Effects of weight loss on stimulated lipolysis.

No registrations found.

Ethical review Positive opinion

Status Pending

Health condition type -

Study type Interventional

Summary

ID

NL-OMON24842

Source

NTR

Brief title

Beta2LIPWL

Health condition

Obesity, beta2-adrenergic sensitivity, lipolysis Obesitas, beta2-adrenerge gevoeligheid, lipolyse

Sponsors and support

Primary sponsor: Maastricht University

Source(s) of monetary or material Support: Not applicable

Intervention

Outcome measures

Primary outcome

Changes in sensitivity to β 2-AR-stimulated lipolysis in adipocytes isolated from subcutaneous adipose tissue biopsies.

Secondary outcome

- 1. Changes in markers of adipose tissue hypoxia;
- 2. Changes in markers of adipose tissue and systemic oxidative stress.

Study description

Background summary

This study investigates the effects of weight loss on beta2-adrenergic stimulated lipolysis in isolated adipocytes. In addition the effects of weight loss on adipose tissue hypoxia and adipose tissue and systemic oxidative stress will be determined.

Study objective

Moderate weight loss increases sensitivity to β 2-AR-stimulated lipolysis in isolated adipocytes from subcutaneous adipose tissue biopsies.

Study design

- 1. Premeasurements will be done before the start of the CO-EUR programme;
- 2. Postmeasurements will be done after 9 months of participation in the CO-EUR programme. Postmeasurements will be done in every subject, who is still participating in the Co-Eur programme after 9 months, regardless of the amount of weight lost or gained in those 9 months.

Intervention

Subjects follow the standard weight loss programme in Heerlen (NL) provided by CO-EUR without interference by the researchers.

CO-EUR is an obesity treatment centre which uses an 18-month-lasting programme with a multi-disciplinary approach, including diet, physical activity and behaviour change, to treat people with obesity.

Contacts

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Eligibility criteria

Inclusion criteria

- 1. All subjects entering the Co-Eur programme;
- 2. Age > 18 years old;
- 3. No more than 3 kg weight change in the past 3 months.

Exclusion criteria

- 1. Smoking;
- 2. Diabetes:
- 3. (Medicinal) use of β -blockers or β -agonists;
- 4. Pregnant women;
- 5. Blood clotting problems or (medicinal) use of anticoagulantia.

Study design

Design

Study type: Interventional

Intervention model: Parallel

Allocation: Non controlled trial

Masking: Open (masking not used)

Control: N/A, unknown

Recruitment

NL

Recruitment status: Pending

Start date (anticipated): 01-07-2011

Enrollment: 37

Type: Anticipated

Ethics review

Positive opinion

Date: 10-08-2011

Application type: First submission

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register ID

NTR-new NL2874 NTR-old NTR3019

Other METC: 11-3-019

ISRCTN wordt niet meer aangevraagd.

Study results

Summary results

N/A