

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

09 Jul 2026

### The effect of telenursing based training on the self-efficacy of patients with stoma.

#### Protocol summary

##### Study aim

Determining the effect of telenursing based training on the self-efficacy of patients with stoma in Zahedan city in 2022

##### Design

First, we select 80 patients with stoma who meet the study entry criteria as available sampling, then, using blue and green color cards that are in the white envelopes, we randomly divide the patients into two groups of 40 people, intervention and control.

##### Settings and conduct

Patient information is obtained from the archives of hospitals covered by Zahedan University of Medical Sciences. Patients will be selected who have had at least one month since their ostomy surgery; Because they have faced problems and limitations.

##### Participants/Inclusion and exclusion criteria

The patient has not participated in stoma care training courses A member of the patient's family should not be part of the treatment staff The patient is not from the treatment staff The patient should not be supported by the Iran Ostomy Association At least one month has passed since the patient's ostomy surgery The age of people should be between 14 and 70 years The patient or their family caregiver has the ability to work with software and smart phones The patient does not have any impairment in speech, hearing or vision The patient has a smart phone The patient must live in Zahedan city

##### Intervention groups

patients are randomly divided into two control and intervention groups after available sampling. The control group only sees routine trainings; the intervention group, in addition to routine trainings, receives the special training of the researcher, which is prepared in the form of an application installed on the smart mobile phone of the patients and they are trained. Patients' self\_efficacy is measured by the stoma self\_efficacy scale questionnaire before the intervention, one and three months after the intervention.

#### Main outcome variables

Self\_efficacy

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20230303057596N1**

Registration date: **2023-07-31, 1402/05/09**

Registration timing: **registered\_while\_recruiting**

Last update: **2023-07-31, 1402/05/09**

Update count: **0**

##### Registration date

2023-07-31, 1402/05/09

##### Registrant information

##### Name

Mostafa Parsa

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 51 5542 1977

##### Email address

parsam3@mums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2023-07-21, 1402/04/30

##### Expected recruitment end date

2023-08-01, 1402/05/10

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

**Trial completion date**

empty

**Scientific title**

The effect of telenursing based training on the self-efficacy of patients with stoma.

**Public title**

The effect of telenursing based training on the self-efficacy of patients with stoma.

**Purpose**

Education/Guidance

**Inclusion/Exclusion criteria****Inclusion criteria:**

The patient has not participated in stoma care training courses A member of the patient's family should not be part of the treatment staff The patient is not from the treatment staff The patient should not be supported by the Iran Ostomy Association At least one month has passed since the patient's ostomy surgery The age of people should be between 14 and 70 years The patient or their family caregiver has the ability to work with software and smart phones The patient does not have any impairment in speech, hearing or vision The patient has a smart phone The patient must live in Zahedan city Also, patients who have a stoma due to cancer with grade 1 and 2 can enter the study

**Exclusion criteria:**

Cancer patients with grade 3 and 4

**Age**

From **14 years** old to **70 years** old

**Gender**

Both

**Phase**

N/A

**Groups that have been masked**

*No information*

**Sample size**

Target sample size: **80**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

Patients will be randomly divided into two groups of intervention and control by selecting colored cards that fall into a white envelope. Green and blue colored cards are prepared and randomly placed inside a white envelope and patients will randomly choose a card. Green is a sign of the patient's entry into the control group and the blue is a sign of the patient's entry into the intervention group.

**Blinding (investigator's opinion)**

Not blinded

**Blinding description****Placebo**

Not used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

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

Zahedan University of Medical Sciences (Research Ethics Committee)

**Street address**

Persian Gulf Highway, Salamat Blvd.

**City**

Zahedan

**Province**

Sistan-va-Balouchestan

**Postal code**

9816913396

**Approval date**

2022-11-20, 1401/08/29

**Ethics committee reference number**

IR.ZAUMS.REC.1401.420

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

stoma

**ICD-10 code**

Z93.3

**ICD-10 code description**

Colostomy status

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

self-efficacy score

**Timepoint**

Before starting the study, one month and three months after the intervention

**Method of measurement**

Stoma self-efficacy questionnaire

**Secondary outcomes**

empty

**Intervention groups****1****Description**

Intervention group: First, the self-efficacy questionnaire is given to patients to complete. Then in this group, in addition to the routine trainings that the patients receive, the researcher installs special trainings using an application on the patients' smart phones and teaches them. Patients have 3 weeks to be trained using the mobile application. One and three months after the intervention, the self-efficacy questionnaire will be given to the patients to be completed.

**Category**

Lifestyle

**2****Description**

Control group: First, the self-efficacy questionnaire is given to patients to complete. The educational program is not installed on the mobile phones of these patients. These patients are not given any special training. After one and three months, the self-efficacy questionnaire is given to the patients to be completed.

**Category**

Lifestyle

**Recruitment centers****1****Recruitment center****Name of recruitment center**

Ali Ibn Abi Talib Hospital

**Full name of responsible person**

DR.Amir Gergij

**Street address**

Ali Ibn Abi Talib Hospital

**City**

Zahedan

**Province**

Sistan-va-Balouchestan

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9816743111

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+98 54 3329 5570

**Email**

nur.mostafa.parsa@gmail.com

**Sponsors / Funding sources****1****Sponsor****Name of organization / entity**

Zahedan University of Medical Sciences, Research assistant

**Full name of responsible person**

Nur Mohammad Bakhshani

**Street address**

Dr. Hasabi Square - Zahedan University of Medical Sciences campus

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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**

Zahedan University of Medical Sciences, Research assistant

**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**

Zahedan University of Medical Sciences

**Full name of responsible person**

Mostafa Parsa

**Position**

Student

**Latest degree**

Bachelor

**Other areas of specialty/work**

Nursery

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Zahedan Nursing and Midwifery Faculty, Fellowship Square

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**Person responsible for scientific inquiries****Contact****Name of organization / entity**

Zahedan University of Medical Sciences

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## Person responsible for updating data

### Contact

**Name of organization / entity**  
Zahedan University of Medical Sciences  
**Full name of responsible person**  
Mostafa Parsa  
**Position**  
Student  
**Latest degree**  
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**Other areas of specialty/work**  
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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

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

Undecided - It is not yet known if there will be a plan to make this available

### Title and more details about the data/document

Questionnaire form and data analysis collected.

### When the data will become available and for how long

Starting the access period year1402

### To whom data/document is available

Researchers working in academic and scientific institutions

### Under which criteria data/document could be used

Statistical result.

### From where data/document is obtainable

Zahedan University of Medical Sciences Library Country  
Theses System Researcher Gmail:  
Nur.mostafa.parsa@gmail.com phone 09033321800

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

With a written request to the Research Vice\_Chancellor of Zahedan University of Medical

### Comments

Does not have