

Fatty Acid Induced Oxidative Stress: its role in preventing hypoglycemia

Submission date 09/01/2006	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 09/01/2006	Overall study status Completed	☐ Statistical analysis plan ☐ Results
Last Edited 01/09/2009	Condition category Other	☐ Individual participant data ☐ Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
NTR517

Study information

Scientific Title

Acronym

FIOS: Fatty acid Induced Oxidative Stress

Study objectives

Elevated levels of Free Fatty Acids during fasting induce oxidative stress and cause insulin resistance to maintain euglycemia.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Non-randomised open label placebo controlled crossover group trial

Primary study design

Interventional

Study type(s)

Other

Health condition(s) or problem(s) studied

No condition, healthy person

Interventions

Subjects will undergo a period of fasting and are assigned to receive either acipimox (inhibitor lipolysis) 250 mg 4dd or placebo. Hereafter insulin sensitivity will be measured using stable isotope technique. Furthermore regulating hormones and lipids will be measured. Muscle specimens (v. lateralis) will be obtained for determination of intramyocellular lipids and transcription factors.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

acipimox

Primary outcome(s)

Insulin resistance, Free fatty acids and oxidative stress with and without acipimox.

Key secondary outcome(s)

Other measures of glucose homeostasis: glucoregulatory hormones, (adipo)cytokines.

Completion date

01/03/2006

Eligibility

Key inclusion criteria

1. 6 healthy men
2. 18-38 years
3. Body mass index (BMI) 20-25
4. Stable weight during the last 3 months

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

38 years

Sex

Male

Key exclusion criteria

1. Diabetes
2. Diabetes first degree relatives
3. Hypercholesterolemia
4. High intensity sport activities
5. Positive oral glucose tolerance testing

Date of first enrolment

01/01/2006

Date of final enrolment

01/03/2006

Locations

Countries of recruitment

Netherlands

Study participating centre

Academic Medical Center
Amsterdam
Netherlands
1100 DD

Sponsor information

Organisation

Academic Medical Centre (AMC) (Netherlands)

ROR

<https://ror.org/03t4gr691>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Academic Medical Centre (AMC) (Netherlands), Department of Endocrinology and Metabolism

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration