

Implicit motor learning in gait training after stroke

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Ethische beoordeling	Positief advies
Status	Anders
Type aandoening	-
Onderzoekstype	Interventie onderzoek

Samenvatting

ID

NL-OMON20141

Bron

NTR

Aandoening

The study is targeted at people after stroke who are in the chronic stage of recovery.

Stroke, CVA

Ondersteuning

Primaire sponsor: This work was supported by Nationaal Regieorgaan Praktijkgericht Onderzoek SIA (RAAKPRO; grant number 2014-01-49PRO).

Overige ondersteuning: <http://www.regieorgaan-sia.nl/contact>

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

The main outcome parameter in work package A is walking speed measured using the 10-Meter Walk Test

The central outcome parameter in work package B is the technology acceptance of the Stappy-system

Toelichting onderzoek

Achtergrond van het onderzoek

Objective: The objective of this study is twofold. The first objective is to study the effectiveness of implicit motor learning on walking speed in clients after stroke in daily practice. The second objective is to determine the technology acceptance, feasibility, and first effects on walking performance of the Stappy-system.

Study design: This research protocol involves two work packages A (objective 1) and B (objective 2). In work package A, a randomized, controlled, single-blinded study design will be adopted. Work package A will be followed by work package B in which a prospective process evaluation of a technology intervention will take place.

Study population: The same participants will take place in both work packages A and B. People will be eligible for participation if they have had a stroke (>6 months after stroke), would like to improve their walking performance, have a slower walking speed than (10 m/s), can communicate in Dutch language (enough to understand the instructions), and can follow a 3-stage command. People will be excluded for participation if they cannot walk a minimum of 10 meters, or have other additional impairments that may influence their gait.

Intervention: Work package A consists of a 3-week intervention period that includes 3 training sessions per week. The experimental group receives gait training based on implicit learning principles and the control group will receive gait training based on explicit learning principles. In work package B, the participants will receive an easy-to-use technology (Stappy-system) to support their gait exercises independently at home.

Main study parameters/endpoints

Demographic data: Age, gender, Length, time post stroke, side of stroke, walking aids, educational level, Montreal Cognitive Assessment, Berg Balance Scale, Rivermead Mobility Index, Fugl-Meyer Assessment, Movement Specific Reinvestment Scale

Work package A: The main outcome parameter is walking speed measured using the 10-Meter Walk Test. Secondary outcome measures include: quality of gait (modified Dynamic Gait Index), dual-task performance, the participants' satisfaction (Global Perceived Effect), the propensity for conscious processing (Movement Specific Reinvestment Scale) and quality of life (Stroke and Aphasia Quality of Life Scale-39).

Work package B: The main outcome parameter for work package B is the technology acceptance of the Stappy-system. Secondary outcomes are based on the feasibility and potential first effects on walking performance.

Doel van het onderzoek

Work package A

The aim of work package A is to examine the effectiveness of analogy learning (based on implicit motor learning principles) compared to explicit motor learning on walking performance in people after stroke.

Work package B

The primary aim of work package B is to evaluate the acceptability of the Stappy-system for independent gait training in daily life in people after stroke. Secondary aims are to assess the feasibility and potential first effects on walking performance of the Stappy-system in people

Onderzoeksopzet

T0: 0-weeks (baseline)

T1: 4-weeks

T2: 8-weeks

Onderzoeksproduct en/of interventie

Depending on randomisation, participants will receive a gait training based on implicit motor learning principles (experimental group) or based on explicit motor learning principles (control group).

Contactpersonen

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Wetenschappelijk

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

In order to be eligible to participate in this study, a subject must meet all of the following criteria. The person

- has had a stroke and is in the chronic stage of recovery (> 6 months after stroke)
- has a walking speed slower than 1.0 m/s (there is limited space for improvement if walking speed would be too high (ceiling effect))
- is able to communicate in Dutch language, at least to understand the verbal instructions of the physiotherapist
- has the ability to complete a 3-stage command

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

A potential subject who meets any of the following criteria will be excluded from participation in this study. The person

- cannot walk a minimal distance of 10 meters (if necessary with manual assistance or walking aid)
- has additional diagnosed impairments, not related to stroke, that can influence the gait pattern e.g. severe osteoarthritis or amputation of the lower limb
- could not ambulate on level surfaces without manual contact of another person (Functional Ambulation Scale (FAC) < 3)

Onderzoeksopzet

Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Parallel
Toewijzing:	Gerandomiseerd
Controle:	Geneesmiddel

Deelname

Nederland	
Status:	Anders
(Verwachte) startdatum:	01-05-2017
Aantal proefpersonen:	80
Type:	Onbekend

Voornemen beschikbaar stellen Individuele Patiënten Data (IPD)

Wordt de data na het onderzoek gedeeld: Nog niet bepaald

Ethische beoordeling

Positief advies	
Datum:	10-03-2017
Soort:	Eerste indiening

Registraties

Opgevolgd door onderstaande (mogelijk meer actuele) registratie

Geen registraties gevonden.

Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

In overige registers

Register	ID
NTR-new	NL6133
NTR-old	NTR6272
CCMO	NL.60338.096.16

Resultaten