

Attentional demand of walking in post-polio patients: the association with incidence of falls

Published: 01-06-2006

Last updated: 14-05-2024

The aims of the study are fourfold: 1) to assess dual-task walking performance in patients with late sequelae of polio and compare this to dual-task walking performance of age-matched healthy controls; 2) to investigate the reproducibility of a dual...

Ethical review	Approved WMO
Status	Pending
Health condition type	Muscle disorders
Study type	Observational non invasive

Summary

ID

NL-OMON29899

Source

ToetsingOnline

Brief title

Dual-task walking in PPS

Condition

- Muscle disorders
- Neuromuscular disorders

Synonym

late effects of poliomyelitis, postpolio syndrome

Research involving

Human

Sponsors and support

Primary sponsor: Academisch Medisch Centrum

Source(s) of monetary or material Support: Ministerie van OC&W

Intervention

Keyword: Poliomyelitis, Postpoliomyelitis Syndrome, Walking

Outcome measures

Primary outcome

Walking performance, assessed with a 2-min walking tests and a dual-task walking test.

The 2-min indoor walking test will be performed on a closed marked indoor trajectory. The indoor oval track has a length of 50 m. Subjects are instructed to walk at a self-preferred walking speed for 2 minutes. The walking distance will be measured.

2) the dual task walking test consists of a 50m walking test combined with a secondary cognitive task. In this test subjects are instructed to walk, on a marked trajectory, a distance of 50m twice at their own speed with their normal walking aid (if needed) without stopping. The first 50m walk will be performed without a secondary task. During the second 50 m walk, subjects will solve orally presented arithmetic problems while walking. Trial duration of each trial will be recorded. The total time of the 2 trials, the difference between the 2 trials and walking velocity at the first 50m walk will be analyzed.

The incidence of falls will be based on the subject*s self report. A questionnaire will be administered, addressing demographic data, frequency of tripping, frequency and description of falls over the last 12 months, their consequences, and community services utilized.

Secondary outcome

Quadriceps muscle strength will be measured with a Biodex dynamometer.

Study description

Background summary

In older subjects, weakness of the lower limbs increases the risk of falls, particularly in situations in which the whole body weight is placed on 1 leg. Persons previously affected by poliomyelitis (polio subjects) commonly have long-standing muscle weakness, often in the lower limbs. These patients are often unstable when standing and walking, and they report frequent falls (Cosgrove et al 1987; Silver and Aiello 2002, Lord et al. 2002). In frail elderly subjects and patients with Parkinson, dual-task walking tests have been used to identify individuals at-risk for falling. It is not known whether dual-task walking tests can predict falls in polio subjects.

Study objective

The aims of the study are fourfold: 1) to assess dual-task walking performance in patients with late sequelae of polio and compare this to dual-task walking performance of age-matched healthy controls; 2) to investigate the reproducibility of a dual-task walking test
3) to determine the frequency of falls over the last 12 months, circumstances surrounding them, and the consequences of falls and 4) to investigate the association between dual-task walking performance and incidence of falls in patients with late sequelae of polio.

Study design

Study design: a cross-sectional study.

Study burden and risks

Patients with late effects of poliomyelitis will visit the department of Rehabilitation AMC twice. During the first visit they will perform the 2 walking tests. Measurement of quadriceps strength will be performed. The questionnaire on fall events will be administered. During the second visit, the walking tests will be repeated.

No risks are associated with performing the walking tests at a self-preferred, comfortable walking speed. Sufficient rest will be provided between the walking tests.

Healthy control subjects will visit the department of Rehabilitation once.
During this visit they will perform the 2 walking tests and quadriceps strength will be measured.

Contacts

Public

Academisch Medisch Centrum

Postbus 22660
1100 DD Amsterdam
Nederland

Scientific

Academisch Medisch Centrum

Postbus 22660
1100 DD Amsterdam
Nederland

Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years)

Elderly (65 years and older)

Inclusion criteria

Patients: 1) diagnosis of poliomyelitis; 2) ability to walk 50 m with or without walking aid.
Controls: good health

Exclusion criteria

1) impaired vision; 2) hearing loss.

Study design

Design

Study type:	Observational non invasive
Intervention model:	Other
Allocation:	Non-randomized controlled trial
Masking:	Open (masking not used)
Control:	Active
Primary purpose:	Prevention

Recruitment

NL	
Recruitment status:	Pending
Start date (anticipated):	15-04-2006
Enrollment:	60
Type:	Anticipated

Ethics review

Approved WMO	
Application type:	First submission
Review commission:	METC Amsterdam UMC

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
CCMO	NL11978.018.06