

Clinical Trial Protocol

Iranian Registry of Clinical Trials

09 Jun 2026

The effect of topical chamomile herbal product on the symptoms of patients with leg cramps in comparison with standard treatment: a double-blind clinical trial

Protocol summary

Study aim

Investigating the effect of chamomile topical product on the severity of symptoms of people with muscle cramps

Design

This study is a double-blind randomized clinical trial study with a control group and a parallel design, which is conducted in the third phase of the trial on 64 people with muscle cramps in the leg area.

Settings and conduct

Every person who meets the criteria for entering the study is evaluated by an Iranian medical specialist using the VAS tool in the first visit (before the start of the study) and subsequent visits (after the start of the intervention). They are evaluated daily and up to a score of less than four in the VAS tool. Referrals to the health school of Babol University of Medical Sciences, Rouhani Hospital Clinic are examined. Chamomile product and diclofenac gel in a single form, with the same label and the same smell and color (with the addition of color and essential oil) by a pharmaceutical specialist in the herbal laboratory. Medicines will be prepared by the Faculty of Iranian Medicine in Babol. The only person who knows the nature of drugs is the pharmacist who makes the drug.

Participants/Inclusion and exclusion criteria

Inclusion criteria: 1. Men and women aged 20 to 60 years
2. Presence of pain with a vas score greater than six
Exclusion: 1. Lack of full assessment to monitor the treatment process
2. Lack of satisfaction during treatment
3. The presence of underlying diseases related to cramps

Intervention groups

1. The intervention group includes people with muscle cramps in the leg area who receive topical chamomile products.
2. The comparison group includes people suffering from muscle cramps in the leg area, receiving 1% diclofenac gel.

Main outcome variables

Treatment duration, pain index

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20200105046009N7**

Registration date: **2023-01-09, 1401/10/19**

Registration timing: **prospective**

Last update: **2023-01-09, 1401/10/19**

Update count: **0**

Registration date

2023-01-09, 1401/10/19

Registrant information

Name

Seyyed Ali Mozaffarpur

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 11 3219 4728

Email address

dr.mozaffarpur@mubabol.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2023-01-20, 1401/10/30

Expected recruitment end date

2023-06-21, 1402/03/31

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date
empty

Scientific title
The effect of topical chamomile herbal product on the symptoms of patients with leg cramps in comparison with standard treatment: a double-blind clinical trial

Public title
The effect of topical chamomile herbal product on the symptoms of patients with muscle cramps

Purpose
Treatment

Inclusion/Exclusion criteria
Inclusion criteria:
Men and women aged 20 to 60 years Presence of muscle cramp in the gastrocnemius muscle The presence of pain with a vas score greater than six
Exclusion criteria:
Lack of complete evaluation to monitor the treatment process Lack of satisfaction during treatment The presence of underlying diseases related to cramps Using other painkillers Performing procedures that lead to the improvement of muscle cramps History of recurrent cramps without clear cause

Age
From **20 years** old to **60 years** old

Gender
Both

Phase
3

Groups that have been masked

- Participant
- Care provider
- Investigator
- Outcome assessor

Sample size
Target sample size: **64**

Randomization (investigator's opinion)
Randomized

Randomization description
Randomization will be done using the permuted block randomization method. The size of the blocks is 4 and the ratio of drug to control (diclofenac gel) in each group is equal to (1:1). In each block, random permutations of 4 combinations of chamomile formulation and diclofenac gel are considered. Random sequence generation will be done by the design methodologist. In order to hide the treatment process, random codes are written on the boxes containing medicine (chamomile product/diclofenac) and after assigning the medicine to the patients, the corresponding code is recorded on the patients' files. The opening of the codes will be done after the end of the study. In case of side effects, the code of that medicine will be opened.

Blinding (investigator's opinion)
Double blinded

Blinding description
Chamomile product and diclofenac gel in a single form, with the same label and the same smell and color (with

the addition of color and essential oil) will be prepared by a pharmaceutical specialist in the medicinal plant laboratory of the Faculty of Iranian Medicine in Babol. The only person who knows the nature of drugs is the pharmacist who makes the drug.

Placebo
Not used

Assignment
Parallel

Other design features

Secondary Ids
empty

Ethics committees

1

Ethics committee

Name of ethics committee
Ethics committee of Babol University of Medical Sciences

Street address
Babol University of Medical sciences, Sargord Ghasemi avenue, Babol

City
Babol

Province
Mazandaran

Postal code
4718647745

Approval date
2023-01-02, 1401/10/12

Ethics committee reference number
IR.MUBABOL.HRI.REC.1401.205

Health conditions studied

1

Description of health condition studied
muscle cramp

ICD-10 code
R25.2

ICD-10 code description
Cramp and spasm

Primary outcomes

1

Description
pain index

Timepoint
The VAS measurement is measured before the intervention and on even days after the start of the intervention, and the treatment continues until the pain index reaches 4 or a change of 1.7 based on the MCID.

Method of measurement
Pain intensity measurement using VAS tool

2

Description

Duration of treatment

Timepoint

VAS is measured before the intervention and on even days after the start of the intervention, and the examination of the pain index continues until the day it reaches the number 4 or the change rate of 1.7 based on the MCID.

Method of measurement

Pain intensity measurement using VAS tool

Secondary outcomes

1

Description

redness

Timepoint

Measurements are taken before the intervention and on even days after the start of the intervention, and the treatment continues until the pain index reaches 4 or a change of 1.7 based on the MCID.

Method of measurement

By patient report or evaluator observation

2

Description

Itching

Timepoint

Measurements are taken before the intervention and on even days after the start of the intervention, and the treatment continues until the pain index reaches 4 or a change of 1.7 based on the MCID.

Method of measurement

By patient report or evaluator observation

3

Description

swelling

Timepoint

Measurements are taken before the intervention and on even days after the start of the intervention, and the treatment continues until the pain index reaches 4 or a change of 1.7 based on the MCID.

Method of measurement

By patient report or evaluator observation

Intervention groups

1

Description

Chamomile topical product (combination of chamomile oil and wax made in the laboratory of the Iranian Faculty of Medicine) in the amount of one knuckle every eight hours is used topically on the leg muscles and the VAS index, before the intervention and on even days after. The start of the intervention is measured and the treatment continues until the pain index reaches 4 or a

change of 1.7 based on the MCID.

Category

Treatment - Drugs

2

Description

1% diclofenac gel of Soban company is used topically on the calf muscles in the amount of one knuckle every eight hours, and the VAS index is measured before the intervention and on even days after the intervention, and the treatment is continued until the day the pain index is reached. Continued to number 4 or a rate of change of 1.7 based on MCID

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Roohani Hospital

Full name of responsible person

Seyyed Ali Mozaffarpur

Street address

Roohani Hospital, Babol University of Medical Sciences, Ganjafrooz Ave, Babol

City

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Postal code

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Phone

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Email

Seyyedali1357@gmail.com

2

Recruitment center

Name of recruitment center

Health College of Iranian Medicine, Babol University of Medical Sciences

Full name of responsible person

Seyyed Ali Mozaffarpur

Street address

Health College of Iranian Medicine,, , Babol University of Medical Sciences, Shahid Keshvari Square, Babol

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Email

Seyyedali1357@gmail.com

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Babol University of Medical Sciences

Full name of responsible person

Mehdi Rajabnia

Street address

Ganjafrooz Street

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Province

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4717647745

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Email

seyyedali1357@gmail.com

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

Yes

Title of funding source

Babol 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

Babol University of Medical Sciences

Full name of responsible person

Seyyed Ali Mozaffarpur

Position

Associate Professor

Latest degree

Ph.D.

Other areas of specialty/work

Traditional Medicine

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+98 11 3219 4728

Email

dr.mozaffarpur@mubabol.ac.ir

Person responsible for scientific inquiries

Contact

Name of organization / entity

Babol University of Medical Sciences

Full name of responsible person

Seyyed Ali Mozaffarpur

Position

Associate Professor

Latest degree

Ph.D.

Other areas of specialty/work

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Person responsible for updating data

Contact

Name of organization / entity

Babol University of Medical Sciences

Full name of responsible person

Seyyed Ali Mozaffarpur

Position

Associate Professor

Latest degree

Ph.D.

Other areas of specialty/work

Traditional Medicine

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Phone

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Email

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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

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

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

Title and more details about the data/document

-

When the data will become available and for how long

-

To whom data/document is available

-

Under which criteria data/document could be used

-

From where data/document is obtainable

-

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

-

Comments