

Clinical Trial Protocol

Iranian Registry of Clinical Trials

07 Jul 2026

Study of the effect of hand reflexology on pain, anxiety and hemodynamic parameters in children who are candidates for surgery during Insertion of the Intravenous (IV) Catheter

Protocol summary

Study aim

The effect of hand reflexology on pain, hemodynamic exercises in children undergoing surgery during intravenous line

Design

The present study is a double blind randomized clinical trial with two intervention and control groups. The number of 88 children eligible for the study were entered into the study using the available method and were randomly divided into two groups of reflexology and control using the block method (blocks of four and with an allocation ratio of 1:1 using a random table).

Settings and conduct

Subjects will be divided into two equal control and intervention groups using block randomization method. Reflexology will be performed for 5 minutes before placing the intravenous line in the intervention group. Two trained nurses will measure pain intensity, anxiety and hemodynamic parameters independently. This study will be conducted in the hospitals of Qom University of Medical Sciences

Participants/Inclusion and exclusion criteria

Inclusion criteria: 3-6 year old children with full consciousness and physiological stability Not having mental disability Decreased sensitivity to pain caused by any other disease No use of painkillers, sedatives, corticosteroids or relaxants Hospitalization for the first time Absence of pain during sampling Exclusion criteria: Inability to place an IV line in the first step and the child's non-cooperation during the implementation of interventions

Intervention groups

In order to perform the intervention in the intervention group, the researcher will perform a massage on the skin of the hand with slow stroking movements with moderate pressure of the palm five minutes before taking the vein and during insertion. In the control group,

the establishment of the venous route will be done without intervention and based on the department's routine

Main outcome variables

Pain, hemodynamic parameters remain stable

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20221205056718N2**

Registration date: **2023-02-04, 1401/11/15**

Registration timing: **prospective**

Last update: **2023-02-04, 1401/11/15**

Update count: **0**

Registration date

2023-02-04, 1401/11/15

Registrant information

Name

Hossein Sharafi

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2023-02-20, 1401/12/01

Expected recruitment end date

2023-04-21, 1402/02/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Study of the effect of hand reflexology on pain, anxiety and hemodynamic parameters in children who are candidates for surgery during Insertion of the Intravenous (IV) Catheter

Public title

The effect of hand reflexology on pain, anxiety and hemodynamic parameters

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

3-6 year old children Full consciousness Physiological stability Not having mental disability Decreased sensitivity to pain caused by any other disease No use of painkillers, sedatives, corticosteroids or relaxants Hospitalization for the first time Absence of pain during sampling

Exclusion criteria:

Inability to place an IV line in the first attempt Child's non-cooperation during interventions

Age

From **3 years** old to **6 years** old

Gender

Both

Phase

N/A

Groups that have been masked

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

Sample size

Target sample size: **88**

Randomization (investigator's opinion)

Randomized

Randomization description

children who are eligible for the study are entered into the study using the available method and are randomly assigned to two intervention and control groups using the block method (blocks of four and with an allocation ratio of 1:1 using a random number table).

Blinding (investigator's opinion)

Double blinded

Blinding description

In this study, the participants will not know about the study due to their young age, and only the child's parents will know. Also, if the child refuses touch and reflexology, he will be excluded from the study. Two nurses who complete the evaluation questionnaires. And the data analyst will not know about the control and intervention groups. After the intervention, the two nurse

assessors will enter the operating room and evaluate the child at the designated times.

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Research Ethics Committees of Qom University of Medical Sciences

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Approval date

2023-01-30, 1401/11/10

Ethics committee reference number

IR.MUQ.REC.1401.227

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

anxiety

ICD-10 code

F41.1

ICD-10 code description

Generalized anxiety disorder

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

Pain related to Reflexology

Timepoint

Before, immediately and 5 minutes after the intervention

Method of measurement

Face, Legs, Activity, Cry, Consolability scale

2**Description**

Anxiety related to Reflexology

Timepoint

Before, immediately and 5 minutes after the intervention

Method of measurement

Modified Yale Preoperative Anxiety Scale

Secondary outcomes

1

Description

Hemodynamic parameters

Timepoint

Before, immediately and 5 minutes after the intervention

Method of measurement

Digital heart monitoring system

Intervention groups

1

Description

Intervention group: Different parts of the hands and arms, with special attention to the areas related to the central nervous system, pituitary gland, spine, solar plexus and head reflexes of patients. For example, the middle of the thumb, which is related to the pituitary reflex, and the side that is related to the spinal reflex. This will be done using a sliding thumb motion, pressing and moving in each area

Category

Treatment - Other

2

Description

Control group: In the control group, the venous route will be established without intervention and based on the routine of the department. The variables in the control group will be measured as before in the intervention group, immediately after vein extraction and 5 minutes after placing the venous catheter by the same two nurses.

Category

N/A

Recruitment centers

1

Recruitment center

Name of recruitment center

Qom Shahid Beheshti Hospita

Full name of responsible person

Majid Moghadam

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Recruitment center

Name of recruitment center

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Full name of responsible person

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3

Recruitment center

Name of recruitment center

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Full name of responsible person

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Ghous University of Medical Sciences

Full name of responsible person

Mehdi Mesri

Street address

Qom - Shahid Lavasani St. (Saheli) - Qom University
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Grant name

Grant code / Reference number

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

No

Title of funding source

Ghous 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

Ghous University of Medical Sciences

Full name of responsible person

Hossein Sharafi

Position

Instructor

Latest degree

Master

Other areas of specialty/work

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Web page address

Sharing plan

Deidentified Individual Participant Data Set (IPD)

Yes - There is 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
Yes - There is a plan to make this available
Data Dictionary
Yes - There is a plan to make this available
Title and more details about the data/document
All data is potentially shareable after de-identifying individuals
When the data will become available and for how long
The second half of 2023
To whom data/document is available

There is no restriction on access to the results
Under which criteria data/document could be used
There are no restrictions after de-identifying people
From where data/document is obtainable
hossein sharafi tell:00989173596990 email:
h_sharafi68@yahoo.com hsharafi@muq.ac.ir
What processes are involved for a request to access data/document
The request for the data that is irrefutably identified will be made through the e-mail introduced by the researcher, and the request will be reviewed in less than 10 days and the requested data will be provided to the requester.
Comments