Dutch validation of Hip Outcome Score in FAI

No registrations found.

Ethical review Positive opinion **Status** Recruiting

Health condition type -

Study type Observational non invasive

Summary

ID

NL-OMON26525

Source

Nationaal Trial Register

Brief title

HOS validation

Health condition

Dutch Validation Hip Outcome Score

Sponsors and support

Primary sponsor: Vakgroep orthopedie, Reinier de Graaf Gasthuis, Delft the Netherlands

Source(s) of monetary or material Support: Initiator = sponsor

Intervention

Outcome measures

Primary outcome

Reliability, construct validity and content validity: high correlation of HOS-NL with mHHS, HAGOS and iHOT-12-NL, no floor-and/or ceiling effects, a good sensitivity to cahnge and a good test-retest reliability with a high intraclass correlation coefficient.

Secondary outcome

Minimal clinical important difference after 6 months, minimal detectable change and responsiveness after 6 months.

Study description

Background summary

Validations of the Dutch translation of the Hip Outcome Score for femoroacetabular impingement patients for reliability, construct validity and content validity and to determine minimal clinical important difference after 6 months.

Study objective

HOS is reliabel, has a good construct validity and content validity.

Study design

Two times with a one-week interval preoperatievely.

One time 6 months post-operatively.

Intervention

Patients will be asked to fill in five questionnaires at three moments: two pre-operative moments with a one-week interval, and one moment six months postoeratively. The required questionnaires are: HOS-NL, mHHS, HAGOS-NL, iHOT-12-NL and NRS for pain.

Contacts

Public

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Eligibility criteria

Inclusion criteria

18-65 years of age

Physical examination and radiological examination suspect for FAI

Inclusion will not interfere with standard care for FAI

Informed consent

Understand dutch language

Exclusion criteria

Prior hip surgery for FAI

Pathological fractures of metastic disease

Refuse to participate

Do not speak Dutch language

Study design

Design

Study type: Observational non invasive

Intervention model: Parallel

Allocation: Non controlled trial

Masking: Open (masking not used)

Control: N/A, unknown

Recruitment

NL

Recruitment status: Recruiting
Start date (anticipated): 01-11-2017

Enrollment: 140

Type: Anticipated

Ethics review

Positive opinion

Date: 25-10-2017

Application type: First submission

Study registrations

Followed up by the following (possibly more current) registration

ID: 47747

Bron: ToetsingOnline

Titel:

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

No registrations found.

In other registers

Register ID

NTR-new NL6018 NTR-old NTR6782

CCMO NL61937.098.17
OMON NL-OMON47747

Study results