A randomised controlled trial to investigate the effect of regular moderate intensity exercise on plasma fibrinogen in people with type two diabetes: a pilot study

Submission date	Recruitment status	Prospectively registered
30/09/2005	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2005	Completed	Results
Last Edited	Condition category	Individual participant data
12/09/2016	Nutritional, Metabolic, Endocrine	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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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0249142478

Study information

Scientific Title

A randomised controlled trial to investigate the effect of regular moderate intensity exercise on plasma fibrinogen in people with type two diabetes: a pilot study

Study objectives

Does moderate intensity exercise modify plasma fibrinogen levels in people with type two diabetes?

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

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Type two diabetes

Interventions

Randomised controlled trial

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

- 1. Plasma fibrinogen
- 2. Plasminogen activator inhibitor-1 (PAI-1)

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/06/2003

Completion date

06/10/2004

Eligibility

Key inclusion criteria

80 patients with type two diabetes

Participant type(s)

Patient

Age group

Not Specified

Sex

Both

Target number of participants

80

Key exclusion criteria

- 1. Smokers
- 2. Taking drugs affecting clotting
- 3. Inability to exercise

Date of first enrolment

01/06/2003

Date of final enrolment

06/10/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre University of Bristol Bristol United Kingdom BS8 1TP

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Government

Funder Name

Taunton and Somerset Research and Development Consortium (UK)

Funder Name

NHS R&D Support Funding (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