

Longlasting adduction-related groin injuries in athletes; regular care or a novel treatment approach

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
19/07/2006	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
19/07/2006	Completed	<input type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
19/07/2006	Signs and Symptoms	<input type="checkbox"/> 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

7502.0005

Study information

Scientific Title

Acronym

LIES

Study objectives

Athletes with longstanding groin pain recover faster and more completely if they are treated with specific pelvic stabilizing muscle training compared with regular care.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled, parallel group trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Groin pain

Interventions

Two different kinds of physiotherapeutic treatments are given for the population. Both treatment strategies are already in use in daily practice.

Patients are randomised to receive either pelvic-stabilizing muscle training or usual care

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

1. Severity of the pain over the last three days (11-point visual analogue scale [VAS])

2. Participation in sports (11-point VAS-scale)

3. General disability (adapted Quebec low back pain disability scale)

4. Global change (six-point Likert scale)

5. How long before return to full athletic activity

6. Recurrences of the same complaints

Parameters 1, 2, 3 are measured before and directly after the period of treatment and 26 and 52 weeks after the start of treatment.

Parameters 4, 5 and 6 are only measured at 26 and 52 weeks after treatment.

Key secondary outcome(s)

1. Hip adduction strength (hand-held dynamometer);
2. Contraction pattern of the abdominal musculature (ultrasound echografie);
3. Active straight leg raise test (ASLR).

Completion date

15/08/2009

Eligibility

Key inclusion criteria

Male athletes, 18-45 years old, hip adduction-related complaints, for a period of at least six weeks, strong desire to compete in sports.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

45 years

Sex

Male

Key exclusion criteria

1. Pain as a result of high-impact trauma, suspicion of fracture, rupture of labrum of the hip, hip arthrosis or arthritis, femoral or inguinal hernia, radicular symptoms, infection of the urinary tract, bursitis, vessel disease, abnormal anatomy
2. Treated for the same (episode of) complaints previously, treated for low back pain with an exercise program in the previous six months, systemic diseases
3. Psychopathology
4. Physical handicaps that make it impossible for the subject to take part of the study

Date of first enrolment

01/03/2005

Date of final enrolment

15/08/2009

Locations

Countries of recruitment

Netherlands

Study participating centre
Heidelberglaan 100
Utrecht
Netherlands
3584 CX

Sponsor information

Organisation

University Medical Center Utrecht (UMCU) (The Netherlands)

ROR

<https://ror.org/0575yy874>

Funder(s)

Funder type

Research organisation

Funder Name

Netherlands Organisation for Health Research and Development (ZonMw)

Alternative Name(s)

Netherlands Organisation for Health Research and Development

Funding Body Type

Private sector organisation

Funding Body Subtype

Other non-profit organizations

Location

Netherlands

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration