

Clinical Trial Protocol

Iranian Registry of Clinical Trials

08 Jul 2026

Evaluation of follow-up nurse plan and its effect on disease severity and dyspnea in patients with chronic obstructive pulmonary disease (COPD)

Protocol summary

Study aim

Evaluation of follow-up nurse plan and its effect on disease severity and dyspnea in patients with COPD

Design

A randomized, parallel- group controlled clinical trial on 88 patients. The randomization list will determine using a web-based randomization program.

Settings and conduct

The study location: Allameh Bohlool Hospital
The Intervention group: The frequency of follow-up of patients is determined based on the severity of the disease and the symptoms from three days to three months after discharge as follows: 1. The "Green area": once a month; 2. The "Yellow area": once a week; 3. The "Red area": Advising to go to the hospital emergency department immediately; follow-up by phone up to 6 hours later; follow-up twice a week after discharge from the hospital until they will be set at the "Green area".

Note: The frequency of follow-up could be determined by the relevant doctor's order. The control group will receive routine hospital training and follow-up. In order to preserve ethics, at least one routine follow-up will be done in the control group.

Participants/Inclusion and exclusion criteria

Definitive diagnosis of COPD; referring to the hospital's self-care and follow-up unit after discharge; being conscious; shortness of breath score more than 3 on the Borg scale; informed consent; resident of Gonabad city; access to landline or mobile; no speech, hearing and vision disorders and the ability to understand and answer questions; no history of mental illness

Intervention groups

This study has two intervention and control groups. In the intervention group, the frequency of follow-up of patients is determined from three days to three months after discharge based on the severity of the disease and the symptoms, or based on the relevant doctor's order. The control group will receive routine hospital training and follow-up.

Main outcome variables

Disease severity; dyspnea

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20191223045868N2**

Registration date: **2022-05-27, 1401/03/06**

Registration timing: **prospective**

Last update: **2022-05-27, 1401/03/06**

Update count: **0**

Registration date

2022-05-27, 1401/03/06

Registrant information

Name

Fatemeh Mohammadzadeh

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 51 5722 5213

Email address

f.mo6325@gmail.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2022-06-10, 1401/03/20

Expected recruitment end date

2022-12-11, 1401/09/20

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluation of follow-up nurse plan and its effect on disease severity and dyspnea in patients with chronic obstructive pulmonary disease (COPD)

Public title

Evaluation of follow-up nurse plan and its effect on disease severity and dyspnea in patients with chronic obstructive pulmonary disease (COPD)

Purpose

Health service research

Inclusion/Exclusion criteria

Inclusion criteria:

Patients with definitive diagnosis of COPD by a specialist Referring to the hospital's self-care and follow-up unit after discharge Being conscious Moderate to severe shortness of breath (Shortness of breath score more than 3 on the Borg scale) Informed consent Resident of Gonabad city Access to landline or mobile

Exclusion criteria:

Speech, hearing, and vision disorders and the disability to understand and answer questions History of mental illness based on self-reporting or patient records data

Age

From **18 years** old

Gender

Both

Phase

N/A

Groups that have been masked

- Data analyser

Sample size

Target sample size: **88**

Randomization (investigator's opinion)

Randomized

Randomization description

Random permutation blocks of sizes 2 and 4 will be used for randomization. The analyzer will determine the randomization list based on a web-based randomization program. The allocation will be concealed using opaque sealed envelopes with a random sequence.

Blinding (investigator's opinion)

Single blinded

Blinding description

Blinding will be done for the data analyzer. For the study groups, A and B codes are considered, and the data analyzer is not told the type of intervention.

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Research Ethics Committee of Gonabad University of Medical Sciences

Street address

Gonabad University of Medical Sciences, Asian Road Border

City

Gonabad

Province

Razavi Khorasan

Postal code

9691793718

Approval date

2022-04-11, 1401/01/22

Ethics committee reference number

IR.GMU.REC.1401.006

Health conditions studied

1

Description of health condition studied

chronic obstructive pulmonary disease

ICD-10 code

J44.9

ICD-10 code description

Chronic obstructive pulmonary disease, unspecified

Primary outcomes

1

Description

Disease severity

Timepoint

Before and three months after intervention

Method of measurement

COPD Assessment Test (CAT)

2

Description

Dyspnea

Timepoint

Before and three months after intervention

Method of measurement

Borg scale

Secondary outcomes

1

Description

Satisfaction of patients

Timepoint

Three months after intervention

Method of measurement
percentage- questionnaire

2

Description
Percentage of unscheduled referrals of discharged patients to hospital emergencies

Timepoint
Three months after intervention

Method of measurement
Hospital file

3

Description
Physical complications after discharge in relation to self-care at home

Timepoint
Three months after intervention

Method of measurement
Percentage of physical complications-questionnaire

4

Description
Re-hospitalization after discharge

Timepoint
Three months after intervention

Method of measurement
Percentage of Re-hospitalization after discharge-questionnaire

Intervention groups

1

Description
The intervention group: The follow-up nurse care plan for the intervention group includes: 1- Safe discharge procedures; 2- Introduce the patient to the follow-up nurse and send a summary of the patient history; 3- Initial evaluation of the patient by the nurse to continue the necessary training after discharge and 4- The main phase of follow-up, which will be called to the patient by phone 3 to 7 days after discharge according on the checklist. Then the nurse introduces him/herself, the appropriate follow-up is performed based on the relevant checklist, and recommendations are offered if the patient requires additional training. Also, if the patient needs support resources such as outpatient clinics, hospital self-care clinics, health centers, psychologists, addiction and smoking cessation centers, nutrition counselors, and home care centers, will refer the patient to the mentioned institutions. The frequency of follow-up of patients is determined based on the severity of the disease and the symptoms from three days to three months after discharge as follows: 1. The "Green area": once a month; 2. The "Yellow area": once a week; 3. The "Red area": Advising to go to the hospital emergency department immediately; follow-up by phone up to 6 hours later; follow-up twice a week after discharge from the hospital until they will be set at the "Green area".

Note: The relevant doctor's order could determine the frequency of follow-up.

Category
Rehabilitation

2

Description
Control group: The control group will receive routine hospital training and follow-up. In order to preserve ethics, at least one routine follow-up will be done in the control group.

Category
Rehabilitation

Recruitment centers

1

Recruitment center

Name of recruitment center
Allameh Bohlool Hospital-Gonabad
Full name of responsible person
Elham Saberi Noghabi
Street address
Parstar Ave., Gonabad University of Medical Sciences
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Fax
+98 51 5723 6160
Email
f.mo6325@gmail.com

Sponsors / Funding sources

1

Sponsor

Name of organization / entity
Gonabad University of Medical Sciences
Full name of responsible person
Leila Sadegh Moghadam
Street address
Gonabad University of Medical Sciences, Asian Road Border
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Province
Razavi Khorasan
Postal code
9691793718
Phone
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Email
is_moghadam@yahoo.com
Grant name

Grant code / Reference number
Is the source of funding the same sponsor organization/entity?

Yes

Title of funding source

Gonabad 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

Gonabad University of Medical Sciences

Full name of responsible person

Elham Saberi Noghabi

Position

Faculty instructor

Latest degree

Master

Other areas of specialty/work

Nursery

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Person responsible for scientific inquiries

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Latest degree

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Person responsible for updating data

Contact

Name of organization / entity

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Full name of responsible person

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Latest degree

Master

Other areas of specialty/work

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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

There is no more information

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

Yes - There is 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

Title and more details about the data/document

There is no more information.

When the data will become available and for how long

There is no more information.

To whom data/document is available

There is no more information.

Under which criteria data/document could be used

There is no more information.

From where data/document is obtainable

There is no more information.

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

There is no more information.

Comments