Development, optimalisation and validation of the exercise diagnostics for functional iliac flow limitations in endurance athletes: blood pressure measurement, pedal force measurement and near infrared spectroscopy

	Prospectively registered
No longer recruiting	☐ Protocol
Overall study status	Statistical analysis plan
Completed	Results
Condition category	Individual participant data
Other	Record updated in last year
	Completed Condition category

Plain English summary of protocolNot provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr G. Schep

Contact details

Maxima Medisch Centrum Dept. Sportgeneeskunde Postbus 7777 Veldhoven Netherlands 5500 MB

Additional identifiers

Protocol serial number NTR526

Study information

Scientific Title

Study objectives

The use of a new protocol of exercise testing with use of blood pressure measurement, near infrared spectroscopy and pedal force measurement will improve the diagnostic value of the current decision algorithm used to diagnose functional iliac flow limitations in endurance athletes.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Received from the local medical ethics committee

Study design

Randomised active controlled parallel group trial

Primary study design

Interventional

Study type(s)

Diagnostic

Health condition(s) or problem(s) studied

Functional iliac flow limitations

Interventions

Blood pressure measurements, pedal force measurements and near infrared measurements during and after exercise testing in both patients and in healthy test subjects

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Improvement of sensitivity and specificity of diagnostic tools in diagnosing sports related flow limitations of the iliac arteries

Key secondary outcome(s))

No secondary outcome measures

Completion date

31/12/2007

Eligibility

Key inclusion criteria

Athletes/patients diagnosed with functional iliac flow limitations and healthy athletes

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Does not comply with the above inclusion criteria

Date of first enrolment

01/09/2005

Date of final enrolment

31/12/2007

Locations

Countries of recruitment

Netherlands

Study participating centre Maxima Medisch Centrum

Veldhoven Netherlands 5500 MB

Sponsor information

Organisation

Maxima Medical Center (Netherlands)

ROR

https://ror.org/02x6rcb77

Funder(s)

Funder type

Not defined

Funder Name

Not provided at time of registration

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration