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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

Registration date
12/09/2003

Overall study status
Completed

Last Edited
21/11/2019

Condition category
Signs and Symptoms

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[ ] Results
[ ] 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
Mr Joshua T Wies

Contact details
Box 194
Rheumatology Research Unit
Addenbrooke's NHS Trust
Cambridge
United Kingdom
CB2 2QQ

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

Study design
Randomised controlled trial

Primary study design
Interventional

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