

Effect of Chitosan on Health and Obesity

Submission date 16/12/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 16/12/2002	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 08/11/2022	Condition category Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number
Nil known

IRAS number

ClinicalTrials.gov number
Nil known

Secondary identifying numbers
HRC 01/416

Study information

Scientific Title

Effect of Chitosan on Health and Obesity

Acronym

ECHO

Study objectives

Results from trials of chitosan as a weight loss treatment have provided conflicting results. A 1998 review of 5 studies suggested that chitosan was an effective treatment for overweight and obesity. However, many more chitosan trials have been carried out since then with variable results. In order resolve the uncertainty surrounding the effectiveness of this dietary supplement, we conducted a large randomised controlled clinical trial of the effect of chitosan on body weight, lipids, and other health outcomes.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Quality of life

Participant information sheet

Health condition(s) or problem(s) studied

Obesity

Interventions

3 g Chitosan daily versus placebo.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Chitosan

Primary outcome measure

The primary aim of the ECHO study was to determine if chitosan was an effective weight loss treatment for people who were overweight or obese.

Measures included changes in body weight (kilograms); body mass index, waist circumference; body fat percentage; systolic and diastolic blood pressure; serum cholesterol and lipids; plasma glucose; fat-soluble vitamins; faecal fat losses and health-related quality of life, from baseline to 24 weeks.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/11/2001

Completion date

30/06/2003

Eligibility**Key inclusion criteria**

1. Overweight/obese male or female volunteers
2. Aged greater than 18 years
3. Body Mass Index (BMI) between 28 - 50 kg/m²

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

250

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/11/2001

Date of final enrolment

30/06/2003

Locations

Countries of recruitment

New Zealand

Study participating centre

Clinical Trials Research Unit

Auckland

New Zealand

-

Sponsor information

Organisation

Health Research Council of New Zealand (New Zealand)

Sponsor details

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Sponsor type

Research council

ROR

<https://ror.org/00zbf3d93>

Funder(s)

Funder type

Research council

Funder Name

The Health Research Council of New Zealand (New Zealand)

Funder Name

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

Not provided at time of registration

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Systematic review	01/09/2004		Yes	No
Results article		17/12/2004		Yes	No
Other publications		01/02/2005		Yes	No