Exercise and vitamin C supplementation in type one diabetes mellitus

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

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 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)

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

Study design

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

Primary study design

Interventional

Study type(s)

Screening

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(s)

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

Key secondary outcome(s))

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.

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

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

Male

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)

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

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
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes