

Metabolic effects of Diabecinn (oral cinnamon extract) in diabetes type 2, a placebo-controlled randomized clinical trial

Submission date 28/04/2006	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 28/04/2006	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 28/04/2006	Condition category Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr J.H. DeVries

Contact details

Academic Medical Center
Department of Internal Medicine, F4-222
P.O. Box 22660
Amsterdam
Netherlands
1100 DD
+31 (0)20 5669111
j.h.devries@amc.uva.nl

Additional identifiers

Protocol serial number

N/A

Study information

Scientific Title

Acronym

Diabecinn trial

Study objectives

The main objective of this randomized, placebo-controlled trial is to determine the effects of cinnamon extract on HbA1c and lipid profiles in type 2 diabetic patients.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics approval received from the local medical ethics committee

Study design

Randomised placebo-controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Diabetes Mellitus type 2 (DM type II)

Interventions

Diabecinn three times a day (tid) or placebo

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Diabecinn

Primary outcome(s)

HbA1c.

Key secondary outcome(s)

1. Lipid profile
2. 6 point glucose profile
3. Hypoglycemia
4. Body weight
5. Free fatty acids
6. C-reactive protein (CRP)

Completion date

01/01/2007

Eligibility

Key inclusion criteria

1. Type 2 diabetes patients
2. Age 35-70 years inclusive
3. HbA1c between 7 and 12% inclusive

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Pregnancy
2. Breast-feeding

Date of first enrolment

01/05/2006

Date of final enrolment

01/01/2007

Locations

Countries of recruitment

Netherlands

Study participating centre

Academic Medical Center

Amsterdam

Netherlands

1100 DD

Sponsor information

Organisation

Academic Medical Center (AMC) (The Netherlands)

ROR

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

Funder(s)**Funder type**

Industry

Funder Name

OTC Pharma

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration