

Clinical Trial Protocol

Iranian Registry of Clinical Trials

07 Jun 2026

Effect of Fenugreek, Cranberries, Chicory and Nettle seed extract on blood glucose in type 2 diabetic patients

Protocol summary

Study aim

A new drug discovery and a lower financial bug for patients with type 2 diabetes. The definition of a new herbal medicine is due to traditional treatments that are more consistent with indigenous culture. Localize the treatment of type 2 diabetes and discover less complicated drugs.

Design

Two arm parallel group randomized trial with blinded postoperative care and outcome assessment

Settings and conduct

The study is conducted in Baqiyatallah Hospital, a tertiary care hospital, in 2017-2018. Lab Tests were calculated before and after study. The level of FBS is checked by the participant by the glucometer at home, daily for 3 days, then 2 days once for 4 days, then on days 15, 30, 45 and 60 and the numbers obtained are recorded in certain forms. Patients will be started at the beginning of the study. At the end of the study, FBS, Lipid profile, BUN, SGOT, SGPT, ALP, HbA1C, Creatinine will be performed. In order to ensure the health of the herbal medicine, liver and kidney function tests for the first and second months are also examined. Patients and investigator were blinded.

Participants/Inclusion and exclusion criteria

Non-insulin-dependent diabetes mellitus treated with a diet or oral medication, Not history of any disease, Non-pregnancy and breastfeeding, The occurrence of any unwanted adverse drug reaction in the requested serial tests and any unwanted gastrointestinal complications and the unwillingness of the participants to continue the study are the main exclusion criteria.

Intervention groups

Patients will receive 2, 1000 mg capsules twice daily containing one quarter of the effective dose of herbs, morning and night after the meal for three months. The control group, the same as the first group, the intervention and the placebo were similar to the treatment with rusks bread powder, which will be in two

capsules of 1000 mg.

Main outcome variables

FBS, LFT, kidney function tests

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20190112042332N1**

Registration date: **2019-01-26, 1397/11/06**

Registration timing: **retrospective**

Last update: **2019-01-26, 1397/11/06**

Update count: **0**

Registration date

2019-01-26, 1397/11/06

Registrant information

Name

Khalil Mohamdzadeh

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2017-09-23, 1396/07/01

Expected recruitment end date

2019-01-20, 1397/10/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date
empty

Scientific title
Effect of Fenugreek, Cranberries, Chicory and Nettle seed extract on blood glucose in type 2 diabetic patients

Public title
Effect of Fenugreek, Cranberries, Chicory and Nettle seed extracts on blood glucose in type 2 diabetic patients

Purpose
Treatment

Inclusion/Exclusion criteria
Inclusion criteria:
Male and female subjects between 20 and 70 years of age Noninsulin-dependent diabetes minimum of 2 years before the study enrollment Patients with diabetes Type 2 and on any antidiabetic medication (oral hypoglycemic agent, insulin) Patients willing to give informed consent Patients with out other comorbid conditions (cardiac, renal, macrovascular and microvascular complication) Non Gangren's diabetic foot ulcer Not Currently pregnant or breastfeeding, fertile women with out using any contraceptives methods Current daily tobacco smoker and Alcohol consumption with out any decision to change diet and physical style Not having a history of cancer less than 5 years and a depression
Exclusion criteria:
Evidence of any unwanted side effects of the drug include increased liver enzymes or functional renal dysfunction or Lower blood glucose control in the requested serial tests Evidence of any unwanted gastrointestinal discomfort, such as diarrhea, nausea, or gastrointestinal intolerance of the combined medication Any unwillingness of participants to continue to participate in the plan

Age
From **20 years** old to **70 years** old

Gender
Both

Phase
N/A

Groups that have been masked

- Participant
- Care provider
- Outcome assessor
- Data analyser

Sample size
Target sample size: **100**
Actual sample size reached: **100**

Randomization (investigator's opinion)
Randomized

Randomization description
Treatment assignment is randomly assigned, in a 1:1 ratio, at each site using a block size of 6. Herbal and placebo medicines are packed in packages that are quite similar to those labeled A and B.To select patients, each patient should first receive a number from 1 to 100 (total sample size) given randomized visits to patients from closed medicines.

Blinding (investigator's opinion)
Double blinded

Blinding description
The treatment assignments are known to the study coordinator. Participants, treating clinicians, and outcome assessors are blinded to intervention group status.

Placebo
Used

Assignment
Parallel

Other design features

Secondary Ids
empty

Ethics committees

1

Ethics committee

Name of ethics committee
Research Ethics Committee of Baqiyatallah University of Medical Sciences

Street address
Baqiyatallah University of Medical Sciences, Mullasadra Street, Vanak Square, Tehran

City
Tehran

Province
Tehran

Postal code
548719395

Approval date
2017-10-29, 1396/08/07

Ethics committee reference number
IR.BMSU.REC.1394.87

Health conditions studied

1

Description of health condition studied
Blood Glucose Control in Type 2 Diabetic patients

ICD-10 code
E11

ICD-10 code description
Type 2 diabetes mellitus

Primary outcomes

1

Description
Fasting Blood Glucose

Timepoint
Before, at the first of study, daily for 3 days, once 2days for 4 days , on the days of 15, 30, 45 and 60

Method of measurement
patients with check fasting blood glucose with Glucometer at Home

Secondary outcomes

1

Description

Liver Function Test

Timepoint

Before, After, At the end of first month, At the end of second month

Method of measurement

Liver Function Tests

Intervention groups

1

Description

Intervention group: Two Capsule containing one fourth of the effective dose of four medicinal herbs

Category

Treatment - Drugs

2

Description

Control group: Two Capsule containing Rye bread powder, 1000 mg twice a day, morning and night after meals

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Baqiyatallah Hospital

Full name of responsible person

Khalil Mohamadzadeh

Street address

Baqiyatallah Hospital, Mullasadra Street, Vanak Square, Tehran

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Bagheiat-allah University of Medical Sciences

Full name of responsible person

Mohsen Saberi

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Grant name

Grant code / Reference number

Is the source of funding the same sponsor organization/entity?

Yes

Title of funding source

Bagheiat-allah University of Medical Sciences

Proportion provided by this source

100

Public or private sector

Public

Domestic or foreign origin

Domestic

Category of foreign source of funding

empty

Country of origin

Type of organization providing the funding

Academic

Person responsible for general inquiries

Contact

Name of organization / entity

Bagheiat-allah University of Medical Sciences

Full name of responsible person

Khalil Mohamadzadeh

Position

Internal Medicine Resident

Latest degree

Medical doctor

Other areas of specialty/work

Internal Medicine

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Other areas of specialty/work

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Undecided - It is not yet known if there will be a plan to make this available

Study Protocol

Undecided - It is not yet known if there will be a plan to make this available

Statistical Analysis Plan

Undecided - It is not yet known if there will be a plan to make this available

Informed Consent Form

Undecided - It is not yet known if there will be a plan to make this available

Clinical Study Report

Undecided - It is not yet known if there will be a plan to make this available

Analytic Code

Undecided - It is not yet known if there will be a plan to make this available

Data Dictionary

Undecided - It is not yet known if there will be a plan to make this available