

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

03 Jun 2026

### Preparation & evaluation of Topical Rosuvastatin & Melatonin vs placebo in Patients with Mild to Moderate Plaque Psoriasis! A Randomized Double-blinded Clinical Trial

#### Protocol summary

##### Study aim

Preparation of melatonin and rosuvastatin creams and comparing their clinical effects on psoriasis patients

##### Design

51 patients aged 18 years and older, with a clinical diagnosis that fills the morphological criteria of stable psoriasis plaque, enter the study with plaques size 100 cm<sup>2</sup> or less. If eligible, patients will be randomized according to a table of random numbers to receive one of the interventions of melatonin or rosuvastatin or placebo creams (17 in each group). Patients, clinical evaluators, and physicians will be blinded by the type of intervention. Before entering the study, all patients undergo a four-week wash-out course of any topical product associated with psoriasis. In addition, systemic therapies, including phototherapy, will be discontinued four weeks before the study. Patients are instructed to use their group's topical cream twice a day and not to use any other moisturizers when using the cream. The amount of cream used in patients is the same as the fingertip unit (FTU), twice a day, which is one use at bedtime. After treatment, patients are monitored for plaque recurrence for 8 weeks.

##### Settings and conduct

Yazd Rheumatology Specialist Office

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Patients were 18 years or older; less than 10% body surface area  
Exclusion criteria: Pregnancy; sensitivity to the components of the cream are the conditions

##### Intervention groups

One group is the Melatonin cream recipient, the other is Rosuvastatin cream, and the last is the placebo cream recipient.

##### Main outcome variables

Psoriasis Area Severity Index (PASI) : Dermatological Sum Score (DSS) : Dermatology Life Quality Index (DLQI)

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20190810044500N16**

Registration date: **2021-06-13, 1400/03/23**

Registration timing: **registered\_while\_recruiting**

Last update: **2021-06-13, 1400/03/23**

Update count: **0**

##### Registration date

2021-06-13, 1400/03/23

##### Registrant information

##### Name

Fatemeh Saghafi

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 35 3820 3419

##### Email address

f.saghafi@ssu.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2021-04-21, 1400/02/01

##### Expected recruitment end date

2021-07-23, 1400/05/01

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

## Scientific title

Preparation & evaluation of Topical Rosuvastatin & Melatonin vs placebo in Patients with Mild to Moderate Plaque Psoriasis: A Randomized Double-blinded Clinical Trial

## Public title

Preparation & evaluation of Topical Rosuvastatin & Melatonin vs placebo in Patients with Mild to Moderate Plaque Psoriasis

## Purpose

Treatment

## Inclusion/Exclusion criteria

### Inclusion criteria:

Patients 18 years and older Less than 10% body surface area (BSA)

### Exclusion criteria:

Pregnancy Sensitivity to cream components

## Age

From **18 years** old

## Gender

Both

## Phase

2

## Groups that have been masked

- Participant
- Care provider
- Outcome assessor

## Sample size

Target sample size: **51**

## Randomization (investigator's opinion)

Randomized

## Randomization description

First, using Random allocation software (version 1.0), we generate a random sequence by a simple random allocation method. In this table, we specify from 1 to 51 and each number is assigned to an intervention group (A or B or C). The first eligible person is referred to as number 1, the second person as number 2, and so on up to 51 patients. To be blind to random allocation, patients are placed in one of the intervention groups (A or B or C) according to the table by a third person who is unaware of the interventions.

## Blinding (investigator's opinion)

Double blinded

## Blinding description

Patients, clinical evaluators, and physicians will be blinded by the type of intervention.

## Placebo

Used

## Assignment

Parallel

## Other design features

## Secondary Ids

empty

## Ethics committees

## 1

### Ethics committee

#### Name of ethics committee

Ethics Committee of Shahid Sadoughi University Medical Sciences

#### Street address

Shahid Sadoughi university of medical sciences, Professor Hesabi Blvd, Yazd, Iran

#### City

yazd

#### Province

Yazd

#### Postal code

8915173143

#### Approval date

2020-11-09, 1399/08/19

#### Ethics committee reference number

IR.SSU.MEDICINE.REC.1399.221

## Health conditions studied

## 1

### Description of health condition studied

Psoriasis

### ICD-10 code

L40

### ICD-10 code description

Psoriasis

## Primary outcomes

## 1

### Description

Psoriasis Area Severity Index

### Timepoint

Weeks 0, 4, 8

### Method of measurement

Score 0\_72

## Secondary outcomes

## 1

### Description

Dermatological Sum Score (DSS)

### Timepoint

Weeks 0,4,8

### Method of measurement

Based on a scale of 4 points (0: absent,1: mild, 2: moderate, 3: severe)

## 2

### Description

Dermatology Life Quality Index (DLQI)

### Timepoint

weeks 0,4,8

### Method of measurement

Based on the questionnaire

## Intervention groups

### 1

#### Description

Intervention group: Patients will use 5% melatonin cream twice daily for 8 weeks

#### Category

Treatment - Drugs

### 2

#### Description

Intervention group: Patients will use 5% rosuvastatin cream twice daily for 8 weeks

#### Category

Treatment - Drugs

### 3

#### Description

Control group: Patients will use placebo cream twice daily for 8 weeks

#### Category

Placebo

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Yazd Rheumatology Specialist Office

##### Full name of responsible person

Dr. Hossein Soleimani Salehabadi

##### Street address

Taleghani St., Mojahedin Alley, after Razi Doctors Building

##### City

yazd

##### Province

Yazd

##### Postal code

8916978477

##### Phone

+98 35 3726 2268

##### Email

h.soleimani@ssu.ac.ir

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Yazd University of Medical Sciences

##### Full name of responsible person

Masoud Mirzaei

##### Street address

Bahonar Square

##### City

yazd

#### Province

Yazd

#### Postal code

8916978477

#### Phone

+98 35 3146 2056

#### Email

mmirzaei@ssu.ac.ir

#### Grant name

#### Grant code / Reference number

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

Yes

#### Title of funding source

Yazd 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

Yazd University of Medical Sciences

##### Full name of responsible person

Fatemeh Saghafi

##### Position

Assistant professor

##### Latest degree

Ph.D.

##### Other areas of specialty/work

Medical Pharmacy

##### Street address

Professor Hesabi Blvd., Yazd Province, Yazd, Iran

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F.saghafi@ssu.ac.ir

## Person responsible for scientific inquiries

#### Contact

##### Name of organization / entity

Yazd University of Medical Sciences

##### Full name of responsible person

fatemeh saghafi

##### Position

Associate professor

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Medical Pharmacy

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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