

Pedometer use as an adjunct to exercise training in peripheral vascular disease

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 27/02/2020	Condition category Circulatory System	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr DC Mitchell

Contact details
Department of Surgery
North Bristol NHS Trust
Southmead Hospital
Westbury-on-Trym
Bristol
United Kingdom
BS10 5NB
+44 (0)117 959 5166
abc@email.com

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0234117604

Study information

Scientific Title

Pedometer use as an adjunct to exercise training in peripheral vascular disease

Study objectives

Does pedometer led exercise have a greater increase in exercise tolerance than targeted exercise alone?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled crossover group trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Other

Participant information sheet

Health condition(s) or problem(s) studied

Peripheral vascular disease

Interventions

Randomised crossover study, group - pedometer led exercise with a diary. Two group exercise alone with diary for 12 weeks - 2 weeks rest then swop over.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/09/2002

Completion date

30/09/2003

Eligibility

Key inclusion criteria

60 Patients.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

60

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/09/2002

Date of final enrolment

30/09/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Surgery

Bristol

United Kingdom

BS10 5NB

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

North Bristol NHS Trust (UK)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration