A new method to objectively observe the function of orthopaedic patients

Submission date	Recruitment status	Prospectively registered
30/05/2007	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/05/2007	Completed	Results
Last Edited	Condition category	Individual participant data
25/10/2021	Musculoskeletal Diseases	Record updated in last year

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 NL942 (NTR967)

Study information

Scientific Title

A new method to objectively observe the function of orthopaedic patients

Study objectives

The use of an objective device (accelerometer) to analyse function is more accurate than the currently used methods (observing by eyes).

Ethics approval required

Old ethics approval format

Ethics approval(s)

Received from the local ethics committee (METC Atrium MC - Maaslandziekenhuis). The committee concluded on the 26th March 2007 that the research is not WMO- obligated and permission was granted.

Study design

Observational study

Primary study design

Observational

Study type(s)

Diagnostic

Health condition(s) or problem(s) studied

Orthopaedic patients, accelerometer

Interventions

An accelerometer, attached to the sacrum of the subject using an elastic belt, measures accelerations in three directions when the doctor observes the patient as usual (walk ability, sit to stand, balancing, etc). The objective outcome will be compared with the doctors observation and questionnaires.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Parameters which give information about the function of the patient like walk speed, asymmetry during walking, postural sway during balancing etc. These will be measured at the end of the doctors investigation.

Key secondary outcome(s))

Patient data like weight, height, date of birth and the health state of the patient is collected. These will be measured at the end of the doctors investigation.

Completion date

01/05/2008

Eligibility

Key inclusion criteria

All orthopaedic patients visiting the policlinic of the orthopaedic department of the Atrium Medical Centre Heerlen.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Unable to perform the tasks (walking, balacing, rising from a chair, step on a block).

Date of first enrolment

15/05/2007

Date of final enrolment

01/05/2008

Locations

Countries of recruitment

Belgium

Netherlands

Study participating centre Hoogstraat 229

Voeren

Belgium

B-3798

Sponsor information

Organisation

Atrium Heerlen Orthopaedic Research and Scientific Education (AHORSE) (The Netherlands)

ROR

https://ror.org/0367sye10

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Atrium Medical Centre (The Netherlands)

Results and Publications

Individual participant data (IPD) sharing plan

Not provided at time of registration

IPD sharing plan summary

Not provided at time of registration