Prospective comparison in new challenging gait and balance tasks in female adolescents with idiopathic scoliosis before and following corrective spinal fusion

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The aim of the present study is to use a more complete analysis of spinal motion (using a 3D multi-segment spine model) and to introduce challenging gait tasks which can optimally reveal the compensation strategies of the spine. Furthermore, static...

Ethical review	Approved WMO
Status	Recruitment stopped
Health condition type	Bone disorders (excl congenital and fractures)
Study type	Observational non invasive

Summary

ID

NL-OMON34137

Source ToetsingOnline

Brief title Scoliosis_gait

Condition

- Bone disorders (excl congenital and fractures)
- Bone and joint therapeutic procedures

Synonym

Adolescent Idiopathic scoliosis, spinal deformity

Research involving

Human

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Sponsors and support

Primary sponsor: Sint Maartenskliniek Source(s) of monetary or material Support: SMK Research BV

Intervention

Keyword: Gait analysis, Postural stability, Scoliosis, Surgical correction

Outcome measures

Primary outcome

In the static balance tasks the primary parameter is the vCOP and the RMS of

the COP. In the challenging gait tasks the main parameter is the axial rotation

of the different spinal segments in relation to each other and in relation to

the pelvis.

Secondary outcome

General information like gender, age, length, weight and menarche will be

collected. The Cobb angle will be determined on X-rays. Furthermore, walking

speed, stride length and joint angles will be calculated. Functional scores

will be related to the Cobb angle.

Study description

Background summary

Patients with disturbed trunk symmetry, such as patients with adolescent idiopathic scoliosis (AIS), have problems with stability control during standing and walking. Often these patients require spinal fusion surgery, which reduces the spinal mobility and affects the trunk biomechanics. So far, one lacks appropriate quantitative measures to assess the functional outcome of these interventions. At best, a simple gait analysis is performed using standard methods (lower limb kinematics and the global spinal kinematics such as acromion-pelvis (C7-S2) angle during simple level walking before and after surgery). Nevertheless, it is essential to study the (functional) changes induced by the surgery, since these are apparently tolerated very well. In addition, one needs to learn how individual patients compensate for the fusion. It is most likely that adjacent vertebrae have to compensate for the loss in spinal mobility in AIS patients. Data of this type are clinically important since this allows us to study the biomechanically important effects of the number of fused vertebrae and the level of the fusion (thoracic or thoracolumbar). Moreover, this will help clinicians in predicting and avoiding additional problems that result from spinal fusion (e.g. adjacent segment degeneration, junctional kyphosis, *adding-on*).

Study objective

The aim of the present study is to use a more complete analysis of spinal motion (using a 3D multi-segment spine model) and to introduce challenging gait tasks which can optimally reveal the compensation strategies of the spine. Furthermore, static balance will be investigated.

Study design

Patients in this experimental study will be tested before and after surgery in demanding balance and gait tasks such as fast walking and walking uphill to test their functional potential and to highlight compensatory strategies. The measurements are non-invasive and also includes a questionnaire (SRS-22).

Study burden and risks

Patients will not be burden with excessive risks in case of participation, nor with benefits. The additional measurements in the gait laboratory will be performed, which will take some extra time. Patients are wearing a safety-belt during walking on the treadmill. Furthermore, no additional visits to the Sint Maartenskliniek will be necessary.

Participation of the patients is completely voluntary and patients have to sign the informed consent procedure. Patients are allowed to withdraw the study at any moment without reason. The privacy of the patients is protected with the following measures: case report forms are filed under code number; all involved investigators have professional responsibility not to compromise the confidentiality of patient information. This is also stated in the patient information. In reports, the patients will be presented anonymously.

Contacts

Public Sint Maartenskliniek Hengstdal 3 6522 JV Nijmegen NL Scientific Sint Maartenskliniek

Hengstdal 3 6522 JV Nijmegen NL

Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adolescents (12-15 years) Adolescents (16-17 years) Adults (18-64 years) Elderly (65 years and older)

Inclusion criteria

-Adolescent Idiopathic Scoliosis -age 12-18 years -on waiting list for surgical correction

Exclusion criteria

-not being barred of any previous spinal surgery -neurological or musculoskeletal disorders -mental retardation

Study design

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Design

Study type: Observational non invasive		
Masking:	Open (masking not used)	
Control:	Uncontrolled	
Primary purpose:	Treatment	

Recruitment

КП

NL	
Recruitment status:	Recruitment stopped
Start date (anticipated):	03-02-2011
Enrollment:	25
Туре:	Actual

Ethics review

Approved WMO	
Date:	02-03-2011
Application type:	First submission
Review commission:	IRB Nijmegen: Independent Review Board Nijmegen (Wijchen)

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 CCMO

ID NL35194.072.10

Study results

Date completed:	21-09-2013
Actual enrolment:	25