

Behavioural medicine for lower-educated patients with type 2 diabetes: the effectiveness and implementation of a tailored intervention in primary care

Submission date 21/03/2022	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 29/03/2022	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 20/01/2025	Condition category Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

This study is focused on behavioural medicine for lower educated type 2 diabetes patients. This study aims to test the effectiveness of a behavioural programme in lower educated type 2 diabetes patients in primary care.

Who can participate?

Eligible participants are between 35 and 70 years old and should be diagnosed at least one year with type 2 diabetes.

What does the study involve?

Patients in the intervention group will receive the behavioural programme 'Powerful together with diabetes' for 27 weeks in weekly group sessions, patients in the control group receive standard care. The 'Powerful together with diabetes' programme aims to support and empower patients to improve their diabetes self-management such as control of blood sugar levels (glycaemic control) and medication adherence. Improvement in glycaemic control will be compared across the intervention and control groups using data from routinely collected blood samples. Participants will also complete questionnaires on their quality of life throughout the study.

What are the possible benefits and risks of participating?

The benefits of participating are that patients' glycaemic control is improved and improvement of their medication adherence and quality of life.

Where is the study run from?

Amsterdam UMC, location AMC (Netherlands)

When is the study starting and how long is it expected to run for?

From February 2021 to October 2025

Who is funding the study?
Netherlands Organisation for Health Research and Development (ZonMw) (Netherlands)

Who is the main contact?
Bedra Horreh
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Study website
<https://www.samensterkmetstuiker.nl/>

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
555003202

Study information

Scientific Title
The DISC-2 (Diabetes in Social Context) study: an effectiveness-implementation hybrid evaluation of a group-based self-management intervention for lower educated T2DM patients

Acronym

Study objectives

Glycaemic control is improved in patients receiving the behavioural programme 'Powerful together with diabetes' as compared to patients not receiving the programme.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 07/02/2022, Medical Ethics Committee of the Academic Medical Center (AMC), (Meibergdreef 9, 1105 AZ, Amsterdam; +310205667389; mecamc@amsterdamumc.nl), ref: 2021_222 - NL79337.018.21

Study design

Single center interventional quasi-experimental non-randomized trial

Primary study design

Interventional

Secondary study design

Non randomised study

Study setting(s)

GP practice

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Type 2 diabetes mellitus

Interventions

Current interventions as of 16/10/2024:

Patients in GP practices receive in the intervention group the 'Powerful together with diabetes' program in 27 weekly group sessions. Patients in the control group receive standard care.

Previous interventions:

Patients in GP practices receive in the intervention group the 'Powerful together with diabetes' program for 1 year in weekly group sessions. Patients in the control group receive standard care.

Intervention Type

Behavioural

Primary outcome measure

Current primary outcome measure as of 16/10/2024:

HbA1c levels are retrieved from GP registries through a database owned by a GP collective at baseline, 12, and 24 months

Previous primary outcome measure:

HbA1c levels are retrieved from GP registries through a database owned by a GP collective at baseline, 6, 12, and 24 months

Secondary outcome measures

Current secondary outcome measures as of 16/10/2024:

1. Prescribed diabetes type 2 medication is retrieved from GP registries from a database owned by a GP collective at baseline, 12, and 24 months
 2. Use of primary and specialist care is retrieved from GP registries from a database owned by a GP collective at baseline, 12, and 24 months
 3. Quality of life is measured through a questionnaire at baseline, 6, 12, and 24 months (in the intervention group only)
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Previous secondary outcome measures:

1. Prescribed diabetes type 2 medication is retrieved from GP registries from a database owned by a GP collective at baseline, 6, 12, and 24 months
2. Use of primary and specialist care is retrieved from GP registries from a database owned by a GP collective at baseline, 6, 12, and 24 months
3. Quality of life is measured through a questionnaire at baseline, 6, 12, and 24 months (in the intervention group only)

Overall study start date

15/02/2021

Completion date

15/10/2024

Eligibility

Key inclusion criteria

1. At least one year since diagnosed with type 2 diabetes
2. Aged >35 and <70 years
3. Current diabetes self-management education is insufficient

Participant type(s)

Patient

Age group

Adult

Lower age limit

35 Years

Upper age limit

70 Years

Sex

Both

Target number of participants

722

Key exclusion criteria

1. Objection against participation from the GP
2. A psychiatric disorder which hampers participation
3. Being unable to come to the intervention location independently
4. Planning to stay abroad for a longer time during the intervention period

Date of first enrolment

14/02/2022

Date of final enrolment

30/04/2025

Locations**Countries of recruitment**

Netherlands

Study participating centre

Amsterdam UMC, location AMC

Meibergdreef 9

Amsterdam

Netherlands

1105 AZ

Sponsor information**Organisation**

Amsterdam University Medical Centers

Sponsor details

Meibergdreef 9
Amsterdam
Netherlands
1105 AZ
+31 205669111
fs-amr@amc.uva.nl

Sponsor type

Hospital/treatment centre

Website

<https://www.amsterdamumc.org/>

ROR

<https://ror.org/05grdyy37>

Funder(s)

Funder type

Research organisation

Funder Name

ZonMw

Alternative Name(s)

Netherlands Organisation for Health Research and Development

Funding Body Type

Private sector organisation

Funding Body Subtype

Other non-profit organizations

Location

Netherlands

Results and Publications

Publication and dissemination plan

Planned publication in a peer-reviewed journal.

Intention to publish date

30/11/2025

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are/will be available upon request from Bedra Horreh (b.horreh@amsterdamumc.nl).

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol article		09/01/2025	20/01/2025	Yes	No