

Clinical Trial Protocol

Iranian Registry of Clinical Trials

11 Sep 2026

Investigating the effect of pelvic floor muscle training exercises along with radiofrequency or biofeedback compared to pelvic floor muscle training exercises alone on urinary incontinence and sexual dysfunction in women with urinary incontinence.

Protocol summary

Study aim

our purpose is to investigate the effectiveness of pelvic floor muscle strengthening exercises with the help of biofeedback or radio frequency compared to pelvic floor muscle exercises alone in the treatment of stress urinary incontinence and sexual disorders in women.

Design

A clinical trial with a control group, with two parallel groups, a blind strain, randomized on 45 patients. Block randomization is used for randomization.

Settings and conduct

Women living in the city of Noorabad, Mamsani, Fars who are eligible to enter the study are invited to participate. After obtaining informed consent and performing clinical examinations, the outcome variables are measured and based on the grouping, the patients in each group receive the desired treatment during the sessions. After the end of the last treatment session and one month after, the outcome variables will be measured.

Participants/Inclusion and exclusion criteria

Women aged 35 to 55 years with stress urinary incontinence and also sexual disorders. Absence of chronic diseases, urinary infections, copper IUD, pregnancy and heart pacemaker and not receiving other incontinence treatments..

Intervention groups

The RF+ PFMT group receives five sessions of radiofrequency therapy and 15 sessions of pelvic floor muscle strengthening exercises (5 weeks). - The Biofeedback+PFMT group receives 15 therapeutic sessions of pelvic floor muscle strengthening exercises with biofeedback during 5 weeks. - The PFMT group receives fifteen therapeutic sessions of pelvic floor muscle strengthening exercises during 5 weeks.

Main outcome variables

the performance (strength and endurance) of the pelvic

floor muscles using perineometer urine lost using the one-hour test pad incontinence symptoms and sexual function using ICIQ-SF, ICIQ-VS, FSFI questionnaires treatment satisfaction using Likert scale

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20241014063359N1**

Registration date: **2024-10-31, 1403/08/10**

Registration timing: **prospective**

Last update: **2024-10-31, 1403/08/10**

Update count: **0**

Registration date

2024-10-31, 1403/08/10

Registrant information

Name

Zahra Ardekani

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2024-11-18, 1403/08/28

Expected recruitment end date

2025-02-16, 1403/11/28 Actual recruitment start date empty Actual recruitment end date empty Trial completion date empty	
Scientific title Investigating the effect of pelvic floor muscle training exercises along with radiofrequency or biofeedback compared to pelvic floor muscle training exercises alone on urinary incontinence and sexual dysfunction in women with urinary incontinence.	
Public title Investigating the effect of radiofrequency and biofeedback in urinary incontinence and impotence	
Purpose Treatment	
Inclusion/Exclusion criteria Inclusion criteria: - Women aged 35 to 55 with urinary incontinence as the main clinical complaint, whose urine loss of more than 1 gram per hour can be confirmed by the test pad, and who have sexual disorders and impotence at the same time. Exclusion criteria: Patients with chronic degenerative diseases that affect muscle and nerve tissues. Presence of any degree of pelvic organ prolapse Active or frequent urinary tract infections Vulvovaginitis Atrophic vaginitis Absence of a copper IUD in the uterus Patients who are pregnant or have given birth less than 6 months ago. Type 1 and 2 diabetes neurological disease Mental illness taking drugs that affect urination History of surgical or drug treatment for urinary incontinence Chronic debilitating diseases such as kidney failure Those who have a pacemaker. If the patient does not want to continue to cooperate at any stage of the study, does not complete the treatment sessions, and the patient's condition changes in such a way that they lose any of the entry criteria, the participants will be excluded from the study.	
Age From 35 years old to 55 years old	Blinding (investigator's opinion) Single blinded
Gender Female	Blinding description The patient receives the type of intervention or control group in sealed envelopes that are coded. Coding is done by one of the colleagues of the project. The evaluator and the person analyzing the data are blind to the grouping of the participants.
Phase N/A	Placebo Not used
Groups that have been masked - Outcome assessor - Data analyser	Assignment Parallel
Sample size Target sample size: 45	Other design features
Randomization (investigator's opinion) Randomized	Secondary Ids empty
Randomization description The grouping method is also a type of random grouping in the block method where people are placed in three groups. In the current clinical trial study (two intervention groups and one control group), it will include	Ethics committees 1
45 samples, which will be done with the block randomization method according to the following process. The size of the used block is 3, and therefore, the combination of these modes for the control group and the patient groups, which are displayed with the letters C, T1 and T2 respectively, will include 6 modes. which will include (T2T1C, T1T2C, T1CT2, CT2T1, and CT1T2, T2CT1,). Blocks will be selected randomly and with the help of Excel software, so that 10 blocks are randomly selected, and therefore 45 samples can be included in the study in a random sequence, which can be included in each control and treatment group. The number of blocks and how they are executed are done by hiding them inside the envelope. In this method, the blocks are numbered based on a random sequence and placed inside the envelopes, and the researcher assigns them to the intervention and treatment groups based on the order of arrival of the patients.	Ethics committee Name of ethics committee Ethics Committee of the University of Rehabilitation Sciences and Social Health
Blinding (investigator's opinion) Single blinded	Street address University of Rehabilitation Sciences and Social Health, kodakyar Dead End, Daneshjoo boulevard, Velenjak
Blinding description The patient receives the type of intervention or control group in sealed envelopes that are coded. Coding is done by one of the colleagues of the project. The evaluator and the person analyzing the data are blind to the grouping of the participants.	City Tehran
Placebo Not used	Province Tehran
Assignment Parallel	Postal code 1985713871
Other design features	Approval date 2024-07-02, 1403/04/12
Secondary Ids empty	Ethics committee reference number IR.USWR.REC.1403.113
	Health conditions studied

1

Description of health condition studied

Urinary incontinence

ICD-10 code

R32

ICD-10 code description

Unspecified urinary incontinence

Primary outcomes

1

Description

Performance (strength and tolerance) of the pelvic floor muscles

Timepoint

Before the start of the intervention, the end of the last session of the intervention, one month later

Method of measurement

using perineometer device

2

Description

The quantitative amount of urine lost

Timepoint

Before the start of the intervention, the end of the last session of the intervention, one month later

Method of measurement

using one hour pad test

3

Description

Incontinence symptoms and sexual function

Timepoint

Before the start of the intervention, the end of the last session of the intervention, one month later

Method of measurement

Using the valid Persian version of ICIQ-SF, ICIQ-VS, FSFI questionnaires

4

Description

Satisfaction with treatment

Timepoint

Before the start of the intervention, the end of the last session of the intervention, one month later

Method of measurement

using Likert scale

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: The RF+ PFMT group receives five

radiofrequency therapy sessions as weekly sessions, with the settings that will be mentioned below, and 15 sessions of pelvic floor muscle strengthening exercises during 5 weeks. At first, patients will perform 3 exercises to strengthen the pelvic floor muscles. In radiofrequency treatment, using a standard technique in which the tip of the intravaginal probe of the device is moved back and forth on the mucosal surface of the vagina and the entire front wall of the vagina using a special probe slowly with wide movements in the treated area, and radiofrequency energy and will remain in direct contact with the tissue for 10 minutes at a temperature of 43°C (between 41 and 45°C based on patient tolerance). Bipolar radiofrequency with a frequency of one megahertz, maximum power of 65 watts will be applied to patients. The radio frequency power will be adjusted to maintain the required temperature.

Category

Rehabilitation

2

Description

Intervention group: The Biofeedback+PFMT group receives fifteen therapeutic sessions of pelvic floor muscle strengthening exercises with biofeedback during 5 weeks. At the beginning of the pelvic floor muscle strengthening exercises, patients are given the necessary anatomical information with the help of biofeedback, and the exercises are taught one-on-one by the therapist. Patients are asked to empty their bladder before the procedure. They lie on their backs with their knees slightly bent and their heads slightly raised. Surface EMG probes are placed on the perineum at the three and nine o'clock positions, an additional neutral probe is placed on the patella, and patients are observed. Patients are asked to contract only their pelvic floor muscles, not their abdominal muscles. They are also asked to track the contraction and relaxation of their pelvic floor muscles on a monitor to make sure they are contracting the correct muscle group. Therefore, it enables active participation in the educational program. In this way, patients are taught how to identify their pelvic floor muscles and how to use their pelvic floor muscles selectively without using their abdominal muscles. After training the correct contraction, the patients are asked to do three pelvic floor muscle strengthening exercises.

Category

Rehabilitation

3

Description

The PFMT group receives fifteen therapeutic sessions of pelvic floor muscle strengthening exercises during 5 weeks. Patients are asked to perform the following three exercises based on the exercise program table. Faucet exercise: repeatedly contract and release your pelvic floor (such as closing and opening the faucet)- Elevator exercise: slowly contract the pelvic floor for 5 counts - hold for 5 counts - release for 5 counts (such as going up in the elevator for 5 counts - holding for 5 counts at the

top floor - coming down with a count of 5) - Coughing or sneezing

Category

Rehabilitation

Recruitment centers

1

Recruitment center

Name of recruitment center

Taskin physiotherapy clinic

Full name of responsible person

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

University of social welfare and rehabilitation sciences

Full name of responsible person

Dr. Mohammad Saied Khanjani

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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

University of social welfare and rehabilitation 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

University of social welfare and rehabilitation sciences

Full name of responsible person

Nahid Rahmani

Position

assistant professor

Latest degree

Ph.D.

Other areas of specialty/work

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Person responsible for updating data

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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

Not applicable

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

Not applicable

Data Dictionary

Not applicable

Title and more details about the data/document

Total potential data after de-identifying individuals

When the data will become available and for how long

The access period starts 6 months after the results are published

To whom data/document is available

Researchers working in academic and scientific institutions

Under which criteria data/document could be used

For use in systematic review studies and meta-analysis

From where data/document is obtainable

Zahraardekanipt@gmail.com

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

Please send your written request and a full description of the reason for the data request to the email address provided. After reviewing your request, your email will be answered within ten working days.

Comments