Do the remaining insulin-producing cells in people with type 1 diabetes help to maintain good glucose control after exercise?

Submission date	Recruitment status No longer recruiting	Prospectively registered		
15/05/2019		☐ Protocol		
Registration date 24/05/2019	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited	Condition category	Individual participant data		
14/06/2023	Nutritional. Metabolic. Endocrine			

Plain English Summary

Background and study aims

When people with Type 1 diabetes exercise, some experience hypoglycaemia (low blood sugar [glucose]), while others do not; in some HbA1c (a marker of diabetes control) gets worse while in others it improves. Exercise is known to increase glucose variability leading to more time with high and low levels. It is now known that many people with long-standing type 1 diabetes can produce small amounts of insulin from the remaining beta-cells in the pancreas. It is unknown if this is important for limiting blood glucose variability at rest and around exercise, and may explain some of the wide variation that is observed in response to exercise in people with Type 1 diabetes. This study aims to examine how residual beta-cell function impacts on glucose control when physically active / exercising in people with Type 1 diabetes.

Who can participate?

Anyone aged 18-65 years old with clinically diagnosed Type 1 diabetes, treated with exogenous insulin (pump or injection), free from diabetes complications can participate.

What does the study involve?

Participants will be required to complete a mixed meal tolerance test and a period of moderate intensity walking exercise for 45 minutes, with blood samples and interstitial glucose recorded before and after exercise

What are the possible benefits and risks of participating?

The benefits of taking part include understanding your own individual responses to exercise, receiving feedback on cardiovascular fitness, and contributing to the care and management of those with Type 1 diabetes. The risks of taking part include experiencing hypoglycaemia, musculoskeletal injury and muscle soreness.

Where is the study run from? Newcastle upon Tyne NHS Foundation Trust, UK. When is the study starting and how long is it expected to run for? October 2016 to May 2019.

Who is funding the study?

- 1. Diabetes Research and Wellness Foundation, UK
- 2. Newcastle University, UK

Who is the main contact?
Dr Daniel West,
daniel.west@newcastle.ac.uk
Mr Gary Taylor,
g.taylor3@newcastle.ac.uk

Contact information

Type(s)

Scientific

Contact name

Dr Daniel West

ORCID ID

http://orcid.org/0000-0003-2246-4925

Contact details

Institute of Cellular Medicine Room M4.077 William Leech Building Newcastle University Newcastle upon Tyne United Kingdom NE2 4HH 0191 2087076 daniel.west@newcastle.ac.uk

Type(s)

Scientific

Contact name

Mr Guy Taylor

ORCID ID

http://orcid.org/0000-0002-5207-1498

Contact details

Institute of Cellular Medicine Room M4.077 William Leech Building Newcastle University Newcastle upon Tyne United Kingdom NE2 4HH 0191 2087076 G.Taylor3@newcastle.ac.uk

Additional identifiers

EudraCT/CTIS number

Nil known

IRAS number

ClinicalTrials.gov number

Nil known

Secondary identifying numbers

C-peptide and exercise in T1D V3 14/07/16

Study information

Scientific Title

The role of residual beta-cell function on post-exercise glycaemic variability in individuals with type 1 diabetes

Study hypothesis

Type 1 diabetes patients with residual beta-cell function demonstrate improved post-exercise glucose control

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 02/09/2016 North East Tyne & Wear South Research Ethics Committee (NHSBT Newcastle Blood Donor Centre, Holland Dr, Newcastle upon Tyne NE2 4NQ; 0207 104 8026; nrescommittee.northeast-tyneandwearsouth@nhs.net), ref: 16/NE/0192

Study design

Acute observational trial

Primary study design

Observational

Secondary study design

Study setting(s)

Hospital

Study type(s)

Quality of life

Participant information sheet

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

Condition

Type 1 diabetes

Interventions

Patients with Type 1 diabetes with a wide range of residual beta-cell function (from negative to clinically significant) will be recruited. Participants will be identified using urinary C-peptide Creatinine Ratio testing, and those eligible will complete a mixed meal tolerance test to establish maximal stimulated serum C-peptide concentrations. Participants will then complete a fixed bout of moderate intensity walking exercise at 60% VO2 peak for 45 minutes, with blood samples and interstitial glucose recorded before and after exercise.

Intervention Type

Behavioural

Primary outcome measure

The amount of time interstitial glucose is spent in euglycaemia measured using blinded interstitial continuous glucose monitoring

Secondary outcome measures

- 1. Glycaemic variability (SD, CV%, MAGE, J-Index, CONGA, MAG, M-value)
- 2. Time spent: hypoglycaemic (<3.9mmol/L, <3.0mmol/L), hyperglycaemic (>10mmol/L, >13. 9mmol/L, >16.7mmol/L)
- 3. Hypoglycaemia stage 1 (<3.9mmol/L for 15+ minutes) and stage 2 (<3.0mmol/L for 15+ minutes) and hyperglycaemia incidence level 1 (>10mmol for 15+ minutes) and level 2 (>13. 9mmol for 15+ minutes)
- 4. Corrective bolus/carbohydrate intake

Overall study start date

01/08/2015

Overall study end date

01/09/2019

Eligibility

Participant inclusion criteria

- 1. Aged 18-65 years old
- 2. Clinically diagnosed Type 1 diabetes
- 3. Treated with exogenous insulin (pump or injection)
- 4. Free from diabetes complications

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Upper age limit

65 Years

Sex

Both

Target number of participants

30

Total final enrolment

30

Participant exclusion criteria

- 1. Type 1 diabetes participants duration of disease less than 1 year
- 2. HbA1c > 10% (86 mmol/mol)
- 3. Unable to complete maximal exercise test

Recruitment start date

01/10/2016

Recruitment end date

31/05/2019

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Newcastle upon Tyne NHS Foundation Trust

Level 1 Regent Point Gosforth Newcastle upon Tyne United Kingdom NE3 3HD

Sponsor information

Organisation

Newcastle University

Sponsor details

Faculty of Medical Sciences
The Medical School
Framlington Place
Newcastle upon Tyne
England
United Kingdom
NE2 4HH
+44 (0)191 208 6000
kay.howes@ncl.ac.uk

Sponsor type

University/education

Website

https://www.ncl.ac.uk/

ROR

https://ror.org/01kj2bm70

Funder(s)

Funder type

Research organisation

Funder Name

Diabetes Research and Wellness Foundation

Alternative Name(s)

Diabetes Research & Wellness Foundation, Diabetes Research and Wellness Foundation UK, DRWF

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

United Kingdom

Funder Name

Newcastle University

Alternative Name(s)

Funding Body Type

Private sector organisation

Funding Body Subtype

Universities (academic only)

Location

United Kingdom

Results and Publications

Publication and dissemination plan

Planned publication in a high-impact peer reviewed journal.

Intention to publish date

01/01/2020

Individual participant data (IPD) sharing plan

The current data sharing plans for this study are unknown and will be available at a later date

IPD sharing plan summary

Data sharing statement to be made available at a later date

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/10/2020	05/10/2020	Yes	No
Results article		10/03/2022	14/06/2023	Yes	No
HRA research summary			28/06/2023	No	No