

Clinical Trial Protocol

Iranian Registry of Clinical Trials

20 Jun 2026

Effect Of Probiotic Supplementation On Some Digestive Problems, Stroke Patients

Protocol summary

Summary

Initially, the purpose and methods of implementation to the patients or their parents will be explained and Written consent to the inclusion criteria of the patients received. The GCS and some indicators such as waist size (the area around the navel) to compare the amount of flatulence in the beginning of the study And frequency of bowel movements (constipation at the beginning and end of the study was to compare the rate) in these patients are checked on arrival. Then based on the method used to assign patients randomly divided into two groups of 35. In addition to the daily diet of the test group 2 probiotic supplements containing Lactobacillus Zhry lactate fermentation biotech company, Bifidobacterium, Streptococcus and Fructose Oligosaccharides are 100 mg; Gavage every 12 hours with a solution for a given week And to control the food is just as routine. At the end of the first week back some indicators such as waist circumference and Tdaddfat stools are examined and measured. The end results will be compared with each other group.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2014111119911N1**

Registration date: **2015-01-06, 1393/10/16**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2015-01-06, 1393/10/16

Registrant information

Name

Sara Jahangiri

Name of organization / entity

Sabzevar University of Medical Sciences and Health Services

Country

Iran (Islamic Republic of)

Phone

+98 51 3604 6516

Email address

sarajahangiri448@yahoo.com

Recruitment status

Recruitment complete

Funding source

Sabzevar University of Medical Sciences

Expected recruitment start date

2014-08-23, 1393/06/01

Expected recruitment end date

2014-12-21, 1393/09/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Effect Of Probiotic Supplementation On Some Digestive Problems, Stroke Patients

Public title

Effect Of Probiotic Supplementation On Constipation and Bloating Stroke Patients

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: Patients aged 80-30; Patients with gastrointestinal problems (constipation and abdominal distension); No obvious bleeding in the upper gastrointestinal tract; Absence of associated diseases that cause constipation, such as diabetes or hypothyroidism; Total parenteral nutrition patients who

do not; Patients with GCS (5-15 have; Patients who have the same die; Hydration in patients who have identical conditions; Not receiving medications that cause constipation; Patients who do not exercise and your daily activities to help others do the work in terms of a level; Stroke patients admitted to the ICU, Taleghani Hospital, Mashhad. Exclusion criteria: The use of probiotics in the following refusal by the patient's physician to the patient; Parents patient's refusal to continue using probiotic supplements; Patients who do not tolerate gavage; Patients with systolic blood pressure lower than 100 millimeters of mercury; The patient died during the study.

Age

From **30 years** old to **80 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **70**

Randomization (investigator's opinion)

Randomized

Randomization description**Blinding (investigator's opinion)**

Not blinded

Blinding description**Placebo**

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

Ethics committees**1****Ethics committee****Name of ethics committee**

Sabzevar University of Medical Sciences and Health Services

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Building number two, Vice Chancellor for Education, University of Sabzevar

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9613873136

Approval date

2014-07-05, 1393/04/14

Ethics committee reference number

medsab.Rec.93.15

Health conditions studied**1****Description of health condition studied**

Stroke

ICD-10 code

I64

ICD-10 code description

Stroke, not specified as haemorrhage or infarction

Primary outcomes**1****Description**

Abdominal circumference

Timepoint

A week before the intervention and after the intervention began.

Method of measurement

Using a tape measure in centimeter.

2**Description**

The frequency of bowel movements

Timepoint

A week before the intervention and after the intervention began.

Method of measurement

Based on the registration number of daily excretion in patient chart

Secondary outcomes**1****Description**

Side Effect

Timepoint

During the intervention and a week after the intervention

Method of measurement

Observation and scrutiny of the patient's general condition

Intervention groups**1****Description**

The control group was routinely given food only / no action

Category

Treatment - Drugs

2**Description**

The intervention group consumed (lactate Zhry Capsules, 500 mg, twice a day after meal for a week)

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Taleghani Hospital, Mashhad

Full name of responsible person

Sara Jahangiri

Street address

Taleghani, Tee Ferdowsi Asian Highway, Mashhad

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Sabzevar University of Medical Sciences

Full name of responsible person

Musa Al-Reza Taday'oun far

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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

Sabzevar University of Medical Sciences

Proportion provided by this source

100

Public or private sector

empty

Domestic or foreign origin

empty

Category of foreign source of funding

empty

Country of origin

Type of organization providing the funding

empty

Person responsible for general inquiries

Contact

Name of organization / entity

Sabzevar University of Medical Sciences and Health Services

Full name of responsible person

Sara Jahangiri

Position

Nursing Graduate Student Sabzevar University of
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Other areas of specialty/work

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Sharing plan**Deidentified Individual Participant Data Set (IPD)***empty***Study Protocol***empty***Statistical Analysis Plan***empty***Informed Consent Form***empty***Clinical Study Report***empty***Analytic Code***empty***Data Dictionary***empty*