

Pilot studies of the effects of custom semi-rigid full-foot orthotics on planta-surface foot pain and anterior knee pain: towards evidence-based practice

Submission date 12/09/2003	Recruitment status No longer recruiting	☐ Prospectively registered
		☐ Protocol
Registration date 12/09/2003	Overall study status Completed	☐ Statistical analysis plan
		☐ Results
Last Edited 21/11/2019	Condition category Signs and Symptoms	☐ Individual participant data
		☐ 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

N0544093555

Study information

Scientific Title

Pilot studies of the effects of custom semi-rigid full-foot orthotics on planta-surface foot pain and anterior knee pain: towards evidence-based practice

Study objectives

Orthotics treatment of foot and knee pain

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Signs and Symptoms: knee/foot pain

Interventions

Subjects will be referred via specialist orthopaedic and rheumatology clinics and will sign informed consent. They will be assessed by a Chartered Physiotherapist (Kristin Giussani) which will include measurements of strength, range of movement, and outcome measures of pain and function. They will then be fitted with either a real or placebo customised shoe insert (an orthotic) which they will use for a 1-month period. The patient will keep a diary of their pain and function over a 1-month period.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

21/10/2003

Eligibility**Key inclusion criteria**

Not provided at time of registration

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

31/10/2000

Date of final enrolment

21/10/2003

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Box 194

Cambridge

United Kingdom

CB2 2QQ

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

Cambridge Consortium - Addenbrooke's (UK)

Results and Publications

Individual participant data (IPD) sharing plan**IPD sharing plan summary**

Not provided at time of registration