

The effect of the implantable two-channel peroneal nerve stimulator as a treatment in stroke patients with a drop foot in comparison with the conventional treatment

Submission date 27/01/2006	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 27/01/2006	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 04/07/2019	Condition category Circulatory System	☐ Individual participant data

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

Protocol serial number

001; NTR494

Study information

Scientific Title

The effect of the implantable two-channel peroneal nerve stimulator as a treatment in stroke patients with a drop foot in comparison with the conventional treatment

Acronym

RCT PNS (peroneal nerve stimulation)

Study objectives

The functional electrical stimulation (FES) group will show in comparison with the conventional therapy group:

1. Increased gait speed (primary outcome)
2. Increased endurance
3. Improved gait kinematics
4. Increased muscle activity level
5. Reduced spasticity
6. Positive effect on passive range of movement (ROM)
7. Reduced disability

Ethics approval required

Old ethics approval format

Ethics approval(s)

Received from the local medical ethics committee

Primary study design

Interventional

Study design

Randomised, active controlled, parallel group trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Dropfoot, stroke

Interventions

The conventional management of dropped foot has been to use a rigid orthosis to maintain the ankle in a neutral position. This has major limitations as a treatment, being both uncomfortable and awkward to use and hence is often rejected by patients and therapists.

Currently, functional electrical stimulation (FES) systems for the treatment of dropped foot are in clinical use in significant numbers. FES is the artificial stimulation of muscles with the purpose of evoking a motor response. Compared with the use of orthosis, electrical stimulation has a number of advantages: it prevents muscle atrophy, the blood flow remains normal or even improves and it is cosmetically better accepted.

An implantable system was developed that stimulates the two branches of the peroneal nerve separately. Results from previous studies indicate that the system is safe to use, well liked by the patients, provides selectivity over moments at the ankle joint and increases both walking

speed and endurance. In the present study the additional value of the two-channel implantable peroneal nerve stimulator in comparison with the conventional treatment will be examined by measuring different parameters.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Walking speed

Key secondary outcome(s)

1. Endurance
2. Spasticity
3. EMG
4. 3D-kinematics
5. Quality of life questionnaires
6. Activity monitoring
7. Carry-over effect

Completion date

31/12/2005

Eligibility**Key inclusion criteria**

1. Dropped foot identified by an inability to achieve a normal heel strike during walking
2. First hemiplegia of at least 6 months as a result of a cerebrovascular accident (CVA) with a stable neurology
3. Successful functional recovery after surface stimulation of the common peroneal nerve
4. Subject is an outdoor walker
5. Able to give informed consent

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Total final enrolment

29

Key exclusion criteria

1. Aged less than 18 years
2. Passive dorsiflexion of the ankle less than 5° with knee in extension
3. Medical conditions limiting the function of walking other than CVA, i.e. neurological, rheumatic, cardio-vascular or systemic disorders (including Diabetes Mellitus)
4. Injury of N. peroneus or N. ischiadicus
5. Not be able to don and doff the equipment
6. Pregnancy

Date of first enrolment

01/09/2002

Date of final enrolment

31/12/2005

Locations**Countries of recruitment**

Netherlands

Study participating centre

Roessingh Research and Development

Enschede

Netherlands

7500 AH

Sponsor information**Organisation**

Roessingh Research and Development B.V. (The Netherlands)

ROR

<https://ror.org/01dmjt679>

Funder(s)**Funder type**

Government

Funder Name

SETER - A branch of the Dutch Ministry of Economic Affairs (The Netherlands)

Results and Publications

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	01/08/2007	04/07/2019	Yes	No