

A Pilot Study to Investigate if Frequency of Therapy Alters Outcomes of the Lee Silverman Voice Treatment (LSVT) Programme for Patients with Parkinson's Disease

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 07/09/2015	Condition category Nervous System Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0016131978

Study information

Scientific Title

A Pilot Study to Investigate if Frequency of Therapy Alters Outcomes of the Lee Silverman Voice Treatment (LSVT) Programme for Patients with Parkinson's Disease

Study objectives

A Pilot Study to Investigate if Frequency of Therapy Alters Outcomes of the Lee Silverman Voice Treatment (LSVT) Programme for Patients with Parkinson's Disease

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)

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

Nervous System Diseases: Parkinson's disease

Interventions

Not provided at time of registration

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Publication

Secondary outcome measures

Not provided at time of registration

Overall study start date

08/08/2003

Completion date

10/01/2005

Eligibility**Key inclusion criteria**

1. Patients entering the study will have volunteered for the study and will have agreed to be randomly allocated to either a control group or one of two treatment groups.
2. A diagnosis of idiopathic PD
3. Categorised between stages 1 to 4 on the 'Functional Rating Scale for Parkinson's Disease', Hoehn & Yahr (1967)
4. Neuropharmacologically stable before and during treatment
5. Fluent English speakers
6. Intelligibility of speech will have received a rating of between 5 and 9 on the Parkinson's disease disability rating scale. (Yorkston 1995)

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

27

Key exclusion criteria

1. Any history of a language deficit caused by a past event such as a cardiovascular event
2. Evidence of a cognitive deficit on the 'Mini-Mental State Examination (score >24), Folstein et al 1975
3. Evidence of depression on the Beck Depression inventory (score <14), Beck & Beck 1972.
4. History of a chronic hearing loss or report/suspected hearing loss
5. History of laryngeal disease and will be free of acute cardiac and respiratory conditions during the study

Date of first enrolment

08/08/2003

Date of final enrolment

10/01/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Hammersmith Hospital

London

United Kingdom

W12 0HS

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House

79 Whitehall

London

United Kingdom

SW1A 2NL

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Hammersmith Hospital NHS Trust (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