Does Vitamin D improve markers of vascular health in type 2 diabetes?

Submission date	Recruitment status	[X] Prospectively registered	
26/07/2005	No longer recruiting	☐ Protocol	
Registration date	Overall study status	Statistical analysis plan	
12/09/2005	Completed	[X] Results	
Last Edited	Condition category	Individual participant data	
29/08/2017	Nutritional, Metabolic, Endocrine		

Plain English summary of protocol

Background and study aims

Low levels of vitamin D have been associated with higher blood pressure and a higher risk of heart attack and stroke. These are particular problems for people with diabetes, and it is possible that giving extra vitamin D to people with diabetes who have low vitamin D levels might help to protect against heart attacks and strokes. The aim of this study is to find out whether giving vitamin D can reduce blood pressure, improve measures of blood vessel health, and improve glucose levels in diabetes.

Who can participate?

Patients aged over 18 with type 2 diabetes and low levels of vitamin D

What does the study involve?

Participants' vitamin D levels are tested and if these are low, the health of their blood vessels is assessed by doing an ultrasound scan of their arm before and after blowing up a blood pressure cuff on their arm. Blood samples are also taken to measure a series of substances that vitamin D might affect, and their blood pressure is measured. Participants are then randomly allocated take either a single large dose of vitamin D or a matching dummy (placebo) dose with no vitamin D in it. This dose acts over a period of several weeks. All participants are asked to return to repeat all of the tests 8 weeks after this single dose.

What are the possible benefits and risks of participating?

Vitamin D is a safe treatment that has been used for decades. There is a small risk of increasing calcium levels in the blood, but these levels are monitored and anyone with high levels at the start of the study is not included.

Where is the study run from? Ninewells Hospital (UK)

When is the study starting and how long is it expected to run for? October 2005 to June 2006

Who is funding the study? Diabetes UK

Who is the main contact? Dr Justine Davies j.i.davies@dundee.ac.uk

Contact information

Type(s)

Scientific

Contact name

Dr Justine Davies

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers JUS004

Study information

Scientific Title

Does Vitamin D improve markers of vascular health in type 2 diabetes? A randomised controlled trial

Study objectives

That Vitamin D improves endothelial function, pulse wave velocity, insulin sensitivity and markers of renin-angiotensin system activation in type 2 diabetics.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Tayside (now East of Scotland) Research Ethics committee, 13/09/2005, ref: 05/S1401/125

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

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

Health condition(s) or problem(s) studied

Type 2 diabetes mellitus

Interventions

100,000 units ergocalciferol versus placebo

Added 29/08/2017:

Intervention: 100,000 units oral Ergocalciferol (vitamin D2), given once at baseline

Comparator: Matching placebo, given once at baseline

Total duration of treatment and follow-up: 8 weeks for all treatment arms

Intervention Type

Supplement

Primary outcome measure

Current primary outcome measures as of 29/08/2017:

Flow-mediated dilatation of the brachial artery in response to reactive hyperaemia, measured by ultrasound at baseline and 8 weeks

Previous primary outcome measures:

Change in flow mediated dilatation of the brachial artery

Secondary outcome measures

Current secondary outcome measures as of 29/08/2017:

- 1. 25-hydroxyvitamin D, serum calcium, serum phosphate, parathyroid hormone levels, measured by blood tests at baseline and 8 weeks
- 2. Glycosylated haemoglobin, insulin sensitivity using HOMA index, measured by blood tests at baseline and 8 weeks
- 3. Office blood pressure, measured by blood pressure cuff at baseline and 8 weeks
- 4. Renin and angiotensin levels, measured by blood tests at baseline and 8 weeks

Previous secondary outcome measures:

Pulse wave velocity, homeostasis model assessment (HOMA) index, renin, angiotensin, Vitamin D, parathyroid hormone (PTH)

Overall study start date

01/10/2005

Completion date

30/06/2006

Eligibility

Key inclusion criteria

- 1. Age ≥18 years
- 2. Vitamin D level <50 nmol/l

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

60

Key exclusion criteria

- 1. Serum creatinine >200 µmol/l
- 2. Liver function tests >3 x upper limit of normal
- 3. Corrected calcium >2.55 mmol/l or <2.15 mmol/l
- 4. Metastatic malignancy
- 5. Unable to give informed consent

Date of first enrolment

01/11/2005

Date of final enrolment

31/03/2006

Locations

Countries of recruitment

Scotland

United Kingdom

Study participating centre Ninewells Hospital

Dundee United Kingdom DD1 9SY

Sponsor information

Organisation

University of Dundee (UK)

Sponsor details

11 Perth Road Dundee Scotland United Kingdom DD1 4HN +44 (0)1382 344664 j.z.houston@dundee.ac.uk

Sponsor type

University/education

ROR

https://ror.org/03h2bxq36

Funder(s)

Funder type

Charity

Funder Name

Diabetes UK

Alternative Name(s)

DIABETES UK LIMITED, British Diabetic Association

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

United Kingdom

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are available upon request from Dr Miles Witham (m.witham@dundee.ac.uk). Anonymised Individual participant data on all those randomised will be made available to bona fide researchers for non-commercial use, subject to sight of an analysis plan and subject to appropriate data sharing agreements and approval from the trial sponsor.

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/03/2008		Yes	No
Basic results		18/08/2017	29/08/2017	No	No