

Clinical Trial Protocol

Iranian Registry of Clinical Trials

12 Jun 2026

The efficacy of *Satoria hortensis* essential oil on pain caused by arteriovenous fistula puncture of patients undergoing hemodialysis: A randomized double-blinded placebo-controlled clinical trial

Protocol summary

Study aim

Reduction of pain caused by arteriovenous fistula needling in dialysis patients using a natural and herbal method. By examining the effect of this essential oil on reducing pain and improving the condition of patients, this research aims to improve the existing treatment methods and introduce new natural treatments with minimal complication methods to manage acute pain.

Design

Randomized double-blind crossover controlled clinical trial on 30 patients and Excel software will be used for randomization.

Settings and conduct

Patients were referred to the dialysis centers of Namazi and Shahid Faqihi hospitals of Shiraz University of Medical Sciences.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Male and female patients with chronic kidney problems between 18 and 70 years old, dialysis at least 2 times a week, a history of dialysis through arteriovenous vessels for at least 3 months, and full consciousness and signing a written informed consent form. Exclusion criteria: Allergy to herbal products, skin disease and injury to the study site, use of oral analgesics in the past 6 hours or topical Arteriovenoussite pain medication in the past 24 hours, diabetic neuropathy or other types of neuropathy, failure of the initial attempt to achieve Arteriovenous vessels.

Intervention groups

For each patient, one tip of finger unit (TUF) of the gel (gel containing essential oil and placebo) will be applied within a radius of 2 cm from the entry point of the arteriovenous fistula (AVF). After ten minutes, an experienced nurse will puncture the AVF.

Main outcome variables

Demographic characteristics such as age, gender, educational status, and number of dialyses per week are

recorded. The main outcome measure is the amount of pain during AVF needling, measured using the video analog scale (AVS). This scale is known as a valid and reliable tool.

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20121031011341N12**

Registration date: **2024-01-03, 1402/10/13**

Registration timing: **prospective**

Last update: **2024-01-03, 1402/10/13**

Update count: **0**

Registration date

2024-01-03, 1402/10/13

Registrant information

Name

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Name of organization / entity

Fasa University of Medical Sciences

Country

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2024-01-15, 1402/10/25

Expected recruitment end date

2024-02-18, 1402/11/29

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The efficacy of Satoria hortensis essential oil on pain caused by arteriovenous fistula puncture of patients undergoing hemodialysis: A randomized double-blinded placebo-controlled clinical trial

Public title

Investigating the effect of Satoria hortensis on hemodialysis needling pain

Purpose

Supportive

Inclusion/Exclusion criteria**Inclusion criteria:**

Patients with chronic kidney disease Twice-weekly dialysis History of dialysis through arteriovenous fistula for a minimum of 3 months Full consciousness and sign a written consent for participation in the study

Exclusion criteria:

Allergy to herbal products Skin injuries to the site of arteriovenous fistula Consumption of oral sedatives in the past 6 hours or Local (arteriovenous fistula) pain medication in the past 24 hours Diabetic or other types of neuropathy Failure of the initial attempt to reach the arteriovenous fistula

Age

From **18 years** old to **70 years** old

Gender

Both

Phase

2-3

Groups that have been masked

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

Sample size

Target sample size: **30**

More than 1 sample in each individual

Number of samples in each individual: **2**

Placebo and Satoria hortensis gel

Randomization (investigator's opinion)

Randomized

Randomization description

To assign people to groups A and B, we will use the block of four method. The randomization list will be generated by a biostatistician using Excel software, and a third party will determine which groups are in treatment and which are placebo by throwing a coin.

Blinding (investigator's opinion)

Double blinded

Blinding description

Using similar non-transparent containers with consecutive numbering will guarantee the lack of

information about the allocation to groups, which will be considered. Both types of gel (gel containing essential oil and placebo gel) will be the same in terms of physical characteristics (including consistency, color, and smell of the product). In addition, they will be packed in containers of the same shape and size. It should be noted that each container has a code that only the person who produced it knows about its contents, so patients, nurses, doctors, and researchers are unaware of the drugs used.

Placebo

Used

Assignment

Crossover

Other design features**Secondary Ids**

empty

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

Ethics Committee of Shiraz University of Medical Science

Street address

Zand Avenue

City

Shiraz

Province

Fars

Postal code

۱۴۳۳۶ - ۷۱۳۴۸

Approval date

2023-12-12, 1402/09/21

Ethics committee reference number

IR.SUMS.MED.REC.1402.400

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

Hemodialysis

ICD-10 code

Z49.0

ICD-10 code description

Preparatory care for renal dialysis

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

Pain

Timepoint

immediately after needle insertion

Method of measurement

analog video scale (AVS)

Secondary outcomes

empty

Intervention groups

1

Description

Satoria hortensis essential oil will be applied on the insertion site of hemodialysis needles and remained for 10 minutes.

Category

Treatment - Drugs

2

Description

Control group: In placebo method, the gel of placebo would be applied on the insertion site of hemodialysis needles and remained for 10 minutes

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Shahid Faghihi Hospital

Full name of responsible person

Aida Iraj

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2

Recruitment center

Name of recruitment center

Namazi hospital

Full name of responsible person

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Sponsors / Funding sources

1

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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

Shiraz 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

Shiraz University of Medical Sciences

Full name of responsible person

Mohammad Hashem Hashempur

Position

Assistant Professor of Shiraz University of Medical Sciences

Latest degree

Specialist

Other areas of specialty/work

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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

Not applicable