

# Study on the benefit of quercetin intake in diabetic patients treated with antidiabetic tablets.

<b>Submission date</b> 20/11/2022	<b>Recruitment status</b> No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 17/01/2023	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 05/02/2025	<b>Condition category</b> Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

### Background and study aims

Quercetin has a scientifically proven anti-oxidant, anti-inflammatory and anti-coagulant effect. Nevertheless, the anti-aging effect of quercetin has been studied exclusively in vitro (test-tube experiments) with promising results. Thus, research is needed to examine quercetin's anti-aging effect in humans. The aim of this study is to investigate the benefit of quercetin in improving health and daily life.

### Who can participate?

The selection of participants will be performed in a random manner from the registered population of responsibility of the 4th Public Health Unit (TOMY) of Heraklion and will include 100 patients with type II diabetes, treated with antidiabetic tablets, over 50 years old, who will be asked to participate in this research.

### What does the study involve?

Participants will be randomly allocated to receive the quercetin supplement (intervention group) or not (control group).

Laboratory tests will be performed at the beginning and end of the study in both groups. The research is divided into the following parts: A pool of potential patients-participants will be established by recording and assessing demographic, anthropometric characteristics and living habits (gender, age, smoking habits, diet, physical activity, medical treatment). Of these, those who after thorough information accept and consent to participate will be included, while particularly vulnerable individuals will be excluded.

The total participation time is 8 months. The study will be conducted in the 4th TOMY (Local Health Unit) of Heraklion, 7th HPE (Health District) of Crete.

The time and dosage of taking quercetin will be based on the indications of supplement in question. Specifically, the intervention group will be given 500 mg of quercetin per day for 12 weeks, then stopped for 8 weeks and re-administered for another 12 weeks.

Blood samples will be taken from participants for laboratory testing at baseline (t=0) and end of the study (t= 8 mo). The sample is necessary to measure telomere length (genetic data), which will give us information about cellular aging.

At the same time, the participants will be asked to consent to the measurement of their anthropometric indicators, such as BMI and waist circumference, while they will also undergo spirometry with a piko-meter in the aforementioned phases.

What are the possible benefits and risks of participating?

The risks associated with quercetin supplementation are considered minimal. However, one potential inconvenience may be the time required to complete the study.

Although there are no direct benefits resulting from the participation in this research, it may help by highlighting the action of quercetin and subsequently facilitating the adoption of targeted interventions. Participants may benefit from the action of this supplement. There will be no personal compensation for participation in the study.

Where is the study run from?

This study will be conducted in the 4th Local Health Unit of Heraklion (TOMY), Crete.

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

January 2021 to January 2024

Who is funding the study?

Investigator initiated and funded

Who is the main contact?

PhD Candidate Aikaterini Mantadaki

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## Contact information

### Type(s)

Principal investigator

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## Additional identifiers

### Clinical Trials Information System (CTIS)

Nil known

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

2934/253

## Study information

### Scientific Title

Study on the benefit of quercetin intake in diabetic patients treated with antidiabetic tablets. A two-arm, prospective randomized control clinical trial (interventional vs observation group) in primary health care.

### Study objectives

In the present study, the hypothesis that the intake of quercetin could be related to an improvement of the body's antioxidant status and an antiaging effect will be examined. The main research question investigates the occurrence of antiaging action. The secondary question concerns the antioxidant effect resulting from the intake of quercetin.

### Ethics approval required

Old ethics approval format

### Ethics approval(s)

Approved 20/08/2021, University of Crete Research Ethics Committee (Rectors Office, University of Crete, Voutes University Campus Heraklion, Crete, 71300, Greece; +302810395206; ehde@uoc.gr), ref: 104/20.08.2021.

### Study design

Interventional randomized controlled clinical trial

### Primary study design

Interventional

### Study type(s)

Other

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

Diabetes mellitus

### Interventions

This is an interventional study, in which a nutritional supplement will be used in a group of diabetic patients treated with antidiabetic tablets, who will receive the supplement in question alongside their treatment (intervention group) and a group which will not be administered to the nutritional supplement alongside the treatment (control group). The nutritional supplement that will be administered is quercetin. The time and dosage of supplementation of quercetin will respect the indications of the nutritional supplement. Specifically, the intervention group will be given 500 mg of quercetin per day for 12 weeks, which will then be ceased for 8 weeks and re-administered for another 12 weeks. A random allocation ratio will be used for the recruitment of the patients in the intervention and control group, taking into account age and gender.

## Intervention Type

Supplement

### Primary outcome(s)

Telomere length will be measured using quantitative Polymerase Chain Reaction (qPCR), before (t=0) and after the intervention (t=32 weeks) for both groups

### Key secondary outcome(s)

1. [CBC] measured using [Beckman Coulter method of counting and sizing], before and after the intervention for both study groups
2. [IL-6] measured using [electrochemiluminescence] before and after the intervention for both study groups
3. [reticulocytes] measured using [Methylene blue stain (NMB)] before and after the intervention for both study groups
4. [hemoglobin] measured using [Beckman Coulter method of counting and sizing] before and after the intervention for both study groups
5. [WBC] measured using [Beckman Coulter method of counting and sizing] before and after the intervention for both study groups
6. [CRP] measured using [latex indirect agglutination (LIA)] before and after the intervention for both study groups
7. [HbA1c] measured using [high-performance liquid chromatography (HPLC)] before and after the intervention for both study groups
8. [TC] measured using [enzymatic method] before and after the intervention for both study groups
9. [LDL] measured using [Friedewald calculation] before and after the intervention for both study groups
10. [TGL] measured using [GPO/POD] before and after the intervention for both study groups
11. [creatinine] measured using [Creatinine amidohydrolase enzymatic method] before and after the intervention for both study groups
12. [serum nitrogen] measured using [Nesslerization] before and after the intervention for both study groups
13. [DHEAS] measured using [DHEA Sulfate Test] before and after the intervention for both study groups,
14. [25-OH-vitamin D] measured using [immunoassay, Liaison 25(OH) Vitamin D Total assay] before and after the intervention for both study groups
15. [FEV1] measured using piko-spirometry before and after the intervention for both study groups

### Completion date

29/01/2024

## Eligibility

### Key inclusion criteria

Eligible participants will be type II diabetes patients treated with antidiabetic tablets, with an age over 50 years who belong to the registered population of responsibility of the 4th Local Health Unit (TOMY) of Heraklion.

### Participant type(s)

Patient

**Healthy volunteers allowed**

No

**Age group**

Other

**Lower age limit**

50 years

**Sex**

All

**Total final enrolment**

100

**Key exclusion criteria**

Highly vulnerable individuals will be excluded, following the relevant legal and ethical rules. The above cases are also excluded:

1. Gestational diabetes (breastfeeding/pregnancy),
2. Maturity-onset diabetes of the young (MODY),
3. Pancreatogenic diabetes (T3CDM),
4. Heart, liver or kidney disease,
5. Serious psychiatric illness or addiction,
6. Patients with comorbidities such as cancer, AIDS, neoplastic diseases,
7. Organ transplant patients, patients under immunosuppressive therapy,
8. Patients who have recently suffered a cardiovascular event, within three months before the start of the study,
9. Patients using other antioxidant supplementations,
10. Patients with a requirement for long-term use of aspirin, other than low dosage for protection against cardiovascular events, or NSAIDs,
11. Patients participating in another interventional clinical study,
12. Patients with planned surgery or other interventional procedure requiring systemic anesthesia during the study.

**Date of first enrolment**

07/02/2023

**Date of final enrolment**

30/05/2023

**Locations****Countries of recruitment**

Greece

**Study participating centre**

**4th Local Health Unit of Heraklion (TOMY)**

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Heraklion Crete

Greece  
71303

## Sponsor information

### Organisation

University of Crete

### ROR

<https://ror.org/00dr28g20>

## Funder(s)

### Funder type

Other

### Funder Name

Investigator initiated and funded

## Results and Publications

### Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are not expected to be made available due to the sensitivity of the genetic, bioclinical and personal data of study participants.

### IPD sharing plan summary

Not expected to be made available

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>		29/08/2024	05/02/2025	Yes	No
<a href="#">Results article</a>		15/06/2024	05/02/2025	Yes	No
<a href="#">Results article</a>		14/04/2024	05/02/2025	Yes	No