

Clinical Trial Protocol

Iranian Registry of Clinical Trials

13 Jun 2026

Evaluate the effect of probiotics on the improvement of autoimmune inflammation in patients with systemic lupus erythematosus

Protocol summary

Study aim

Evaluation of probiotic administration on immune responses and clinical manifestations in patients with systemic lupus erythematosus

Design

Clinical trial with control group, community-based, with parallel groups, thruple blind, randomized

Settings and conduct

Before and 4 months after the intervention, 5 cc of blood is taken from the brachial vein of patients for examinations, real time PCR and preparation of serum for ELISA test. In monthly visits, in addition to the activity of the disease, a checklist of the number and dose of drugs used by patients in both groups is recorded to be compared at the end of the study and the confusing points of treatment changes are removed from the study.

Participants/Inclusion and exclusion criteria

Inclusion criteria: All patients with systemic lupus erythematosus between the ages of 18 to 45 years living in Mashhad and surrounding cities with a SLEDAI score between 4.5 to 10. Be willing to cooperate in the project. Exclusion criteria: Patients who are not satisfied to continue cooperation.

Intervention groups

Patients are divided into two equal groups (23 people in each group) using block block randomization method and in one group, in addition to common treatments for lupus, oral probiotic Lact 2 Plus is prescribed, and in the other group, in addition to common treatments, A placebo is prescribed.

Main outcome variables

Evaluation of IL17 protein levels, Evaluation of FOXP3, T-bet, GATA-3 and ROR γ t transcription factor gene expression by SYBR Green real-time RT-PCR at the beginning of the process, four months after the course

General information

Reason for update

Acronym

SLE(systemic lupus erythematosus)

IRCT registration information

IRCT registration number: **IRCT20201019049073N1**

Registration date: **2021-01-08, 1399/10/19**

Registration timing: **registered_while_recruiting**

Last update: **2021-01-08, 1399/10/19**

Update count: **0**

Registration date

2021-01-08, 1399/10/19

Registrant information

Name

Arezoo Faridzadeh

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 51 3802 2229

Email address

faridzadeha961@mums.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2020-10-28, 1399/08/07

Expected recruitment end date

2022-12-10, 1401/09/19

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluate the effect of probiotics on the improvement of autoimmune inflammation in patients with systemic lupus erythematosus

Public title

Evaluate of effect probiotic on treatment of systemic lupus erythematosus

Purpose

Basic science

Inclusion/Exclusion criteria

Inclusion criteria:

All patients with SLE have a SLEDAI score of 4.5 to 10. Patients should be between 18 and 45 years old. Patients are willing to cooperate in the project.

Exclusion criteria:

Patients will be excluded from the study if there is a change in the dose or number of medications taken by patients.

Age

From **18 years** old to **45 years** old

Gender

Both

Phase

3

Groups that have been masked

- Participant
- Care provider
- Investigator

Sample size

Target sample size: **46**

Randomization (investigator's opinion)

Randomized

Randomization description

Method of block randomization Unit of randomization individual Tools used in randomization computer software The type of randomization in this study is simple random sampling. To select the participants in this study, 50 patients from whom the diagnosis of lupus has been confirmed are randomly included in the study. Then 25 patients are randomly divided into probiotic group and 25 patients into placebo group.

Blinding (investigator's opinion)

Triple blinded

Blinding description

Participants will receive the same drug or placebo. The main researcher is unaware of the effect of the drug on improving quality of life and effective immune responses.

Placebo

Used

Assignment

Parallel

Other design features

Our review of past studies showed that no similar studies have been conducted in this area. Therefore, this study can provide new results on the effect of probiotics on systemic lupus erythematosus disease.

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Mshhad University of medical Sciences

Street address

Ferdowsi University of Medical Sciences , Azadi Sq, Mashhad-Iran

City

mashhad

Province

Razavi Khorasan

Postal code

9177948564

Approval date

2020-12-28, 1399/10/08

Ethics committee reference number

IR.MUMS.MEDICAL.REC.1399.577

Health conditions studied

1

Description of health condition studied

SLE(Systemic lupus erythematosus)

ICD-10 code

ICD-10 code description

kinds of autoimmune diseases

Primary outcomes

1

Description

IL17, Evaluation of FOXP3, T-bet, GATA-3 and ROR γ t transcription factor gene expression by SYBR Green real-time RT-PCR method

Timepoint

Before intervention, 4 months after intervention

Method of measurement

Elisa test, SYBR Green real-time RT-PCR

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: In addition to common therapies, one oral probiotic FamiLact 2plus daily includes the strains of Lactobacillus acidophilus, Lactobacillus casei, Lactobacillus rhamnosus, Lactobacillus salivarius, Lactobacillus reuteri, Bifidobacterium lactis, Bifidobacterium longum, Bifidobacterium bifidum. The duration of the intervention is 4 months. FamiLact 2Plus is from Zist Takhmir Company, made in Iran.

Category

Treatment - Drugs

2**Description**

Control group: In addition to common treatments, one placebo is given daily. The duration of the intervention is 4 months. The placebo is from Zist Takhmir Company, made in Iran.

Category

Placebo

Recruitment centers**1****Recruitment center****Name of recruitment center**

Emam Reza hospital

Full name of responsible person

Doctor Seyedeh Zahra Mirfeizi

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Rheumatology Research Center of Imam Reza, mashhad city Hospital

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Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

Mashhad University of Medical Sciences

Full name of responsible person

Doctor Mohsen Tafaghodi

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faculty of medical, Mashhad Ferdowsi University, Ferdowsi square, Mashhad city

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Grant name

Mashhad University of Medical Sciences

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

Yes

Title of funding source

Mashhad 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**

Mashhad University of Medical Sciences

Full name of responsible person

Arezoo Faridzadeh

Position

MD-Ph.D candidate

Latest degree

Medical doctor

Other areas of specialty/work

Immunology

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Person responsible for scientific inquiries**Contact****Name of organization / entity**

Mashhad University of Medical Sciences

Full name of responsible person

Arezoo Faridzadeh

Position

medical doctor

Latest degree

Medical doctor

Other areas of specialty/work

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Person responsible for updating data**Contact****Name of organization / entity**

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Full name of responsible person

Arezoo faridzadeh

Position

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Latest degree

Medical doctor

Other areas of specialty/work

Immunology

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Mashhad city**City**

mashhad

Province**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