# A Preliminary Investigation Into The Effect Of Home Standing Programs On Lower Limb Spasticity, Spasm, Pain and Well-being For Individuals With Progressive Multiple Sclerosis (MS)

Submission date	Recruitment status	Prospectively registered
30/09/2005	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2005	Completed	Results
Last Edited	Condition category	Individual participant data
12/09/2013	Nervous System Diseases	<ul><li>Record updated in last year</li></ul>

## Plain English summary of protocol

Not provided at time of registration

## Contact information

## Type(s)

Scientific

#### Contact name

Miss Karen Baker

#### Contact details

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## Additional identifiers

EudraCT/CTIS number

IRAS number

## ClinicalTrials.gov number

## Secondary identifying numbers

N0515143146

## Study information

#### Scientific Title

## **Study objectives**

The principal research questions are that daily standing in a frame will:

- 1. Decrease passive resistance to movement in lower limb (lower limb stiffness know as hyper tonicity)
- 2. Enhance range of motion at ankles.
- 3. Decrease frequency of lower limb spasms.
- 4. Lead to self-reported decrease in pain.
- 5. Lead to self-reported improvement in well-being.

## 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

Nervous System Diseases: Multiple sclerosis (MS)

#### **Interventions**

The study proposed to use a crossover design of a minimum of six participants with primary or secondary MS these participants will be randomly selected to either a three-week exercise group or a three week standing group. The participants in the standing group will stand daily in a Standing Frame for 30 minutes if able. These participants will be shown how to use and stand in

the frame, given exercises to perform whilst standing and show how to correctly position themselves in the frame. Preliminary research results will be feedback to the participants at the end of the six week study period. They will also be informed that final results will be available from the researchers, and the local MS society will also be made aware of the results. Measurements will be taken at assessment, three and six weeks, with the groups crossing over after three weeks.

## Intervention Type

Other

#### Phase

**Not Specified** 

## Primary outcome measure

Not provided at time of registration

#### Secondary outcome measures

Not provided at time of registration

## Overall study start date

01/05/2004

## Completion date

01/05/2005

## **Eligibility**

### Key inclusion criteria

Participants are to be identified from clinical team members (physiotherapists) working in either Harrow PCT, Northwick Park Hospital or Harrow MS Therapy Centre.

### Participant type(s)

Patient

#### Age group

**Not Specified** 

#### Sex

**Not Specified** 

### Target number of participants

6

### Key exclusion criteria

Does not match inclusion criteria

#### Date of first enrolment

01/05/2004

#### Date of final enrolment

## Locations

#### Countries of recruitment

England

**United Kingdom** 

Study participating centre
Physical Disability Support Team
Harrow
United Kingdom
HA1 3UJ

## Sponsor information

## Organisation

Department of Health

## Sponsor details

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**

North West London Hospitals NHS Trust (UK) - NHS R&D Support Funding

## **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