Exercise and vitamin C supplementation in type one diabetes mellitus

Submission date 16/04/2008	Recruitment status No longer recruiting	Prospectively registered	
		[] Protocol	
Registration date 12/06/2008	Overall study status Completed	Statistical analysis plan	
		[X] Results	
Last Edited 27/03/2012	Condition category Nutritional, Metabolic, Endocrine	Individual participant data	

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s) Scientific

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N/A

Study information

Scientific Title

Molecular detection of exercise-induced free radicals following ascorbate prophylaxis in type one diabetes mellitus: a randomised controlled trial

Study objectives

Consistent with the human literature, we hypothesise that compared to healthy controls: 1. Exercise would compound basal oxidative stress in type one diabetics, and 2. Ascorbic acid would provide effective prophylaxis

A randomised, double-blind, placebo-controlled experimental design incorporating an electron paramagnetic resonance (EPR) spin-trapping technique combined with a comprehensive assessment of lipid peroxidation and non-enzymatic antioxidants will be employed to test these hypotheses.

Ethics approval required

Old ethics approval format

Ethics approval(s)

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Ethics approval received from The Bro Taf (Wales) Research Ethics Committee on the 18th December 1996 (ref: 96/1649).
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Study design

A balanced single-centre randomised double-blind placebo-controlled trial

Primary study design Interventional

Secondary study design Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Screening

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Type one diabetes mellitus

Interventions

Subjects were randomised to receive: 1. 1 g ascorbic acid two hours prior to exercise 2. Placebo two hours prior to exercise

Supplementation took place once only for both treatment groups, and the follow up was conducted immediately post-exercise and 24 hours later via telephone.

Intervention Type

Drug

Phase Not Specified

Drug/device/biological/vaccine name(s)

Ascorbic acid

Primary outcome measure

Free radical species in human blood, measured before supplementation, two hours after supplementation (before exercise) and immediately after exercise.

Secondary outcome measures

Blood biochemical markers such as lipid hydroperoxides, vitamin C and vitamin E, measured before supplementation, two hours after supplementation (before exercise) and immediately after exercise.

Overall study start date

01/01/2001

Completion date

01/06/2001

Eligibility

Key inclusion criteria

Diabetic volunteers were recruited from the adult diabetic clinic at the University Hospital of Wales, based on the following inclusion criteria:

- 1. Males aged 18 30 years
- 2. Glycosylated haemoglobin (HbA1c) of between 7 10%
- 3. Microalbuminuria negative
- 4. No underlying vascular complications

Non-diabetic control subjects were recruited from the student population of the University of Glamorgan, based on the following inclusion criteria:

- 1. Matched for age
- 2. Matched for fitness
- 3. No metabolic or circulatory medical condition
- 4. Had no family history of diabetes as confirmed via interview and medical history questionnaire

Participant type(s)

Patient

Age group Adult

Lower age limit 18 Years **Sex** Male

Target number of participants 12 male type one diabetic patients; 14 apparently healthy male volunteers

Key exclusion criteria

1. Cardiovascular disease

2. Hypertension

3. Any other known cardiac complication

Date of first enrolment 01/01/2001

Date of final enrolment 01/06/2001

Locations

Countries of recruitment United Kingdom

Study participating centre University of Ulster Newtownabbey United Kingdom BT37 OQB

Sponsor information

Organisation University of Glamorgan (UK)

Sponsor details

Pontypridd Cardiff Wales United Kingdom CF37 1DL

Sponsor type University/education

Website http://www.glam.ac.uk/ ROR https://ror.org/02mzn7s88

Funder(s)

Funder type Hospital/treatment centre

Funder Name University of Glamorgan (UK)

Funder Name University Hospital of Wales (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/11/2008		Yes	No