

Kinematic Motion Analyses of Upper Extremity After Intensive Training with Spring-assisted Dynamic Hand Orthosis in Patient with Stroke

Submission date 11/10/2010	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 19/10/2010	Overall study status Completed	☐ Statistical analysis plan ☐ Results
Last Edited 19/10/2010	Condition category Circulatory System	☐ Individual participant data ☐ Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
2009-44

Study information

Scientific Title

Effects of Spring Assisted Dynamic Hand Orthosis Training on Functions and Movement Smoothness of the Hemiparetic Upper Extremity: A randomised controlled trial

Acronym

SADHO

Study objectives

The purpose of this experiment was to evaluate the effectiveness of training using a spring assisted dynamic hand orthosis on smoothness of movement, clinical assessment score, and grip strength of the affected limb in hemiparetic patients. To determine resultant velocity and jerkiness score for movement smoothness, the Box and Block Test (BBT), Action Research Arm Test (ARAT), and Fugl-Meyer Assessment (FMA) for functions of the hemiparetic upper extremity were conducted.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Yonsei University, College of Medicine approved on the 25th of March 2010 (ref: 2009-44)

Primary study design

Interventional

Study design

Randomised comparative interventional pretest-posttest control group design

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Hemiparesis due to Stroke

Interventions

Patients will undergo training activities while each wore a spring assisted dynamic hand orthosis. Training will be guided by a physiotherapist for 1 hour per day per week for 4 weeks.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

3 clinical assessments and the measurement of grip strength. Fugl-Meyer Assessment and grip strength and Action Research Arm Test and Box and Block Test.

All outcomes were measured at baseline and after 4 weeks of training.

Key secondary outcome(s)

Spatiotemporal parameters were collected using a 3-D motion analysis system and workstation software pre- and post-test (VICON MX system, Oxford Metrics, U.K.). Data collection was

conducted at the Motion Analysis Research Laboratory in the Yonsei University.
All outcomes were measured at baseline and after 4 weeks of training.

Completion date

15/03/2010

Eligibility

Key inclusion criteria

1. 18 years or older
2. Unilateral hemiparesis more than 6 months post-stroke duration
3. No current or previous orthopedic or surgical histories affecting the hemiparetic upper extremity
4. Mini-Mental State Examination (MMSE) Korean version - score ≥ 23 (to ensure that they fully understood the study procedure)
5. Patients should have at least some active voluntary movement of the upper extremity (i.e., 10 degrees of shoulder flexion/abduction, 10 degrees of elbow flexion/extension, and 30 degrees of interphalangeal proximal joints / 20 degrees of interphalangeal distal joints of volitional finger flexion when the hand is positioned in wrist and finger extension)
6. No flaccidity of the affected limb
7. No severe contracture or spasticity of the affected wrist or hand

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 Years

Sex

All

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

23/12/2009

Date of final enrolment

15/03/2010

Locations

Countries of recruitment

Korea, South

Study participating centre
Department of Physical Therapy
Gyeongsangbuk-do
Korea, South
760-709

Sponsor information

Organisation
Yonsei University College of Medicine (South Korea)

ROR
<https://ror.org/01wjejq96>

Funder(s)

Funder type
Other

Funder Name
Investigator funded (South Korea) - Dissertation project

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration