

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

29 Jun 2026

### Investigating the effect of non-pharmacological treatment including diet adjustment and increased physical activity in the treatment of functional constipation

#### Protocol summary

##### Study aim

Investigating the effect of non-pharmacological treatment (diet adjustment and increased physical activity) in the treatment of functional constipation

##### Design

Clinical trial without control group (single group), phase 2 on 64 patients.

##### Settings and conduct

The present study is a clinical trial that will be conducted in Isfahan. Patients' demographic information is first collected by a questionnaire. The severity of constipation in patients at the beginning of the study is measured by a questionnaire based on the Rome III criteria. Patients will then be scheduled to adjust their diet and increase physical activity for 12 weeks. Again, the constipation questionnaires for all patients will be filled in weeks 2, 4, 8, and 12 after the intervention begins, and scoring will be done.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Having functional constipation based on Rome III criteria; Providing informed written consent to participate in the study. Non-inclusion criteria: Any history of chronic inflammatory disease or structural disease of the gastrointestinal tract; Any serious physical illness or illnesses such as inflammation or malignancy; Any physical problem that prevents from doing physical activity; Addiction to the drug or sleeping pills; History of chronic diseases such as diabetes.

##### Intervention groups

Patients will be scheduled to adjust their diet and increase physical activity for 12 weeks. Diet adjustment is done by designing a diet containing 3 to 6 grams of fiber and 2 glasses of water or other liquids other than tea on a daily basis and by a nutritionist, as well as according to the daily needs of each person. Activity levels are defined as half an hour of brisk walking daily, which increases heart rate by 50%.

#### Main outcome variables

Constipation severity score

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20200601047621N2**

Registration date: **2020-06-23, 1399/04/03**

Registration timing: **prospective**

Last update: **2020-06-23, 1399/04/03**

Update count: **0**

##### Registration date

2020-06-23, 1399/04/03

##### Registrant information

##### Name

Maryam Soheilipour

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 31 3729 4502

##### Email address

maryamsoheilip@med.mui.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2020-08-25, 1399/06/04

##### Expected recruitment end date

2020-10-24, 1399/08/03

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

**Trial completion date**  
empty

**Scientific title**  
Investigating the effect of non-pharmacological treatment including diet adjustment and increased physical activity in the treatment of functional constipation

**Public title**  
Non-pharmacological treatment of constipation

**Purpose**  
Treatment

**Inclusion/Exclusion criteria**  
**Inclusion criteria:**  
Having functional constipation based on Rome III criteria  
Free and informed written consent to participate in the study  
**Exclusion criteria:**  
Any history of chronic inflammatory disease or structural disease of the gastrointestinal tract  
Any serious physical problems or illness such as inflammation or malignancy  
Any physical problem that prevents from doing physical activity  
Drug or sleeping pills addiction  
History of chronic diseases such as diabetes

**Age**  
From **18 years** old to **50 years** old

**Gender**  
Both

**Phase**  
2

**Groups that have been masked**  
*No information*

**Sample size**  
Target sample size: **64**

**Randomization (investigator's opinion)**  
N/A

**Randomization description**  
**Blinding (investigator's opinion)**  
Not blinded

**Blinding description**  
**Placebo**  
Not used

**Assignment**  
Single

**Other design features**

## Secondary Ids

empty

## Ethics committees

1

**Ethics committee**  
**Name of ethics committee**  
Ethics committee of Isfahan University of Medical Sciences  
**Street address**  
No. 8, Hezar Jarib Ave., Daneshgh Blvd., Isfahan

**City**  
Isfahan  
**Province**  
Isfahan  
**Postal code**  
6719674255  
**Approval date**  
2020-05-23, 1399/03/03  
**Ethics committee reference number**  
IR.MUI.MED.REC.1399.106

## Health conditions studied

1

**Description of health condition studied**  
constipation  
**ICD-10 code**  
K59.0  
**ICD-10 code description**  
Constipation

## Primary outcomes

1

**Description**  
Functional constipation  
**Timepoint**  
Weeks 2, 4, 8 and 12 after the intervention begins  
**Method of measurement**  
Questionnaire for constipation

## Secondary outcomes

empty

## Intervention groups

1

**Description**  
Intervention group: The severity of constipation in patients at the beginning of the study is measured by a questionnaire based on Rome III criteria, then patients will be scheduled to adjust their diet and increase physical activity for 12 weeks. Diet adjustment is done by designing a diet containing 3 to 6 grams of fiber and 2 glasses of water or other liquids other than tea on a daily basis and by a nutritionist, as well as according to the daily needs of each person. The rate of activity is defined as half an hour of brisk walking daily, which increases the heart rate by 50%. Again, constipation severity questionnaire will be filled out for all patients at weeks 2, 4, 8, and 12 after the intervention begins.  
**Category**  
Treatment - Drugs

## Recruitment centers

## 1

### Recruitment center

**Name of recruitment center**

Khorshid Hospital

**Full name of responsible person**

Maryam Soheilipour

**Street address**

Khorshid hospital, Ostandari Blvd., Isfahan

**City**

Isfahan

**Province**

Isfahan

**Postal code**

6719674255

**Phone**

+98 31 3729 4225

**Email**

maryamsheilip@med.mui.ac.ir

## Sponsors / Funding sources

### 1

#### Sponsor

**Name of organization / entity**

Esfahan University of Medical Sciences

**Full name of responsible person**

Shaghayegh Haghjoo

**Street address**

No. 8, Hezar Jarib Ave., Daneshgh Blvd., Isfahan

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**Email**

haghjoo.sh@med.mui.ac.ir

**Grant name****Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

**Title of funding source**

Esfahan 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**

Esfahan University of Medical Sciences

**Full name of responsible person**

Maryam Soheilipour

**Position**

Assistant Professor

**Latest degree**

Subspecialist

**Other areas of specialty/work**

Internal Medicine

**Street address**

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## Person responsible for scientific inquiries

### Contact

**Name of organization / entity**

Esfahan University of Medical Sciences

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**Position**

Assistant Professor

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## Person responsible for updating data

### Contact

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Esfahan University of Medical Sciences

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**Phone**

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**Email**

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**Web page address****Sharing plan****Deidentified Individual Participant Data Set (IPD)**

Yes - There is a plan to make this available

**Study Protocol**

Yes - There is a plan to make this available

**Statistical Analysis Plan**

Yes - There is a plan to make this available

**Informed Consent Form**

Yes - There is a plan to make this available

**Clinical Study Report**

Yes - There is a plan to make this available

**Analytic Code**

Yes - There is a plan to make this available

**Data Dictionary**

Yes - There is a plan to make this available

**Title and more details about the data/document**

The information of this research will be available after deleting the names and details of the patients, through the research deputy of Isfahan University of Medical Sciences. These data include age, sex, duration of illness, and treatment information.

**When the data will become available and for how long**

Late 2020

**To whom data/document is available**

Health researchers

**Under which criteria data/document could be used**

In order to use the information in future research and under the permission of the Vice Chancellor for Research

**From where data/document is obtainable**

Information will be available through the Research Vice Chancellor of Isfahan University of Medical Sciences

**What processes are involved for a request to access data/document**

Apply online through the university website and issue a license within about 2 weeks and apply for data access

**Comments**