Effects of cinnamon on postprandial blood glucose, and insulin in subjects with type 2 diabetes or impaired glucose tolerance.

No registrations found.

Ethical review Positive opinion

Status Recruiting

Health condition type -

Study type Interventional

Summary

ID

NL-OMON21422

Source

NTR

Brief title

N/A

Health condition

type 2 diabetes or impaired glucose tolerance

Sponsors and support

Primary sponsor: Department of Medicine, Malmö University Hospital **Source(s) of monetary or material Support:** Supported by Hans-Gabriel and Alice Trolle-Wachtmeister's Foundation for Medical Research.

Intervention

Outcome measures

Primary outcome

This study was therefore designed to determine whether cinnamon lower postprandial blood glucose, and insulin levels in subjects with type 2 diabetes or impaired glucose tolerance.

Secondary outcome

N/A

Study description

Background summary

N/A

Study objective

This study was therefore designed to determine whether cinnamon lower postprandial blood glucose, and insulin levels in subjects with type 2 diabetes or impaired glucose tolerance.

Study design

The study started on 11 May 2009 and will probably end on 11 September 2009.

Intervention

The test meal consisted of 50 g oral glucose tolerance test with added 6.9 g lactose ingested with 15 capsules with cinnamon. The reference meal consisted of 50 g oral glucose tolerance test ingested with 15 placebo capsules. The meals were served in random order at intervals of 1 week. Randomization was performed using a table of random numbers.

Contacts

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Eligibility criteria

Inclusion criteria

Patients were selected for the study on the basis of the following inclusion criteria:

- 1. Diagnosis of type 2 diabetes for < 6 months before enrollment;
- 2. Diagnosis of impaired glucose tolerance for < 6 months before enrollment.

Exclusion criteria

Patients who had thyroid disorders, or used insulin, oral hypoglycemics, insulin-sensitizing drugs, and â-blockers within 60 days before enrollment were excluded.

Study design

Design

Study type: Interventional

Intervention model: Crossover

Allocation: Non controlled trial

Masking: Double blinded (masking used)

Control: Placebo

Recruitment

NL

Recruitment status: Recruiting
Start date (anticipated): 11-05-2009

Enrollment: 15

Type: Anticipated

Ethics review

Positive opinion

Date: 20-05-2009

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 NL1711 NTR-old NTR1821

Other : Dnr 353/2008

ISRCTN wordt niet meer aangevraagd

Study results

Summary results

N/A