# Personalized diets for treatment of fatty liver and central obesity

| Submission date<br>20/06/2024 | <b>Recruitment status</b> No longer recruiting | <ul> <li>Prospectively registered</li> </ul> |  |  |
|-------------------------------|--|--|--|--|
|                               |  | ☐ Protocol                                   |  |  |
| Registration date 26/06/2024  | <b>Overall study status</b> Completed          | Statistical analysis plan                    |  |  |
|                               |  | [X] Results                                  |  |  |
| <b>Last Edited</b> 25/06/2024 | <b>Condition category</b><br>Digestive System  | Individual participant data                  |  |  |

# Plain English summary of protocol

Background and study aims

Dietary interventions are a new and promising way to treat hepatic steatosis, which is the buildup of fat in the liver, and visceral adiposity, which is the buildup of fat around the central organs in the body.

One type of dietary treatment involves prebiotics. Prebiotics are types of fiber and resistant carbohydrates that affect the gut microbiome. The gut microbiome consists of trillions of bacteria, yeast, and other microbes living in the intestine.

Our research focuses on the potential of xylo-oligosaccharides (XOS), a type of fiber, to treat these conditions. We also study how this prebiotic affects the gut microbiome and identify who might benefit the most from these interventions.

Who can participate?

Our study recruited overweight or obese adults aged 18 - 75 years

What does the study involve?

Participants ingest a dose of XOS daily for four months, preceded by 1 month without XOS. We measured their body composition and liver fat content in three time points and also collected blood and fecal samples to study the gut microbiome.

What are the possible benefits and risks of participating?

Participants receive potential health benefits from the dietary intervention and receive comprehensive information about their health and wellbeing. A possible adverse effect of XOS is gastrointestinal distress. Participants are inquired about adverse effects weekly and can opt out at any time.

Where is the study run from? Research Council of Finland

When is the study starting and how long is it expected to run for? January 2019 to June 2020

Who is funding the study? Research Council of Finland Juho Vainio Foundation (Finland) Sydäntutkimussäätiö (The Finnish Foundation for Cardiovascular Research)

Who is the main contact? satu.p.pekkala@jyu.fi jukka.e.hintikka@jyu.fi

# Contact information

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Public, Principal investigator

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Scientific

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# Additional identifiers

Clinical Trials Information System (CTIS)

# ClinicalTrials.gov (NCT)

Nil known

#### Protocol serial number

jyu-resd.172576021

# Study information

#### Scientific Title

Personalized diet intervention to treat fatty liver and visceral adiposity

#### Acronym

**MAKSA** 

## **Study objectives**

XOS intervention manipulates the gut microbiome and benefits hepatic health in overweight responsive individuals

## Ethics approval required

Ethics approval required

## Ethics approval(s)

approved 27/11/2019, Ethics Committee of the Hospital District of Southwest Finland (P.O. Box 52, Turku, 20251, Finland; +358 504383708; eettinen.toimikunta@varha.fi), ref: ETMK 72/2019

# Study design

Single group controlled experimental study

# Primary study design

Interventional

# Study type(s)

Other, Quality of life

# Health condition(s) or problem(s) studied

Overweight, non-alcoholic fatty liver disease

#### Interventions

Prebiotic nutritional supplementation: xylo-oligosaccharides 2.8 g daily in powder form. Each participant went through a 1-month control period, followed by 4 months of dietary intervention.

#### Intervention Type

Supplement

#### Primary outcome(s)

Change in liver fat, assessed with MRI at baseline (0), pre-intervention (4 wks), and post-intervention (16 wks)

## Key secondary outcome(s))

- 1. Responses in gut microbial composition or diversity, measured with 16S rRNA sequencing at baseline (0), pre-intervention (4 wks), and post-intervention (16 wks)
- 2. Determinants of response to the prebiotic, measured with microbiome, metabolome, GWAS at baseline (0), pre-intervention (4 wks), and post-intervention (16 wks)

# Completion date

01/06/2020

# **Eligibility**

#### Key inclusion criteria

- 1. Age 18<75years
- 2. Being overweight (body mass index [BMI] >25 kg/m²
- 3. High waist circumference (>102cm for males, >88cm for females)

# Participant type(s)

Healthy volunteer

# Healthy volunteers allowed

No

# Age group

Adult

# Lower age limit

18 years

# Upper age limit

75 years

#### Sex

All

#### Total final enrolment

49

#### Key exclusion criteria

- 1. Antibiotic treatment 1 month prior to the study
- 2. Excessive alcohol consumption (>20 g/day for females, 30 g/day for males)
- 3. Inflammatory bowel disease
- 4. Celiac disease
- 5. Major eating disorders
- 6. Hypothyroidism

#### Date of first enrolment

01/01/2020

#### Date of final enrolment

# Locations

#### Countries of recruitment

Finland

Study participating centre
University of Jyvaskyla, Faculty of Sport and Health Sciences
Rautpohjankatu 8
Jyvaskyla
Finland
40700

# Sponsor information

# Organisation

Academy of Finland

#### **ROR**

https://ror.org/05k73zm37

# Funder(s)

# Funder type

Research council

#### **Funder Name**

Research Council of Finland

# Alternative Name(s)

Academy of Finland, Suomen Akatemia, Finlands Akademi, AKA

# **Funding Body Type**

Government organisation

# **Funding Body Subtype**

Research institutes and centers

#### Location

Finland

#### **Funder Name**

Juho Vainio Foundation

# Alternative Name(s)

Academy of Finland, Suomen Akatemia, Finlands Akademi, AKA

#### **Funding Body Type**

Government organisation

## **Funding Body Subtype**

Research institutes and centers

#### Location

Finland

#### **Funder Name**

Sydäntutkimussäätiö

#### Alternative Name(s)

Finnish Foundation for Cardiovascular Research

#### **Funding Body Type**

Private sector organisation

#### **Funding Body Subtype**

Trusts, charities, foundations (both public and private)

#### Location

Finland

# **Results and Publications**

# Individual participant data (IPD) sharing plan

Restricted use due to personal information protection. You can still contact author to ask for a copy of the material.

Metadata for the project is shared at https://doi.org/10.17011/jyx/dataset/85068

# IPD sharing plan summary

Available on request

#### Study outputs

| Output type               | Details | Date created | Date added | Peer reviewed? | Patient-facing? |
|---------------------------|---------|--------------|------------|----------------|-----------------|
| Interim results article   |         | 30/01/2023   | 20/06/2024 | Yes            | No              |
| Other unpublished results |         |              | 20/06/2024 | No             | No              |