diagnosis of mental health illness.

### Date of first enrolment
21/02/2005

### Date of final enrolment
30/04/2006

# Locations

### Countries of recruitment
England

United Kingdom

### Study participating centre
**Teignbridge PCT**
Newton Abbot
United Kingdom
TQ12 4PT

# Sponsor information

### Organisation
Record Provided by the NHSTCT Register - 2006 Update - Department of Health

### 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.dh.gov.uk/Home/fs/en

# Funder(s)

### Funder type
Government

### Funder Name
Exeter Primary Care Trust (UK) - Own account funding

**Funder Name**
NHS R&D Support Funding (UK)

# Results and Publications

## Publication and dissemination plan
Not provided at time of registration

## Intention to publish date

## Individual participant data (IPD) sharing plan
Not provided at time of registration

## 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 | | 01/07/2007 | | Yes | No |