

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

09 Jun 2026

### Evaluation of protective effect of Royal Jelly on Serum Oxidative Stress markers and Interleukin-1Beta of Patients with Multiple Sclerosis.

#### Protocol summary

##### Study aim

The evaluation of the protective effect of Royal Jelly capsule consumption on serum oxidative stress markers and level of Interleukin-1 beta of patients with Multiple sclerosis.

##### Design

Dividing 60 patients (random and double-blind) into intervention and control groups, phase 3 trial, 45 half-gram capsules of Royal Jelly or placebo to the patient (45 days), blood sampling at the beginning and after the 45th day, assessment (MDA, SOD, GPx, IL-1) and EDSS after the intervention.

##### Settings and conduct

In Dezful University of Medical Sciences: Determining the EDSS level of patients by a neurologist, dividing 60 patients (random and double-blind) into two intervention and control groups, 45 half-gram capsules of Royal Jelly or placebo per patient (daily consumption of one), then blood sampling at the beginning and after the day 45th, measurement of MDA, SOD, GPx, catalase and IL-1, and EDSS after the intervention.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria of participants:: Patients' subtype of M.S should be the "relapsing-remitting MS" (RRMS). The patient's age should be 20- 45 years old and all of them treated with interferon for at least 6 months. conditions for not entering the study: The Patients with anemia, chronic heart disease, lung disease, diabetic Mellitus, autoimmune diseases such as rheumatoid arthritis, SLE, liver diseases, and a history of asthma and allergies. Patients are treated with corticosteroids or ACTH drugs, or consumption of supplements.

##### Intervention groups

Dividing 60 patients (random and double-blind) into intervention and control groups, 45 half-gram capsules of Royal Jelly or placebo to each patient (daily consumption of one).

##### Main outcome variables

Decreased MDA, IL1: Increased SOD, GPx, CAT: improved

EDSS in intervention groups.

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20220116053732N1**

Registration date: **2022-05-08, 1401/02/18**

Registration timing: **retrospective**

Last update: **2022-05-08, 1401/02/18**

Update count: **0**

##### Registration date

2022-05-08, 1401/02/18

##### Registrant information

##### Name

Zahra Eslamifar

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 61 4242 9531

##### Email address

eslamifar.z@gmail.com

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2022-03-06, 1400/12/15

##### Expected recruitment end date

2022-04-04, 1401/01/15

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

### Scientific title

Evaluation of protective effect of Royal Jelly on Serum Oxidative Stress markers and Interleukin-1Beta of Patients with Multiple Sclerosis.

### Public title

Evaluation of protective effect of Royal Jelly on Serum Oxidative Stress markers and Interleukin-1 Beta of Patients with Multiple Sclerosis.

### Purpose

Treatment

### Inclusion/Exclusion criteria

#### Inclusion criteria:

Age of patients should be 20-45 The sub type of M.S should be "Relapsing-Remitting MS" (RRMS). The diagnosis of "RRMS" is confirmed by a neurologist based on neurological examination and imaging. Patients should be treated with "Interferon" for at least 6 months.

#### Exclusion criteria:

Patients with anemia, Chronic Heart Disease, lung disease, Diabetic mellitus, Autoimmune diseases such as Rheumatoid Arthritis, SLE, ... Patients treated with corticosteroids or ACTH. Use of any supplements by patients. History of asthma and allergies in the patient. Occurrence of allergic symptoms during the study Occurrence of a "MS" attack during the study period. The patient' s refuse to continue study.

### Age

From **20 years** old to **45 years** old

### Gender

Both

### Phase

3

### Groups that have been masked

- Participant
- Care provider
- Investigator
- Outcome assessor
- Data analyser

### Sample size

Target sample size: **60**

### Randomization (investigator's opinion)

Randomized

### Randomization description

Permuted block randomization method will be used and randomization unit is individual. According to the sample size of 60 and two treatment groups, 10 blocks with volume 6 are prepared, for this, a list of six combinations of Royal Jelly and placebo is prepared and randomly allocated each compound to one block. In order to reduce the prediction rate, the therapist is not aware of the permutation block method and especially the size of the block.

### Blinding (investigator's opinion)

Double blinded

### Blinding description

Patients were invited to participate in the study and after obtaining informed consent, they were blindly classified into control or intervention groups. The clinician,

researcher, outcome evaluator, and data analyst all play their role blindly.

### Placebo

Used

### Assignment

Other

### Other design features

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Ethics committee of Dezful University of Medical Sciences

##### Street address

Azadegan boulevard, deputy of training, school of medicine, Dezful, Iran.

##### City

Dezful

##### Province

Khuzestan

##### Postal code

6461665145

#### Approval date

2020-09-28, 1399/07/07

#### Ethics committee reference number

IR.DUMS.REC.1399.038

## Health conditions studied

### 1

#### Description of health condition studied

MS disease

#### ICD-10 code

G35

#### ICD-10 code description

Multiple sclerosis (of):NOS brain stem cord disseminated generalized

## Primary outcomes

### 1

#### Description

The rate of change of oxidative stress state

#### Timepoint

Evaluation of "oxidative stress status" at the beginning of the study and on day 30 after the use of Royal Jelly

#### Method of measurement

The assay of Malondialdehyde , Superoxide dismutase, Catalase, Nitric oxide

## Secondary outcomes

## 1

### Description

Expanded Disability Status Scale

### Timepoint

Study of "Expanded Disability Status Scale" at the beginning of the study and on day 30 after the use of Royal Jelly

### Method of measurement

Scoring from zero (best condition) to 10 (death due to MS) according to the neurologist

## Intervention groups

## 1

### Description

The intervention group included people with MS who on the first day of the study and on the 50th day of the study "after swallowing a capsule containing 350 mg of royal jelly lyophilized daily", "Extensive disability status scale" and blood sampling to assess the status of "oxidative stress "It will be measured. The study of oxidative stress is done by measuring factors such as malondialdehyde, catalase, superoxide dismutase and nitric oxide.

### Category

Treatment - Drugs

## 2

### Description

Control group included people with MS who on the first day of the study and on the 50th day of the study "after swallowing a capsule containing 350 mg of wheat flour daily", "Extensive disability status scale" and blood sampling to assess the status of "oxidative stress "It will be measured. The study of oxidative stress is done by measuring factors such as malondialdehyde, catalase, superoxide dismutase and nitric oxide.

### Category

Placebo

## Recruitment centers

## 1

### Recruitment center

#### Name of recruitment center

Neshat Clinic 2

#### Full name of responsible person

Dr. Mohammad Reza Zandi

#### Street address

No. 1, Neshat St., Neshat Clinic 2, Dezful Town

#### City

Dezful

#### Province

Khuzestan

#### Postal code

6461795759

#### Phone

+98 61 4227 2201

### Email

eslamifar.z@gmail.com

## Sponsors / Funding sources

## 1

### Sponsor

#### Name of organization / entity

Dezful University of Medical Sciences

#### Full name of responsible person

Dr. Alireza Khosropanah

#### Street address

No. 107, School of Paramedical, Dezful University of Medical Sciences, next to the Traffic Department, Azadegan Blvd, Dezful.

#### City

Dezful

#### Province

Khuzestan

#### Postal code

6461665145

#### Phone

+98 61 4242 9531

#### Email

eslamifar.z@gmail.com

### Grant name

### Grant code / Reference number

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

Yes

### Title of funding source

Dezful 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

Dezful University of Medical Sciences

#### Full name of responsible person

Dr. Susan Sabbagh

#### Position

Faculty member

#### Latest degree

Ph.D.

#### Other areas of specialty/work

Anatomy

#### Street address

No. 37, Fajr St., Shahid Beheshti St., Dezful

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**Email**

sabbaghsusan@yahoo.com

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

Dezfoul University of Medical Sciences

**Full name of responsible person**

Dr. Susan Sabbagh

**Position**

Faculty member

**Latest degree**

Ph.D.

**Other areas of specialty/work**

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Dezfoul University of Medical Sciences

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**Sharing plan****Deidentified Individual Participant Data Set (IPD)**

Yes - There is a plan to make this available

**Study Protocol**

No - There is not a plan to make this available

**Statistical Analysis Plan**

No - There is not a plan to make this available

**Informed Consent Form**

No - There is not a plan to make this available

**Clinical Study Report**

No - There is not a plan to make this available

**Analytic Code**

No - There is not a plan to make this available

**Data Dictionary**

No - There is not a plan to make this available

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

No

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

No

**To whom data/document is available**

No

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

No

**From where data/document is obtainable**

No

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

No

**Comments**