

Functional strength training to improve walking and upper limb function in stroke patients

Submission date 12/01/2009	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 30/01/2009	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 15/08/2014	Condition category Circulatory System	<input type="checkbox"/> Individual participant data

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
TSA 2008/08

Study information

Scientific Title

Functional strength training to improve walking and upper limb function in people at least one year after stroke: a phase II trial

Study objectives

The hypothesis is that providing functional strength training (FST) to people who are at least one year after stroke will improve motor function and ability to perform everyday functional activity. The first step towards testing this hypothesis in a definitive phase III clinical trial is to address the following questions:

1. Is there sufficient efficacy to justify subsequent trials of FST for upper and lower limb motor recovery in people at least one year after stroke?
2. Is FST delivered in the community acceptable to stroke survivors one year and more after stroke?
3. What is the probable recruitment rate to a subsequent phase III trial?
4. What sample size is needed for subsequent phase III trial (effect size, attrition rate, response variation)?
5. What cost-effectiveness data should be collected in subsequent trials?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Norfolk Ethics Committee approval pending as of 12/01/2009

Study design

Randomised placebo-controlled observer-blind phase II trial with embedded qualitative investigation

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

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

Stroke

Interventions

Functional strength training is hands-off progressive resistive exercise during functional activity whilst directing participants attention to the activity being performed and providing appropriate verbal feedback to enhance knowledge of performance. FST is designed to increase ability to

produce voluntary muscle force throughout joint range, increase ability to modulate force in muscles/muscle groups appropriate for the activity being trained and improve functional ability. Activities are progressed by increasing the number of repetitions, increasing range of joint motion required and increasing the load to be moved. Intervention will be provided by a Research Physiotherapist for 1 hour/day, 4 days/week. Portable equipment (e.g. free weights and steppers) will be used as appropriate. Participants will be encouraged to use the paretic limb (upper or lower as allocated) in everyday functional activity. Standardised treatment schedules will be used (as in our earlier trials) to record the amount and type of intervention provided (compliance).

FSTLL will focus on functional activities e.g. sit-to-stand using body weight as resistance and progressively lowering the height of the supporting surface as participants improve. Equipment such as portable steppers will be used as appropriate but only with physiotherapy supervision for safety reasons.

FSTUL will focus on reaching for, retrieval of and manipulation (use) of everyday objects for activities such as pouring water from a jug, opening jars and lacing a shoe.

Both treatment schedules will include resistive exercise to strengthen specific muscles/muscle groups where weakness is hampering functional activity. The scientific papers describing these treatment schedules are currently in preparation.

Intervention Type

Other

Phase

Phase II

Primary outcome measure

1. Functional Ambulatory Category for lower limb function
2. Action Research Arm Test for upper limb function

All measured on days 43 and 88.

Secondary outcome measures

1. Modified Rivermead Mobility Index for lower limb function
2. Nine Hole Peg Test for upper limb function

All measured on days 43 and 88.

Overall study start date

01/03/2009

Completion date

01/03/2012

Eligibility

Key inclusion criteria

1. Adults aged 18 years and above, either sex, one to five years after stroke in anterior circulation (infarct or haemorrhage) not receiving formal therapy for their upper or lower limb

2. Have a walking ability on the Functional Ambulatory Category between 1 and 49 and have some voluntary activity in the paretic upper limb scoring 12+/57 on the Action Research Arm Test but unable to complete the Nine Hole Peg Test in 50 seconds or less
3. Can follow a one-stage command, i.e. sufficient communication/orientation for interventions in this trial

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

58

Key exclusion criteria

Known pathology precluding them participating in functional strength training

Date of first enrolment

01/03/2009

Date of final enrolment

01/03/2012

Locations**Countries of recruitment**

England

United Kingdom

Study participating centre

School AHP

Norwich

United Kingdom

NR4 7TJ

Sponsor information

Organisation

The Stroke Association (UK)

Sponsor details

Stroke House
240 City Road
London
United Kingdom
EC1V 2PR
research@stroke.org.uk

Sponsor type

Charity

Website

<http://www.stroke.org.uk/>

ROR

<https://ror.org/05mgfq941>

Funder(s)

Funder type

Charity

Funder Name

The Stroke Association (UK) (ref: TSA 2008/08)

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?
Protocol article	protocol	01/07/2013		Yes	No

[Results article](#)

results

12/08/2014

Yes

No