

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 Jul 2026

Comparison of the results of chronic anal fissure surgery between conventional internal partial sphincterotomy and tailored sphincterotomy of string suture anoscope

Protocol summary

Study aim

Comparison of the results of chronic anal fissure surgery between conventional internal partial sphincterotomy and tailored sphincterotomy of string suture anoscope

Design

A single-blind, randomized clinical trial study with parallel groups and phases 3 on 60 patients. Randomization will be done with the block randomization method using Random allocation software.

Settings and conduct

This study will be conducted in patients with chronic anal fissure undergoing surgery in Urmia Imam Khomeini Hospital. Patients will be randomly assigned to one of two groups: conventional partial internal sphincterotomy or tailored sphincterotomy. The study will be a single-blinded study.

Participants/Inclusion and exclusion criteria

Patients with a history of chronic anal fissure lasting more than one month will be included in the study. The main exclusion criteria will be acute fissures, coexisting fissure with fistula or inflammatory bowel disease, a history of systemic diseases, a history of anal surgery (for fissure, fistula, or hemorrhoids), fecal incontinence.

Intervention groups

Patients will be randomly assigned via block randomization into Conventional partial internal sphincterotomy or tailored sphincterotomy. In conventional Sphincterotomy, 30% of the exposed internal sphincter will be cut. The incision site will be repaired with absorbable sutures and dressed. In tailored Sphincterotomy group, a No. 34 purse-string anoscope will be inserted and rotated circumferentially. Any areas of tightness will be incised using electrocautery. The fissure will be assessed, and thickened edges or apical skin tags (if present) will be excised. The surgeon will ensure that the electrocautery does not extend beyond the dentate line. The extent of sphincterotomy will be

precisely adjusted based on fissure height.

Main outcome variables

Complete treatment of Fischer; fecal incontinence, gas incontinence

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20210919052515N6**

Registration date: **2025-06-20, 1404/03/30**

Registration timing: **prospective**

Last update: **2025-06-20, 1404/03/30**

Update count: **0**

Registration date

2025-06-20, 1404/03/30

Registrant information

Name

Naser Masoudi

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 44 3337 9924

Email address

masoudi.n@umsu.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2025-07-23, 1404/05/01

Expected recruitment end date

2025-12-22, 1404/10/01

Actual recruitment start date

empty

Actual recruitment end date
empty

Trial completion date
empty

Scientific title
Comparison of the results of chronic anal fissure surgery between conventional internal partial sphincterotomy and tailored sphincterotomy of string suture anoscope

Public title
Comparison of the two methods of conventional internal partial sphincterotomy and tailored sphincterotomy of string suture anoscope in chronic anal fissure surgery

Purpose
Treatment

Inclusion/Exclusion criteria
Inclusion criteria:
 Age between 18-65 old years Chronic anal fissure lasting more than one month Exhibiting signs of chronicity upon anal examination such as the presence of a skin tag Visible fibers of the internal anal sphincter at the base of the wound
Exclusion criteria:
 Patients with acute fissures Coexisting fissure with fistula or inflammatory bowel disease A history of systemic diseases A history of anal surgery (for fissure, fistula, or hemorrhoids) Fecal incontinence Perineal tears following delivery Prior radiotherapy Use of corticosteroids or other immunosuppressive drugs pregnant women

Age
From **18 years** old to **65 years** old

Gender
Both

Phase
N/A

Groups that have been masked

- Participant

Sample size
Target sample size: **60**

Randomization (investigator's opinion)
Randomized

Randomization description
Patients will be divided into two methods of surgery using block randomization based on generated numbers by random allocation software. So that, in this software, the number of groups and the total number of the sample size will be entered and then in the block section, the block randomization method will be implemented. Patients will be allocated to two groups based on generated numbers.

Blinding (investigator's opinion)
Single blinded

Blinding description
The study will be conducted as a single-blind clinical trial. The patient will be blinded about surgery methods.

Placebo
Not used

Assignment
Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics Committee of Imam Khomeini Hospital, Urmia University of Medical Sciences

Street address

Imam Khomeini hospital, Ershad street, Modarres Blvd., Urmia, Iran.

City

Urmia

Province

West Azarbaijan

Postal code

57157-81351

Approval date

2025-03-12, 1403/12/22

Ethics committee reference number

IR.UMSU.HIMAM.REC.1403.139

Health conditions studied

1

Description of health condition studied

Chronic anal fissure

ICD-10 code

K60.1

ICD-10 code description

Chronic anal fissure

Primary outcomes

1

Description

Complete treatment of Fischer

Timepoint

Weeks one, four, eight, and 6 months after surgery

Method of measurement

Clinical examination

2

Description

Gas incontinence

Timepoint

Weeks one, four, eight, and 6 months after surgery

Method of measurement

Questioning the patient

3

Description

Fecal incontinence
Timepoint
Weeks one, four, eight, and 6 months after surgery
Method of measurement
Questioning the patient

Secondary outcomes

1

Description
Pain score
Timepoint
After surgery during recovery
Method of measurement
Visual Analogue Scale (VAS)

2

Description
Dosage of analgesia
Timepoint
24 hours after surgery
Method of measurement
Milligram

Intervention groups

1

Description
Intervention group: In tailored Sphincterotomy group , a No. 34 purse-string anoscope will be inserted and rotated circumferentially. Any areas of tightness will be incised using electrocautery. The fissure will be assessed, and thickened edges or apical skin tags (if present) will be excised. The surgeon will ensure that the electrocautery does not extend beyond the dentate line. The extent of sphincterotomy will be precisely adjusted based on fissure height.
Category
Treatment - Surgery

2

Description
Control group: . In conventional partial Sphincterotomy, 30% of the exposed internal sphincter will be cut. The incision site will be repaired with absorbable sutures and dressed.
Category
Treatment - Surgery

Recruitment centers

1

Recruitment center
Name of recruitment center
Urmia Imam Khomeini hospital
Full name of responsible person
Dr. Naser Masoudi

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Imam Khomeini hospital., Ershad street., Modarres Blvd., Urmia, Iran.
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masoudi.n@umsu.ac.ir

Sponsors / Funding sources

1

Sponsor
Name of organization / entity
Oroumia University of Medical Sciences
Full name of responsible person
Dr. Saber Gholizadeh
Street address
Urmia University of Medical Sciences, Resalat street, Jihad Blvd., Urmia, Iran.
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5714783734
Phone
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Email
saber@umsu.ac.ir
Grant name
Grant code / Reference number
Is the source of funding the same sponsor organization/entity?
Yes
Title of funding source
Oroumia 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
Oroumia University of Medical Sciences
Full name of responsible person
Dr. Naser Masoudi

Position

Associate professor

Latest degree

Subspecialist

Other areas of specialty/work

General Surgery

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Person responsible for updating data

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

No - There is not a plan to make this available

Justification/reason for indecision/not sharing IPD

No further information available.

Study Protocol

No - There is not a plan to make this available

Statistical Analysis Plan

Not applicable

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

Not applicable

Data Dictionary

Not applicable