

Evaluation of pancreas composition and function during return to normal glucose tolerance in type 2 diabetes

Submission date

21/06/2011

Recruitment status

No longer recruiting

Prospectively registered

Protocol

Registration date

21/06/2011

Overall study status

Completed

Statistical analysis plan

Results

Last Edited

21/06/2016

Condition category

Nutritional, Metabolic, Endocrine

Individual participant data

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

10274

Study information

Scientific Title

Evaluation of pancreas composition and function during return to normal glucose tolerance in type 2 diabetes

Acronym

DRN 591

Study objectives

The overall objective is to elucidate the pathophysiology of type 2 diabetes by defining the contribution of change in beta cell function and insulin sensitivity to the return to normal glucose homeostasis during definitive therapeutic intervention. The compelling in vitro and in vivo animal data for a direct adverse effect of fatty acid exposure on beta cell function and hepatic insulin sensitivity will be tested in vivo in people with type 2 diabetes by use of a newly devised magnetic resonance technique. The specific hypothesis will be tested that the early effect of bariatric surgery in normalising blood glucose in type 2 diabetes is brought about by the same mechanistic changes as seen during hypocaloric dieting.

Ethics approval required

Old ethics approval format

Ethics approval(s)

10/H0906/78

Study design

Randomised interventional and observational treatment case-controlled study

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Topic: Diabetes Research Network, Metabolic and Endocrine; Subtopic: Type 2, Metabolic and Endocrine (all Subtopics); Disease: Metabolic

Interventions

1. Subjects with and without type 2 diabetes will be investigated before and after gastric bypass surgery
2. Very low calorie diet (VLCD)
3. Subjects with T2DM will be randomised into 'surgery only' or 'VLCD+surgery' groups

Intervention Type

Mixed

Primary outcome(s)

First phase insulin response at baseline, 1 week after intervention and 8 weeks after surgery

Key secondary outcome(s)

1. Liver fat and hepatic insulin sensitivity at baseline, 1 week after intervention and 8 weeks after surgery
2. The treatment induced changes in pancreatic fat content at baseline, 1 week after intervention and 8 weeks after surgery

Completion date

01/08/2012

Eligibility

Key inclusion criteria

1. Type 2 diabetes duration less than 15 years
2. On treatment with diet alone or diet plus oral agents
3. BMI 35-45m/kg²
4. Listed for gastric bypass surgery
5. HbA1c < 10%

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Waist circumference >150cm
2. Kidney dysfunction (serum creatinine >150 µmol/l)
3. Liver dysfunction (serum ALT > 2.5 times the upper limit of normal)
4. Contraindications to MRI (metal implants and claustrophobia)
5. Alcohol consumption >14 units per week
6. Treatment with thiazolidinediones, insulin, GLP1 analogues or steroids
7. History of bowel surgery or inflammatory bowel disease

Date of first enrolment

01/02/2011

Date of final enrolment

01/08/2012

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Institute for Ageing and Health

Newcastle upon Tyne

United Kingdom
NE4 5PL

Sponsor information

Organisation

Newcastle upon Tyne Hospitals NHS Foundation Trust

ROR

<https://ror.org/05p40t847>

Funder(s)

Funder type

Research organisation

Funder Name

European Foundation for the Study of Diabetes (Germany)

Alternative Name(s)

The European Association for the Study of Diabetes, European Association for the Study of Diabetes (EASD), EFSD

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

Germany

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
	results				

[Results article](#)

01/01/2016

Yes

No